TOXICOLIGICAL AND BIOCHEMICAL STUDIES ON THE EFFECT OF SOME INSECT GROWTH REGULATORS ON SPODOPTERA LITTORALIS (BOISD.) LARVAE

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Abstract

The toxicological and biochemical effects of five insect growth regulators (IGRs) namely Cascade, Atabron, Consult, Match and Mimic against 2nd and 4th instar larvae of *Spodoptera littoralis* (Boisd.) were studied under laboratory conditions. Larvae were fed on castor bean leaves treated with 9 successive concentrations. The obtained results indicated that: Atabron proved to be the most potent insect growth regulator, whereas Mimic was the least toxic one among the tested IGRs against both 2nd and 4th instar larvae. The 4th instar larvae proved to be more sensitive to all the tested IGRs than the 2nd one at all tested concentrations.

In concerning the biochemical effects the obtained results indicated that:

The tested IGRs increased the total soluble protein after 2 days of treating 2nd and 4th S. littoralis larvae except Cascade at the two lower concentrations. However, a marked decrease was detected after 5 days of treatment both 2nd and 4th instars with Consult, Match and Cascade.

The tested IGRs decreased the activity of glutamic oxaloacetic transaminase (GOT), while it increased (glutamic pyruvic transaminase (GPT) activity at 2 days post treatment the 2nd instar larvae. The inverse was true at 5 days post treatment. In case of the 4th instar, the tested IGRs increased the activity of the two enzymes after 2 and 5 days of treatment.

Exposing the 2^{nd} and 4^{th} instar larvae to the tested IGRs increased markedly the activity of trehalase, invertase and amylase in both 2^{nd} and 4^{th} instars after 2 and 5 days except Consult and Mimic in case of invertase of the 4^{th} instar.

INTRODUCTION

The toxicological effects of some IGRs have been thourghly investigated against *S. littoralis,* Bayoumi *et al.* (1998) and Badr (2000). Using synthetic IGRs which interfere with cuticle deposition by inhibition of chitin synthesis, may become important in controlling insect pest populations that are resistant to conventional insecticides, due to the action of enzymes which are either insensitive to the

insecticide or able to degrade it to non-toxic metabolites. The reduced level of chitin in the cuticle seems to result from inhibition of biochemical process leading to chitin formations. Carbohydrates are contributed to the structures and functions of all insect tissues. Metabolism of carbohydrates are controlled mainly by invertase, amylase and trehalase enzymes which play a principal role in the digestion and utilization of carbohydrates by insects (Salem *et al.*, 1995). Trehalase is activated during moulting to generate production of glucose for chitin build-up. Invertase and amylase are two important digestive enzymes. The amino-transferases enzymes especially GPT is one of the components of oxidative metabolism of protein which in certain insects is utilized during the initial periods of flight (Bursell, 1963) and acts as a catalytic agent in the metabolism of carbohydrate (Katunuma *et al.*, 1968)

This study aimed to elucidate and find some biochemical relationship between treated IGRs and activities of some enzymes and total soluble protein in *S. littoralis*.

MATERIALS AND METHODS

1. Tested compounds:

- Flufenoxuron (Cascade 10 % EC)
- [4-[2-chloro 4-(trifluoromethyl phenoxy)] 2-flurophenyl 3-(2,6-difluorophenzoyl) urea. The recommended rate is 200 ml/feddan.
 - Chlorfluazuron (Atabron, IKI- 7899 5 % EC)
- 1- (2, 6, difluorobenzoy I3 [4 (chloro 5 trifluoromathyl-2-pyridyloxy)3,5,-dichlorophenyl] urea. The recommended rate is 400 ml/ feddan.
 - Benzoylphenyl urea (Consult 10 % EC)
- N (3, 5- dichloro 4 (1,1,2,2, tetrafluoroethoxy)-phenylamino) carbanyl 2,6-difluorobenzamide. The recommended rate is 200 ml/feddan.
 - Lufenuron (Match 5 % EC)
- $\label{eq:N-continuous} $$N-[[[2,5-dichloro-4-(1,1,2,3,3,3-hexafluoropropoxy]-(phenyl]amino]$ carbonyl]-2,6-difluoro-benzamide. The recommended rate is 400 ml/feddan.$
 - Tebufenozide (Mimic 24 % EC)
- 3,5- dimethyl benzoic acid 1- (1,1 dimethyl ethyl) 2-(4- ethylbenzoyl) hydrazide. The recommended rate is 350 ml/feddan.

Rearing technique: Susceptible strain of *S. littoralis* was reared on castor bean leaves according to El-Defrawi *et al.* (1964) and some modification of Khedr (2002) under laboratory conditions 25 ± 2 °C and 65 ± 5 % R. H.

Oral toxicity of the tested insect growth regulators against the second and the fourth instars of Spodoptera littoralis larvae;

The leaf dipping technique was applied to evaluate the insecticidal activity of tested insect growth regulators against the 2nd and 4th instar larvae. Successive concentrations, i.e. 100, 50, 25, 12.5, 6.25, 3.13, 1.56, 0.78 and 0.39 ppm for Cascade, Atabron, Consult and Match were prepared. However, the tested concentrations of Mimic were 420, 210, 105, 52.5, 26.25, 13.13, 6.56, 3.28 and 1.64 ppm.

