

EFFECT OF GIBERELLIC ACID AND STIMUFOL FERTILIZER ON GROWTH AND FLOWERING OF Hemerocallis aurantiaca PLANT.

Journal

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ABSTRACT

A two-year trial was carried out at the Experimental Farm of Horticulture and the Laboratory of Agricultural Botany Dept., Fac. of Agric., Benha Univ., during 2005/2006 and 2006/2007 seasons, to study the effect of sixteen treatments represented the combination between four concentrations of GA₃ (0.0, 50, 100 and 200 ppm.) and four levels of Stimufol fertilizer (0.0, 2, 4 and 6 g/L) on the vegetative growth, flowering, anatomical measurements and chemical composition of *Hemerocallis aurantiaca* plant.

GA₃ treatments positively affected vegetative growth parameters (number of leaves/clump, fresh and dry weights of leaves/clump and number of offshoots/clump), flowering growth parameters (number of days from planting to start flowering, number of flowers/clump, length and diameter of flower stalk, fresh and dry weights of flower, duration of flower on plant and flower vase life) and chemical composition parameters (leaf total carbohydrates, N, P and K contents) with superiority to the medium and high rates. Also, all levels of Stimufol fertilizer significantly improved all the aforementioned parameters, especially the medium and high levels. Additionally, most the combinations of GA₃ concentrations and Stimufol fertilizer levels enhanced all the studied parameters, particularly the combinations of GA₃ at 100 or 200 ppm with Stimufol fertilizer at 4 or 6 g/L in both seasons.

As for the anatomical studies, several alternations were existed with different applied treatments. Most of the combinations between GA3 concentrations and levels of Stimufol fertilizer showed the highest values in transfer sections of leaf blade and flower stalk

compared with the control. The most effective treatment was GA₃ at 200 ppm provided with Stimufol fertilizer at 6 g/L.

Conclusively, it is preferable to spray *Hemerocallis aurantiaca* plant three times at three weeks interval with GA₃ at 200 ppm supplemented with Stimufol at 6 g/L to obtain the best vegetative and flowering characteristics with high quality.

Key word: *Hemerocallis aurantiaca*, GA3, Stimufol fertilizer, vegetative growth, flowering, anatomy and chemical composition.

INTRODUCTION

The flowering bulbous plants are considered as a group of the most beautiful adjuncts for garden decoration. They are used particularly in landscape, production of commercial cut-flowers and act as a source of glorious colors and perfumes. *Hemerocallis aurantiaca*, Baker (Hemero, a day; kallos, beauty; in reference to the flowers being fresh only for only day or so) belongs closely to Fam. Asphodlaceae not to Fam. Liliaceae, is a very short rhizome with numerous more or less tuberous and fleshy roots, native to Japan and commonly known as golden summer daylily or orange daylily. It is well known for its hardiness and showy blooms produced during spring and summer (Oganejova, 1990).

Hemerocallis is an indispensable choice for woodland and landscape gardens, when grown in the background or herbaceous borders, in front of a shrubbery or a long the sides of ponds or streams, a highly delightful effect is produced. Daylily is also used as a cut flowers. (Bose and Yadav, 1989). Most bulbs need more than two applications of fertilizers during the growing season, but the most important point is that the greatest increase in size and weight of the new developing bulb takes place in the period during and mostly after flowering, as long as the foliage remains in good condition. Thus, fertilization must continue for good vegetative growth to produce a good flower and large new mature bulbs (Rees, 1992). In this concern, Manoly (1996) and Naglaa and Kandeel (2001) they demonstrated that spraying iris plants with foliar fertilizers (containing macro and micro elements) produced the best results for flowering parameters. In addition, Abou-Dahab (1996) reported that most of foliar commercial fertilizers used (Irral, All-Grow, Kristalon, Stimofort and Sangoral)

improved the growth of schefflera plants. Shahin (1998) claimed that spraying Hemerocallis aurantioca plants with greenzit (foliar fertilizer containing macro and micro elements) at the rates of 1, 3 or 5cm³ per liter caused a considerable increase in vegetative and flowering growth as well as leaf chemical composition determinations. Plant growth and development are greatly influenced by chemical growth substances. The stimulatory effect of gibberellins application on growth and flowering of the different ornamental plants has been indicated by several researchers. In this concern, Singh et al., (1994) on Dahlia variables, Naglaa and Kandeel (2001) on iris and Youssof (2004) on Strelitzia reginae plants found that spraying with GA₃ at 100 or 200 ppm improved vegetative growth parameters (number of leaves and offsets, plant height, fresh and dry weights of leaves) as well as flowering growth parameters (number, length, fresh and dry weights of flowers/plant). In addition, Abou El-Ella (2007) showed that spraying Acanthus mollis plant with GA₃ enhanced vegetative and flowering growth measurements.