Mortality counts were made after 1, 2, 3, 4 and 5 days of treatment. Mortality data were corrected according to Abbott's formula (1925).

The dosage mortality regression lines were statistically analyzed according Finney (1952).

- **3. Preparation of samples for biochemical assay** Samples of tested 4th instar larvae were prepared after 2 and 5 days of application. For each concentration applied, 5 larvae were picked up and placed in clean jars, then starved for 4 hr. The starved larvae were homogenized in distilled water (5 larvae/5 ml) using a teflon homogenizer surrounded with jacket of crushed ice for 3 minutes. The homogenate was centrifuged at 3500 rpm for 10 minutes at 5 °C. The supernatant was immediately assayed to determine total soluble protein and the activities of glutamic oxaloacetic transaminase (GOT), glutamic pyruvic transminase (GPT), trehalase, amylase, and invertase enzymes.
- 4. Determination of total soluble protein Colourimetric determination of total soluble protein in total homogenate of larvae of S. littoralis was carried out as described by Gornall et al. (1949).
- 5. Determination of enzymes activities:
- **5.1. Transaminase enzymes (GOT and GPT)** The activities of glutamic oxaloacetic transaminase (GOT) and glutamic pyruvic transminase (GPT) enzymes were determined colourimetically according to Reitman and Frankle (1957).

5.2. Carbohydrate hydrolyzing enzymes (trehalase, invertase and amylase)

Determination of trehalase, amylase and invertase enzymes activities in digesting trehalose, starch and sucrose, respectively were determined according to Ishaaya and Swiriski (1976).

RESULTS AND DISCUSSION

Oral toxicity of the tested insect growth regulators against the second and fourth instars of *Spodoptera littoralis* larvae:

i. The 2nd instar larvae:

Comparison on the basis of LC₅₀ **and LC**₉₀ **values** The tested toxicants could be arranged discerningly according to their potency against 2^{nd} instar larvae of *S. littoralis* at the LC₅₀ level as follows: Atabron, Cascade, Consult, Match and Mimic. The corresponding concentrations (LC₅₀) are: 0.18, 0.22, 0.26, 0.34 and 1.60 ppm. (Table 1) whereas the toxicity lines are drawn in Figures 1 and 2.

The descending order of the tested toxicants at LC_{50} level is Atabron, Cascade, Match, Consult and Mimic. The respective LC_{90} values were 5.40, 5.80, 7.00, 80.00 and 105 ppm (Table 1). It is clear that Consult and Mimic changed their places at the LC_{90} level. This allocation of steepness of Mimic being less steeper than the line of the other compound.

However, Atabron proved to be the most potent insecticide, whereas Mimic was the least toxic one among the tested compounds against $2^{\rm nd}$ instar larvae.

-Toxicity index Obtained results show that the toxicity index at LC_{50} level is as follows: Cascade (81.82 %), Consult (69.23 %), Match (52.94 %) and Mimic (11.25 %) as toxic as Atabron.

Basing on the LC_{90} , the tested insect growth regulators could be divided into two groups according to their toxicity to 2^{nd} instar larvae as follows:

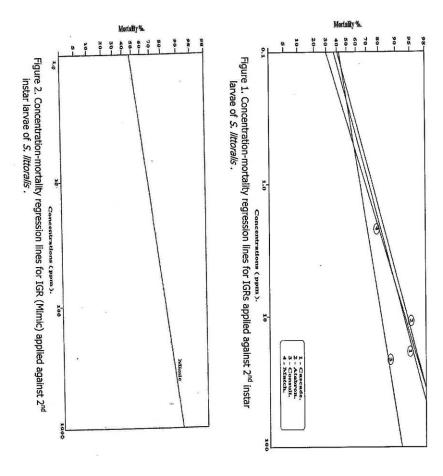
- 1- The first and the most toxic group includes, Atabron, Cascade and Match.
- 2- The second and the least toxic group include Consult and Mimic (Table 1).
 Similar data were also reported by EL-Ghareeb (1992) they stated that Atabron proved to be highly effective against S. littoralis.

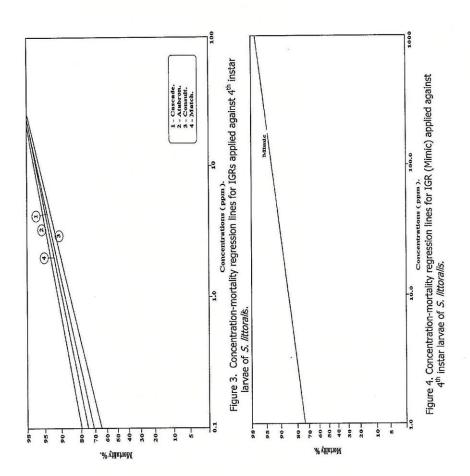
Table 1. Toxicity data of the tested insect growth regulators applied orally to 2nd and 4th instar larvae of S. littoralis under laboratory conditions.

	_	_					
*	LC ₉₀	4 th instar	100	57.14	47.05	30.76	2.67
Toxicity index *	07	2 nd instar	100	93.10	6.75	77.14	5.142
ř	LC ₅₀	2 nd instar	100	81.82	69.23	52.94	11.25
ъе.	4 th	instar	1.8549	1.9359	2.0287	1.4285	0.4802
Slope	2 _{nd}	instar	1.6579	1.7063	0.6589	1.7382	0.4477
5	4 th	instar	5.40	5.80	80	7.0	1050
3	2 nd	instar	0.80	1.40	1.70	2.60	30
LC ₅₀	2 nd instar		0.18	0.22	0.26	0.34	1.60
Field	200	7	100	100	100	100	420
Treatments			Atabron	Cascade	Consult	Match	Mimic

* =Toxicity index (Sun, 1950) = LC₅₀ or LC₅₀ of the other compound LC₅₀ or LC₅₀ of the efficient compound

× 100





ii. The 4th instar larvae:

- **-Comparison on the basis of LC₉₀ values** The tested toxicants could be arranged discerningly according to their potency against 4^{th} instar larvae of *S. littoralis* as follows: Atabron, Match, Cascade, Consult and Mimic. The corresponding lethal concentrations (LC₉₀) are 0.80, 1.40, 1.70, 2.60 and 30.0 ppm (Table 1), whereas the toxicity lines are shown in Figures 3 and 4. Atabron proved to be the most potent insect growth regulators, meanwhile Mimic was the least toxic one to 4^{th} instar larvae at LC₉₀ level.
- **-Toxicity index** Data of the toxicity index at LC_{90} levels are presented in Table 1. It is clear that the toxicity index is as follow: Match (57.14 %), Cascade (47.05 %), Consult (30.76 %) and Mimic (2.67 %) as toxic as Atabron.

Generally, the tested insect growth regulators could be divided into three groups according to their toxicity to 4th instar larvae. The first and the more toxic group include Atabron. The second group includes Match, Cascade and Consult, whereas the third and the least toxic one include Mimic.

Generally, the 4^{th} instar larvae proved to be more sensitive to all the tested IGRs than the 2^{nd} one at all concentrations.

One of the explanations of this phenomenon is that the food uptake of the 4^{th} instar larvae is greater than the 2^{nd} one that means greater uptake of the IGRs.

On the contrary, Badr (2000) reported that the 2nd instar larvae of *S. littoralis* are more sensitive than the 4th instar ones when fed on cotton leaves sprayed with Consult at 5 and 10 days after spray.

Due to the IGRs are chitin synthesis inhibitors, therefore their effect would appear after suppressing moulting process. As shown in Table 1, LC_{90} of the tested IGRs is greatly reduced as compared to the applied field rate. Subsequently, the malformations in the following stages (pupa and adult) would not exist as all the treated larvae died. Therefore Figure 5 a - c manifested the symptoms of death due to the exposure of larvae to the tested IGRs. The symptoms of death usually being at three days post treatment with different IGRs at different concentrations, larvae became inactive, alive without feeding, appeared to be wet, completely paralyzed, dark colour spread all over the body and viscous excretion comes out the larval body, then it became unable to ecdyse (Figure 5 a). In addition, (in case of Mimic at different concentrations) a partially exit of the posterior part of the alimentary canal

from the anus Figure 5. (b and c). Similarly Abd El-hakim (1996) showed some malformation effects in cotton bollworms treated with IGRs.

Biochemical effects insect growth regulators on Spodoptera littoralis larvae:

a- Total soluble protein Data in Table 2 show the level in the total soluble protein detected in the supernatant homogenate of 2nd and 4th instar larvae treated with different concentrations of the tested IGRs. It is obvious that all the tested soluble protein in both 2nd and 4th *S. littoralis* larvae after two days of treatment with the exception of Cascade, especially at lower concentrations.

The 4th instar larvae proved to be more sensitive to the tested IGRs than the 2^{nd} instar ones. For instance, the total soluble protein level ranged between 150 - 450 % in 4^{th} instar larvae while it ranged between 120 - 300 % in the 2^{nd} instar. Atabron was the most effective IGR.

The total soluble in the treated larvae after five days of treatment was greatly reduced in general with all the tested IGRs, expect Atabron down to a concentration equivalent to 1/128 as that of field rate on 4th instar.

In addition, Mimic at higher concentration caused slight increase in this parameter especially on the 4^{th} instar larvae.

In this connection, Abdel-Hafez *et al.* (1983) found that the IGRs, difflubenzuron and triflumuron caused reduction in the levels of protein and free amino acids of the treated *S. littoralis* larvae. Similarly, Mostafa (1993) showed significant decrease in the level of total soluble protein as affected by IGRs. Both flufenoxuron and teflubenzuron were more effective on the 6th instar larvae than 4th instar ones of *S. littoralis*.

Generally, the tested IGRs increased markedly the total soluble protein in the treated larvae in both instars after two days of exposure. On the contrary, the level of total soluble protein in the treated larvae was severely decreased after five days of application with the exception of Atabron against only 4th instar larvae and the higher concentrations of Mimic on 2nd and 4th instars.