The present work is an attempt to enhance the production of this plant, to prolong the flowering period and improve the quality and quantity of flowers, especially during summer months.

MATERIALS AND METHODS

This investigation was carried out in the open field at the Floriculture Nursery of the Horticulture Department and Anatomical Laboratory of Th Agriculture Botany Dept., Faculty of Agriculture at Moshtohor, Benha University, Kalubia Governorate, during the seasons of 2005/2006 and 2006/2007 to study the ability of GA₃ treatments (0, 50, 100 and 200 ppm) and Stimufol fertilizer treatments (0, 2, 4 and 6 g/L water) in combining treatments to lump their benefits in extending the flowering season provided with high flower quality, as well as increasing the plant productivity of the new hemerocallis offshoots.

Plant material:

Divisions of *Hemerocallis aurantiaca*, Baker clumps "local variety" at the weight of 75-80 g were obtained freshly from Floriculture Nursery of the Horticulture Department, Faculty of Agriculture at Moshtohor.

Planting procedure:

Divisions of hemerocallis clumps were planted in a clay loam soil "the analyses of the used soil are presented in Tables (a & b)" in mid September in beds 1x1m² as each bed contain 4 divisions planted at 50x50 cm in between in both seasons.

Table (a): Mechanical analysis of the experimental soil

D	T I : 4	Seas	ons
Parameters	Unit	2005/2006	2006/2007
Coarse sand	%	4.93	5.15
Fine sand	%	16.41	15.57
Silt	%	27.33	26.98
Clay	%	51.33	52.30
Textural class		Clay loam	Clay loam

Table (b): Chemical analysis of the experimental soil.

Parameters	Unit	Sea	sons
rarameters	Ont	2005/2006	2006/2007
Caco ₃	%	2.21	2.32
Organic matter	%	2.41	2.39
Available nitrogen	%	0.94	0.92
Available phosphorus	%	0.45	0.49
Available potassium	%	0.79	0.81
EC	ds/m	1.21	1.33
рН		7.73	7.77

Stimufol fertilizer treatments:

Hemerocallis plants received Stimufol fertilizer (as a complete foliar fertilizer containing macro and micro nutrients, Table c) at the rates of 0, 2, 4 and 6 g/L as a foliar spray three times a year after 50, 65 and 80 days from planting in both seasons.

Table (c): Chemical composition of Stimufol fertilizer (%)

Total N	P	K	Mg	В	Fe	Zn	Mn	Cu	Co	Mo	Amino acids
25	16	12	0.20	0.044	0.17	0.03	0.08	0.08	0.01	0.001	0.20

Giberellic acid treatments:

Hemerocallis plants were subjected to foliar spray with GA_3 at the rates of 0, 50, 100 and 200 ppm three times, each at two weeks interval, the first one was after 55 days from planting in both seasons.

Layout of the Experiment:

The design of this experiment was a split plot design with 16 treatments represented the combinations between Stimufol fertilizer at the rates of 0, 2, 4 and 6 g/L and GA₃ at the rates of 0, 50, 100 and 200 ppm (4 Stimufol fertilizer levels x 4 GA₃ concentrations) replicated three times (each replicate consisted of three beds, with four plants/bed). GA₃ treatments were employed to the main plot, while Stimufol fertilizer treatments were assigned to the sub plot.

The following measurements were recorded:

1- Vegetative growth parameters:

Number of leaves/clump, number of offshoots/clump and fresh and dry weights of leaves/ clump.

2-Flowering parameters:

Flowering start (days), number of flowers/clump, length and diameter of flower stalk, fresh and dry weights of flower (g), duration of flower on plant (days) and flower vase life (days).

3- Chemical composition determinations:

-Total nitrogen percentage was determined in the dried leaves by using wet digestion according to Piper (1947), using micro-kjeldahl method as described by Horneck and Miller (1998).

-Phosphorus was determined calorimetrically in spectronic (20) spectrophotometer using the method described by sandell (1950).

-Potassium content was determined by flame photometer according to Horneck and Hanson (1998).

-Total carbohydrates content was determined in dry leaf powder according to Herbert *et al.*, (1971).

All samples of chemical analyses were taken at the flowering start.