Fourth instar larvae was more susceptible to the IGRs application than the $2^{\rm nd}$ instar ones.

b- Transaminase enzymes:

i. GOT Data in Table 3. indicate that the activity of GOT increased after two days as affected by Cascade (300-150 %) as control and Consult (350-175 %) and the

higher 2-3 concentrations of Match and Mimic induced, slight and the other concentrations had no effect or became under the control level.

On the contrary, all concentrations of Atabron at all concentrations and the lower concentration of Match and Mimic decreased the activity of GOT severely (80 - 205 %) as control. After five days of treatment the picture was completely inverse for all the tested IGRs. For instance, Cascade and Consult at all concentrations and Match and Mimic at the lower concentrations showed severe reduction in the activity for GOT enzyme (as 25-83.3 % as the control).

On the contrary, all concentrations of Atabron and Match induced marked increase in GOT activity recording between as 150 - 366.67 %, 116.67 - 316.67 %, and 133.33 - 200 % as the control, respectively.

In case of 4^{th} instar larvae treatment, data in Table 3 indicate that most concentrations of the tested IGRs increased activities of GOT in 4^{th} instar larvae at two days after treatment. Atabron was the most potent, as it recorded the highest activity as it ranged between 166.67 - 3333.3% as the control.

However, the first three concentrations of Match decreased the activity of GOT to 66.67 % for f.r. (recommended Field rate), 60 % for 1/2 f.r. and 40.33 % for 1/4 f.r. as that of the control (100 %). In case of Mimic, the last four concentrations decreased the enzyme activity, recording as 80, 50, 66.67 and 66.67 % as control for 1/32 f.r., 1/64 f.r. 1/128 f.r. and 1/256 f.r., respectively. The same trend was nearly noticed after five days, all IGRs at different concentrations increased the activity of GOT with the exception of Cascade at 1/256 f.r. and Match at 1/16 and 1/3 f.r. that recorded as 66.67, 50 and 66.67 % as the control, respectively. However, the higher two concentrations of Mimic did not cause any change in the enzyme activity.

ii. GPT Data in Table 4. indicate that all IGRs increased GPT activity as compared to control after two days of treating 2^{nd} instar larvae, expect Mimic at the least concentration that recorded 50 %. Atabron recorded the highest increase in GPT activity that ranged between 150.0 - 366.67 %.

Meanwhile, five days post treatment, Consult and Atabron caused decreasing activity that ranged between 20-75% and 25-83.33 %, respectively. Other IGRs increased GPT activity at the higher concentrations and decrease it at the lower ones.

Table 2. Total soluble protein content % in the supernatant of the homogenated *S. littoralis* larvae as affected by insect growth regulators treatment.

Time Time Attack of the control of t									2 nd i	2 nd instar									
f.r.* f.r. f.r. <t< td=""><td>Time</td><td></td><td></td><td></td><td>Aft</td><td>er 2 da</td><td>ays</td><td></td><td></td><td></td><td></td><td></td><td></td><td>Aft</td><td>er 5 da</td><td>ıys</td><td></td><td></td><td></td></t<>	Time				Aft	er 2 da	ays							Aft	er 5 da	ıys			
180 220 140 160 200 180 200 100 - - 80 80 50 280 280 260 300 240 250 150 - - 90 60 50 200 280 280 260 300 240 200 140 120 50 40 80 50 260 280 300 250 240 200 160 100	Conc. Treat.	f.r.*	12 f.r.	1/4 f.r.	1/8 f.r.	1/16 f.r.	¹ / ₃₂ f.r.	1/64 f.r.	1/ ₁₂₈ f.r.	1/256 f.r.	f.r.	1/2 f.r.	1/4 f.r.	1/8 f.r.	¹ / ₁₆ f.r.	¹ / ₃₂ f.r.	¹ / ₆₄ f.r.	¹ / ₁₂₈ f.r.	¹ / ₂₅₆ f.r.
280 300 280 300 440 250 150 150 60 60 60 50 200 200 180 200 180 160 180 160 180 160 180 160 180 160 180 160 180 160 180 160 180 160 180 160 180 160 180	Cascade	180	220	140	160	200.	180	200	100	100			80	80	20	09	40	40	20
200 200 150 160 180 200 140 120 50 80 70 140 120 50 80 70 100 80 260 280 280 250 240 200 180 160 120 - - 50 40 80 100 180 140 220 240 200 250 140 100<	Atabron	280	300	280	260	300	240	250	200	150			90	09	20	20	80	40	20
260 280 300 250 240 200 180 160 120 - - 50 40 80 200 180 100	Consult	200	200	150	160	180	200	200	140	120	20	80	70	100	80	90	20	20	50
200 180 140 220 240 200 140 140 160 <td>Match</td> <td>260</td> <td>280</td> <td>300</td> <td>250</td> <td>240</td> <td>200</td> <td>180</td> <td>160</td> <td>120</td> <td>t</td> <td></td> <td>20</td> <td>40</td> <td>80</td> <td>09</td> <td>100</td> <td>80</td> <td>100</td>	Match	260	280	300	250	240	200	180	160	120	t		20	40	80	09	100	80	100
100 100	Mimic	200	180	140	220	240	200	250	140	140	100	120	80	20	9	9	40	40	50
After 2 days 350 250 300 300 250 250 200 150 100	Control	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
350 250 300 250 300 150 <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>4th i</td> <td>nstar</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>									4 th i	nstar									
350 250 300 350 250 250 150 <td></td> <td></td> <td></td> <td></td> <td>Aft</td> <td>er 2 da</td> <td>sys</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>Aft</td> <td>er 5 da</td> <td>ıys</td> <td></td> <td></td> <td></td>					Aft	er 2 da	sys							Aft	er 5 da	ıys			
400 450 350 250 300 150 200 250 -	Cascade	350	250	300	300	250	250	200	150	100	1					80	40	80	09
200 180 150 120 130 150 200 120 200 120 40 40 200 150 200 200 250 250 350 300 -	Atabron	400	400	450	300	250	300	150	200	250	1	1	1	1	ı	220	160	140	80
200 150 200 200 250 250 350 300 -	Consult	200	200	180	150	120	130	150	200	120	·	ı		80	40	40	50	40	20
300 300 300 300 150 150 150 150 150 - 120 100 100 120 120 120 100 100 100 100	Match	200	150	200	200	200	250	250	350	300	,	1	ï	ī	1	20	20	20	20
100 100 100 100 100 100 100 100 100 100	Mimic	300	300	300	300	300	150	150	150	150	ř	120	100	100	120	80	80	9	40
	Control	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100