4-Anatomical studies

The samples of leaf were taken from the internodes lower the inflorescence from top and samples from flowering stalk were taken from the main inflorescence. The samples were taken from effected treatments added with the control. The specimens were taken then killed and fixed in FAA (5ml. formalin, 5ml. glacial acetic acid and 90ml. ethyl alchohol 70%), washed in 50% ethyl alcohol, dehydrated in series of ethyl alchohols 70,90,95 and 100%, infiltrated in xylene, embedded in paraffin wax with a melting point of 60-63°C, sectioned to 20 microns in thickness (Sass 1951), stained with the double stain method (fast green and safranin), cleared in xylene and mounted in Canada balsam (Johanson, 1940). Sections were read to detect histological manifestation of noticeable responses resulted from other treatments.

Statistical analysis:

All data obtained in both seasons of study were subjected to analysis of variance as factorial experiments in split plot design. L.S.D. method was used to differentiate means according to Snedecor and Cochran (1989).

RESULTS AND DISCUSSION

Effect of GA₃ and Stimufol:-

I- Vegetative growth measurements:

I-1- Number of leaves and offshoots/ clump:

Data in Table (1) indicate that all GA₃ concentrations significantly increased the number of leaves and offshoots/clump when compared with untreated plants "control" in both seasons. In this respect, the medium concentration of GA₃ (100ppm) gave the highest number of leaves and offshoots/ clump in both seasons. On the other hand, there was a positive correlation between the number of leaves

Table(1): Effect of GA₃ and Stimufol fertilizer treatments on number of leaves and offshoots/clump of Hemerocallis aurantiaca plants (2005/2006 and 2006/2007 seasons).

F	arameters		Numbe	r of leaves	/ clump			Number	of offshoo	ots/ clump)
			(GA ₃				G	A ₃		
Treatments		0.0	50 ppm	100 ppm	200 ppm	Mean	0.0	50 ppm	100 ppm	200 ppm	Mean
						First s	eason		·		
	0.0	15.3	16.8	18.6	18.1	17.2	2.83	3.11	3.51	3.21	3.17
Stimufol fertilizer	2g/L	15.9	17.2	19.2	18.6	17.7	3.01	3.41	3.81	3.19	3.36
Stimuloi lei tinzei	4g/L	16.7	19.7	22.6	21.9	20.2	3.21	3.92	4.51	3.92	3.89
	6g/L	17.2	21.2	23.8	20.7	20.7	3.18	3.82	4.63	4.18	4.00
Mean		16.3	18.7	21.1	19.8		3.1	3.57	4.12	4.00	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer trea Interaction (GA ₃ X fertil				2.13 ² 1.86 2.64					0.43 0.36 0.62		
						Second	season				
	0.0	16.2	18.3	19.7	18.0	18.1	2.69	3.28	3.93	3.23	3.28
Stimufol fertilizer	2g/L	17.8	20.2	20.3	19.3	19.4	3.21	3.64	4.21	3.52	3.65
~	4g/L	18.5	19.8	23.6	20.2	20.5	3.64	4.21	4.81	3.98	4.16
	6g/L	19.7	22.1	24.2	21.7	21.9	3.72	4.26	4.93	4.19	4.28
Mean		18.1	20.1	22.0	19.8		3.32	3.85	4.47	3.73	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer treat Interaction (GA ₃ X fertil				1.98 1.92 2.83					0.44 0.41 0.70		

and offshoots/ clump and Stimufol fertilizer levels, so the number of leaves and offshoots were increased as the level of Stimufol fertilizer increased. In this concern, spraying the plants with high Stimufol level (6g/L) statistically induced the highest number of leaves and offshoots/ clump in both seasons.

Moreover, data in Table (1) show that all the interactions between GA₃ concentrations and Stimufol fertilizer levels statistically increased the number of leaves and offshoots/ clump as compared with control in both seasons. However, the combined treatment between GA₃ at 100ppm and Stimufol fertilizer at 6g/L significantly produced the highest number of leaves and offshoots/ clump as compared with the other combinations or control in both seasons.

I-2- Fresh and dry weights of leaves/ clump (g)

It is interest to note from Table (2) that there was a positive relationship between the fresh and dry weights of leaves and GA₃ and Stimufol fertilizer treatments. Since, as the levels of GA₃ or Stimufol fertilizer were increased, the fresh and dry weights of leaves increased until reach to the maximum increases at the high level. This trend was true in both seasons.