* f.r. = recommended field rate

Table 3. GOT enzyme % in the supernatant of the homogenated S. littoralis larvae as affected by insect growth regulators treatment.

					_			_	_			_		-		
		1/256 f.r.	33.33	366.67	25	250	166.67	100			29.99	166.67	109.33	200	333.33	100
		1/128 f.r.	33.33	366.67	33.33	266.67	133.33	100			109.33	250	140	160.67	320	100
		1/64 f.r.	33.33	333.33	20	116.67	166.67	100			120	300	180	120	320	100
	ays	1/32 f.r.	83.33	350	40	200	200	100		ays	166.67	333.33	250	29.99	300	100
	After 5 days	1/16 f.r.	29.99	150	50	316.67	150	100		After 5 days	200	350	266.33	20	266.67	100
		1/8 f.r.	75	166.67	29.99	233.33	133.33	100					300	-	200	100
		7.4 f.r.	83.33	200	75	250	100	100					300		100	100
		7. T.			33.33	1	20	100			ī	-	-		100	100
		f.r.	1	-	100		20	100			,		,	,	i.	100
star		1/256 f.r.	150	20	200	20	25	100	ıstar		200	180	133.33	200	66.67	100
2 nd instar		1/128 f.r.	175	25	212.50	25	20	100	4 th instar		120	166.67	140	250	29.99	100
		1/64 f.r.	275	25	212.50	25	37.50	100			166.67	220	120	266.67	20	100
		1/32 f.r.	237.50	35	250	20	100	100		10	200	266.67	120	166.67	80	100
	After 2 days	1/16 f.r.	300	20	300	62.50	100	100		After 2 days	166.67	240	109.33	266.67	230	100
	AH	1/8 f.r.	262.50	40	300	75	75	100		Afte	266.67	250	29.99	100	220	100
		1/4 f.r.	237.50	20	237.50	100	125	100			100	300	66.67	40.33	166.67	100
		½ f.r.	300	75	350	112.50	112.50	100			166367	300	266.67	09	233.33	100
		f.r.*	200	80	175	175	100	100			200	333.33	100	66.67	200	100
	Time	Conc.	Cascade	Atabron	Consult	Match	Mimic	Control			Cascade	Atabron	Consult	Match	Mimic	Control

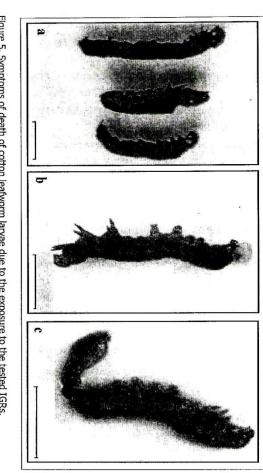


Figure 5. Symptoms of death of cotton leafworm larvae due to the exposure to the tested IGRs.

In case of 4^{th} instar, an increase in the activity of GPT that observed after two days of treatment with Cascade and Atabron at all concentrations. However, a gradual decrease in the activity of GPT was noticed after treatment with Consult, Match and Mimic, that ranged between 220-50 %, 266.67-75 % and 350-75 %, respectively.

Five days after treating 4th instar larvae with Cascade and Atabron the activity of GPT was increased at all tested concentrations. The higher concentrations of Consult, Match and Mimic increased GPT activity, while the lower ones caused a noticeable decrease in the activity.

In conclusion, nearly all or most concentrations of tested IGRs increased GOT and GPT enzymes activities in the treated 4^{th} instar larvae of *S. littoralis*. Atabron was the most effective IGR.

These results are in agreement with those of Salem *et al.* (1995) working on the chitin synthesis inhibitor diafenthiuron, buprofezin and pyriproxyfen against *S. littoralis* 4^{th} instar larvae after 24, 48 and 72 hours.

Most of the tested concentrations applied for all IGRs against 2nd instar larvae after two days of treatment showed decrease in GPT activity, while the inverse was true after five days of treatment. In this respect, Abdel-Hafez *et al.* (1993) found reduction in GOT activity and increased GPT after treatment of laboratory strain larvae of *S. littoralis* with diflubenzuron and flufenoxuron. On the other hand, Mostafa (1993) and EL-Kordy *et al.* (1995) showed significant increase in GOT activity and significant reduction in GPT activity in the 4th and 6th instar larvae after treatment with three IGPs

c- Carbohydrate hydrolyzing enzymes:

i. Trehalase Data given in Table 5. summarize the changes of trehalase in both 2nd and 4th instars of *S. littoralis* larvae. The 2nd instar larvae showed an increase in the trehalase activity two days after treating with Mimic at all concentrations nearly and Match till 1/16 f.r., whereas rest of IGRs at most concentrations reduce the enzyme activity. The highest activity was recorded at f.r. for Mimic, whereas the least value (40 %) was recorded at 1/256 and 1/128 f.r. consult and Match, respectively.

After five days of treatment the 2^{nd} instar larvae treated with the different concentrations of tested IGRs increased trehalase activity. The highest value (375 %) was recorded at 1/16 f.r. for Atabron.

Table 4. GPT enzyme % in the supernatant of the homogenated S. littoralis larvae as affected by insect growth regulators treatment.

							2 nd instar	star									
			Afte	After 2 days									After 5 days	lays			
, <u>;</u>	½ f.r.	4.r.	1/8 f.r.	1/16 f.r.	1/32 f.r.	1/64 f.r.	1/128 f.r.	1/256 f.r.	j.	۲. کړ ۲. ت.	% 7. °. °. °. °. °. °. °. °. °. °. °. °. °.	1/8 f.r.	1/16 f.r.	¹ / ₃₂ f.r.	1/64 f.r.	1/128 f.r.	1/256 f.r.
250	200	220	100	100	180	175	200	150	ı	,	175	150	125	75	20	20	25
366.67	350	350	333.3.3	300	250	200	150	150	ı	,	83.33	66.67	75	20	33.33	20	25
150	200	300	350	120	175	150	220	100	100	100	75	29.99	20	29.99	33.33	40	20
28	150	100	100	220	175	180	200	250	1		225	200	125	100	75	20	20
150	250	300	200	175	150	100	100	20	150	250	225	175	166.67	120	100	100	20
100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
							4 th ins	instar									
			Affe	After 2 days									After 5 days	days			
366.66	350	350	333.33	300	250	280	266.67	250	,	,			283.33	266.67	200	166.67	166,67
300	280	266.67	266.67	250	233.33	220	200	175		1			316.67	350	333.33	250	266.67
220	200.33	180	166.67	120	109.33	75	29.99	20		1	250	233.33	133.33	116.67	29'99	20	33.33
266.67	233.33	200	180	166.67	150	100	100	75	,	,			216	150	266.67	133.33	29.99
320	333.33	300	275	250	250	233.33	75	75		150	166.67	100	133.33	120	100	83.33	66.67
8	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100

Table 5. Trehalase enzyme % in the supernatant of the homogenated *S. littoralis* larvae as affected by insect growth regulators treatment.

								2 nd	2 nd instar									
Time	1			Aft	After 2 days								100 No. 100	After 5 days	days			
Conc.	fr.*	7. T.	4 J.	1/8 f.r.	1/16 f.r.	¹ / ₃₂ f.r.	1,64 f.r.	1/128 f.r.	1/256 f.r.	'I'J	½ f.r.	1/4 f.r.	1/8 f.r.	1/16 f.r.	1/32 f.r.	1/64 f.r.	¹ / ₁₂₈ f.r.	1/256 f.r.
Cascade	7.5	105	66.67	109.33	150	80	150	8	20			120	266.67	105	109.33	105	100	108
Atabron	91.66	80	09	29.99	200	250	75	20	29.99			275	200	375	166.67	233.33	200	175
Consult	175	06	75	66.67	80	72.33	33.33	20	40	200	225	270	150	109.33	105	120	100	100
Match	150	200	29.99	180	175	06	56	40	99			250	270	280	266.67	233.33	109.33	100
Mimic	333.33	300	350	266.67	250	266.67	109.33	105	20	266.67	290	300	270	300	270	320	250	200
Control	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
								44	4 th instar					5000				
				AR	After 2 days									After 5 days	days			
Cascade	200	133.33	166.67	105	109.33	109.33	100	100	100	,	·			166.67	233.33	100	250	200
Atabron	366.67	333.33	350	275	250	150	109.33	80	29.99					20	75	80	66.67	09
Consult	200	100	100	75	20	29.99	20	33.33	20	,	,	250	266.67	333.33	300	275	200	150
Match	220	200	166.67	200	106.67	96	66.67	75	20		,			275	200	330	300	333.33
Mimic	250	200	230	175	150	166.67	09	20	40		275	220	266.67	250	175	150	100	20
Control	100	100	100	100	100	100	100	100	100	100	100	100	18	100	100	100	100	100

Table 6. Invertase enzyme % in the supernatant of the homogenated S. Iittoralis larvae as affected by insect growth regulators treatment.