In addition, all interactions between GA_3 concentrations and Stimufol fertilizer levels caused remarkable increments in the fresh and dry weights of leaves in both seasons. However, the heaviest fresh and dry weights of leaves were obtained by using the combined treatment between GA_3 at 200ppm and Stimufol fertilizer at 6 g/L in both seasons.

The aforementioned results of GA₃ regarding vegetative growth measurements may be due to the fact that GA₃ may cause cell elongation by the induction of enzymes that weaken the cell walls. Also, the mechanism by which gibberellins might stimulate cell elongation is that the hydrolysis of starch resulting from the production of GA₃ induced α-amylase which might increase the concentration of sugars, thus raising the osmotic pressure in the cell sap so that water enters the cell and tends to stretch it (Macleod and Millar, 1962). The aforementioned results of vegetative growth measurements i.e. number of leaves and offshoots / clump and fresh and dry weights of leaves/clump as affected by GA₃ are in parallel with those of Singh *et al.*, (1994) on dahlia, Preeti *et al.*, (1997) and Ved *et al.*, (1998) on tuberose Dantuluri *et al.*, (2002) on *Lilium*

Table(2): Effect of GA₃ and Stimufol fertilizer treatments on fresh and dry weights of leaves/clump of Hemerocallis aurantiaca plants (2005/2006 and 2006/2007 seasons).

]	Parameters	F	resh weig	ht of leave	s/clump (g	g)	I	ry weigh	t of leave	s/clump (g)	
				A ₃				G	A ₃			
Treatments		0.0	50 ppm	100 ppm	200 ppm	Mean	0.0	50 ppm	100 ppm	200 ppm	Mean	
,				· · · · · · · · · · · · · · · · · · ·		First s	eason		· · · · · · · · · · · · · · · · · · ·		<u> </u>	
	0.0	166.7	189.8	225.0	237.1	204.7	21.3	24.4	29.2	30.6	26.4	
Stimufol fertilizer	2g/L	174.9	194.3	236.2	249.2	213.7	21.8	24.4	30.2	31.4	27.0	
Summer of the control	4g/L	187.0	234.4	293.9	297.8	253.3	22.8	28.8	37.0	36.9	31.4	
	6g/L	206.4	267.2	285.6	310.2	267.4	22.7	32.0	36.2	36.9	32.0	
Mean		183.8	221.4	260.2	273.6		22.2	27.4	33.2	34.0		
LSD at 5 % for GA ₃ treatments Stimufol fertilizer trea Interaction (GA ₃ X ferti				32.6 45.2 57.6	;				5.13 3.93 5.96			
			Second season									
	0.0	179.8	223.2	240.3	237.6	220.2	24.9	28.7	30.9	31.6	29.0	
Stimufol fertilizer	2g/L	213.6	232.3	255.7	279.8	245.4	28.8	29.0	31.9	33.5	30.8	
Simuloi ici inizci	4g/L	205.3	257.4	304.4	272,7	260.0	27.5	31.6	37.4	35.4	33.0	
	6g/L	221.2	282.8	314.6	323.3	285.5	28.7	33.9	37.7	40.1	35.1	
Mean		205.0	248.9	278.8	278.4		27.5	30.8	34.5	35.2		
LSD at 5 % for GA3 treatments Stimufol fertilizer trea Interaction (GA3 X ferti	1			30.2 33.0 50.7					3.18 2.80 5.14			

moculatum and Gomaa (2003) who mentioned that using GA₃ at 50, 100 and 150 ppm increased the plant height, stem diameter, number of branches and foliage fresh weight of Dahlia pinnata plants. Whereas, the results of Stimufol fertilizer are in parallel with those obtained by Manoly (1996) on Iris, Shahin (1998) on Hemerocallis aurantiaca, who mentioned that spraying the plants with greenzit (foliar fertilizer containing macro and micro elements) at the rates of 1, 3 or 5cm³ per liter increased the number of leaves and offshoots /plant and fresh and dry weights of leaves, Atta-Alla and Zaghloul (2002) and Youssef and Goma (2007) on Iris tingitana and Abou-El-Ella (2007) showed that spraying Acanthus mollis plants with Kristalon at 2, 3 or 4 g/L and New-Star fertilizer at 3, 4 or 5 g/L increased plant height, number of leaves and fresh and dry weights of leaves.