									2 nd instar	star								
Time				¥	After 2 days	Ş							500	After 5 days	skı	S4		
Conc. Treat.	f.r.*	7.5 f.r.	½. f.r.	1/8 f.r.	1/16 f.r.	1/32 f.r.	1/64 f.r.	1/ ₁₂₈ f.r.	¹ /256 f.r.	f.r.	½ f.r.	7.4 f.r.	¹ /8 f.r.	1/16 f.r.	¹ / ₃₂ f.r.	1/64 f.r.	1/ ₁₂₈ f.r.	¹ /256 f.r.
Cascade	200	166.67	150	120	200	109.33	100	100	100			105	109.33	120	100	100	100	100
Atabron	109.33	150	175	200	133.33	100	100	100	100		1	109.33	118	112	109.33	100	100	100
Consult	100	100	100	100	233.33	220	100	300	200	100	120	100	100	133,33	105	109.33	109.33	100
Match	109.33	109.33	100	100	100	100	100	100	100	a,	=	120	133,33	105	109.33	100	100	100
Mimic	333,33	333.33	275	200	220	100	100	166.67	175	300	200	100	166.67	120	120	133.33	100	100
Control	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
	CONTRACTOR ACCOUNTS							C C C C C C C C C C C C C C C C C C C	4 th instar	star								
				Ā	After 2 days	ñ								After 5 days	sys			
Cascade	275	300	250	300	200	175	150	120	100	ı	1			200	150	33.33	133.33	33.33
Atabron	400	375	300	350	200	200	250	150	100	п	-	ı	ť	266.67	200	233,33	150	166.67
Consult	350	300	300	300	200	250	275	200	100	æ	1	233.33	166.67	150	50	100	66.67	33.33
Match	300	275	200	200	250	150	175	175	100				-	75	133.33	150	200	166.67
Mimic	300	150	100	225	200	250	150	200	100	п	233.33	200	100	66.67	100	20	33.33	33.33
Control	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100

* f.r. = recommended field rate

It is clear that the 4th larvae showed increased trehalase activity after two days of treatment with the higher concentration followed by decreased activity with lower concentration. Atabron recorded the highest values (366.67 to 109.33 %), meanwhile Consult caused the lowest values (33.33 to 75.0 %). All the tested IGRs showed increased activity of trehalase except Atabron at all the tested concentrations and Mimic at the least concentration.

ii. Invertase Data in Table 6. clear that at two days after application of the 2nd instar larvae, the tested IGRs affected the enzyme in three patterns, i.e. Cascade, Atabron and Mimic induced greatly the enzyme activity at the higher concentrations being ineffective at lower ones. Meanwhile, Consult manifested an inverse effect. On the other hand, Match seemed to be ineffective.

After five days of treatment, Mimic at higher concentrations induced noticeable increase in invertase activity then it decreased with decreasing the concentrations to be normal at 1/128 f.r. Other IGRs caused slight increase with the higher concentrations and no effect with lower concentrations.

In case of 4th instar larvae, all the applied concentrations, but not the least tested IGRs increased greatly the activity of invertase enzyme two days post treatment for 4th instar larvae, Atabron followed closely by Consult recorded the highest increase of activity ranged between as 400 to 150 % and as 350 to 200 % as the control, respectively.

After five days of treatment, irregular enzyme activity patterns were noticed due to the IGRs application. However, Atabron increased the enzyme activity that ranged between as 150-266.67% as control.

iii. Amylase Data in Table 7 show the changes in activity of amylase enzyme after two days of treating 2^{nd} instar larvae with the tested IGRs. It is clear that Match at the higher five concentrations caused noticeable increase in the enzyme activity (as 166.67 - 266.67% as the control). The other four IGRs induced slight or no effect on the amylase activity.

Similarly, the amylase enzyme after five days of treatment showed slight or no response to the IGRs application. Regarding the effect of the tested IGRs to 4th larval instar on the activity of amylase after two days of treatment, it is clear that there was noticeable increase in the enzyme activity especially at higher concentrations of all tested IGRs.

On the other hand, the lower concentrations of Cascade, Consult and Match and all the concentrations of Mimic except f.r. caused sever reduction in amylase activity.

The alive larvae of the 4th instar larvae after five days of treatment showed increased amylase activity with the higher concentrations of Match, Consult and Cascade and nearly all the concentrations of Atabron and Mimic.

Discussing the data concerning, the activity of carbohydrate hydrolyzing enzymes (trehalase, invertase and amylase) revealed the following:

- 1- The 4th larval instar proved to more sensitive than the 2nd instar.
- 2- Amylase activity seemed to be more tolerant to the IGRs application whereas trehalase was the most sensitive.
- 3- Atabron proved to be the most potent IGR.

It is obvious that there are some factors affected the efficiency of the tested IGRs on carbohydrate hydrolyzing enzymes. For instance, the tested IGRs them selves has its specific effect therefore some compounds were found to be highly effective, others were less effective. Moreover, some IGRs compounds affected severely the activity of one enzyme and others less effective another enzyme such as Match.

In addition, the higher concentration used in Match, Mimic and Consult caused the highest effect, while the lower concentration caused the lowest effect in case of trehalase. Moreover, the treated instar as the 4th proved more susceptible than the 2nd instar larvae. Also, the time of evaluation affected the efficiency of the tested IGRs, for instance after two days of treatment most the IGRs increased the enzymes activity while after 5 days the activity was decreased. The role of these factors could explain the varied data obtained and those documented in the literature where some authors reported an increase and/or decrease others reported no effect. For instance, Abdel-Fattah *et al.* (1986) found that diflubenzuron and triflumuron decreased greatly the activity of trehalase, invertase and amylase enzymes.

On contrary, AL-Elimi and Eid (1998) found an increase in carbohydrate hydrolyzing enzymes activity after treating larvae of *S. littoralis* with different concentrations of the IGRs diffubenzuron and buprofezin.

Table 7. Amylase enzyme % in the supernatant of the homogenated 5. Ilitoralis larvae as affected by insect growth regulators treatment.

																_
		1/256 f.r.	100	100	100	100	100	100			100	100	109.33	100	330	100
		1/128 f.r.	100	103	100	100	100	108			100	100	133.33	100	275	100
		1/64 f.r.	109.33	100	105	166.67	112	. 001			100	100	266.67	250	300	100
	lays	1/32 f.r.	105	100	100	100	120	100		days	133.33	166.67	266.67	330	100	100
	After 5 days	1/16 f.r.	112	103	111.33	105	109.33	100		After 5 days	150	120	350	275	100	100
		1/8 f.r.	103	106.67	109	133.33	109.33	100			-		333.33	1	166.67	100
		1/4 f.r.	111.33	105	106.67	109.33	100	100				-	330	-	200	100
		72 f.r.	ı.	1	100		112	100			5	1			225	100
		f.r.		1	112	105	í.	100						•	-	100
2 nd instar		1/256 f.r.	100	100	100	100	100	100	4 th instar		33.33	20	22	20	25	100
2 nd ii		¹ / ₁₂₈ f.r.	100	100	120	100	100	100	₽.		20	200	20	33.33	20	100
		1/64 f.r.	100	100	100	100	109.33	100			20	150	80	29.99	33.33	100
	After 2 days	¹ /32 f.r.	111.67	100	109.33	105	100	100			09	20	29'99	75	40	001
		1/16 f.r.	103.33	109	112	166.67	105	108		After 2 days	75	330	75	20	20	100
		1/8 f.r.	100	109.33	100	175	112	100			66.67	333.33	06	99	80	100
		7.4 f.r.	105	120	100	200	100	100			80	250	133.33	166.67	29.99	100
		% f.r.	109.33	105	111.66	250	111.33	100			120	300	109.33	133.33	09	100
		f.r.*	100	103.33	106.33	79.997	100	100			133.33	333.33	200	150	100	100
E.	Time	Conc. Treat.	Cascade	Atabron	Consult	Match	Mimic	Control			Cascade	Atabron	Consult	Match	Mimic	Control

* f.r. = recommended field rate

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دراسات سمية وبيوكيميائية على تأثير بعض منظمات النمو الحشرية على دودة ورق القطن

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درس تأثير خمسة منظمات نمو حشرية هى كاسكيد، أتابرون، كونسلت، ماتش ومميك على السمية وبعض القياسات (الأنشطة) البيوكيميائية للعمرين الثاني والرابع لدودة ورق القطن على فترات مخففة من مقد عنيت اليرقات الحديثة الانسلاخ على أوراق خروع معاملة بتسعة تركيزات مخففة من التركيز الحقلى وقد أشارت النتائج إلى:

ظهر مركب أتابرون كأكثر منظمات النمو الحشرية فعالية بينما كان مركب مميك أقلها تأثيراً ضد كــل مــن العمر اليرقى الثاني والرابع، وقد أظهر العمر اليرقى الرابع حساسية أكبر لكل المركبات المختبرة عن العمر اليرقى الثاني عند كل التركيزات المختلفة.

- زادت منظمات النمو الحشرية محتوى البروتين الكلى الذائب بعد يومين من معاملة العمرين البرقيين الثاني والرابع عدا كاسكيد عند التركيزات الأقل، وقد سجل انخفاضاً ملحوظا بعد خمسة أيام من معاملة العمرين البرقيين الثاني والرابع بكل من كونسات، ماتش وكاسكيد.
- خفضت المركبات المختبرة نشاط إنزيم GOT بينما زاد نشاط إنزيم GPT بعد يومين من
 معاملة العمر اليرقى الثاني وكان العكس صحيحا بعد خمسة أيام من المعاملة، وفي حالة
 العمر الرابع زادت منظمات النمو نشاط كلا الإنزيمين بعد ٢و٥ يوم من المعاملة.
- زيادة نشاط إنزيمات تريهاليز، انفرتيز وأميليز في العمرين اليرقيين الثاني والرابع بعد ٢و
 و يوم من المعاملة ما عدا مركبي كونسلت ومميك في حالة إنزيم الانفرتيز في يرقات العمر الرابع.