II- Flowering growth measurements:

II-1. Flowering start "days"

Data in Table (3) indicate that number of days to start flowering was decreased as the concentration of GA₃ increased in both seasons. So, 200 ppm GA₃- sprayed plants significantly induced earlier flowering after 250 and 249 days as compared with 260 and 278 days for control from planting in the first and second seasons, respectively. In addition, all levels of Stimufol fertilizer led to early flowering of this parameter, especially the medium level (4g/L). On the other hand, all combinations between GA₃ concentrations and Stimufol fertilizer levels induced a remarkable precocity in this parameter, especially the combinations of GA₃ at 200 ppm in both seasons. However, the earliest flowering was recorded by the combined treatment between GA₃ at 200 ppm and Stimufol fertilizer at 4g/L as it recorded 246 and 242 days in the first and second seasons, respectively.

II-2. Number of flowers/clump.

Data in Table (3) show that all GA₃ treatments statistically increased the number of flowers/ clump as compared with control in both seasons. In this concern, the highest number of flowers/clump was obtained by medium GA₃ concentration as it gave 4.69 and 5.18 flowers/ clump in the first and second seasons, respectively.

On the other side, there was a positive relationship between the number of flowers/ clump and Stimufol fertilizer levels, hence the number of flower/clump increased as the level of Stimufol fertilizer increased. Thereupon, 6g/L Stimufol fertilizer- sprayed plants showed

Table(3): Effect of GA₃ and Stimufol fertilizer treatments on flowering start and number of flowers/clump of Hemerocallis aurantiaca plants (2005/2006 and 2006/2007 seasons).

	Parameters		Flowe	ring start	(days)			Number	r of flowe	rs/clump	
				A ₃				G	A ₃		
Treatments		0.0	50 ppm	100 ppm	200 ppm	Mean	0.0	50 ppm	100 ppm	200 ppm	Mean
				All the state of t		First s	eason				
	0.0	264	261	258	254	259	3.02	3.41	3.92	3.87	3.56
Stimufol fertilizer	2g/L	258	254	252	251	254	3.41	3.86	4.83	3.93	4.01
Sumuloi ierunzer	4g/L	255	252	249	246	251	3.81	4.13	4.79	4.28	4.25
•	6g/L	262	259	258	249	257	3.96	4.63	5.21	5.18	4.75
Mean		260	257	254	250		3.55	4.01	4.69	4.32	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer trea Interaction (GA ₃ X ferti				2.38 3.21 5.97					0.56 0.63 0.86		
						Second	season				
	0.0	261	259	256	252	257	3.21	3.82	4.32	4.35	3.93
Stimufol fertilizer	2g/L	257	256	253	250	254	3.93	4.15	4.82	4.39	4.32
Stimulor for thizer	4g/L	253	252	251	242	250	4.50	4.11	5.67	5.37	4.91
	6g/L	259	258	256	251	256	4.39	4.60	5.92	5.82	5.18
Mean		278	256	254	249		4.01	4.17	5.18	4.98	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer trea Interaction (GA ₃ X ferti				2.38 3.21 5.97		_			0.65 0.72 0.91		

to be the most effective level for producing the highest number of flowers/ clump as it gave 4.75 and 5.18 flowers/ clump in the first and second seasons, respectively. Moreover, all combinations between GA₃ concentrations and Stimufol fertilizer levels succeeded in increasing the number of flowers/ clump in both seasons. However, the highest number of flowers/ clump was registered by using the combined treatment between GA₃ at 100ppm and Stimufol fertilizer at 6g/L, followed descendingly by the combined treatment between GA₃ at 200ppm and Stimufol fertilizer at 6g/L in both seasons. The differences between the aforementioned two combined treatments were non significant in both seasons from the statistical standpoint.

II-3. Length and diameter of flower stalk (cm).

Data in Table (4) show that flower stalk length was increased with increasing GA₃ concentration. On the opposite, flower stalk diameter was decreased with increasing GA₃ concentration in both seasons. Besides, all studied Stimufol fertilizer levels statistically increased the length and diameter of flower stalk with superior for the high level in both seasons. However, the longest flower stalk was scored by using the combined treatment between GA₃ at 200 ppm and Stimufol fertilizer at 6 g/L, whereas the thickest flower stalk was registered by the combined treatment between GA₃ at 50 ppm and Stimufol fertilizer at 6 g/L in both seasons.

II-4. Fresh and dry weights of flowers (g):

Data in Table (5) reveal that all GA₃, Stimufol fertilizer treatments and their combinations increased the fresh and dry weights of flower in both seasons. In this concern, the increments in fresh and dry weights of flower were in parallel with the applied concentrations of GA₃ or Stimufol fertilizer, so the highest concentration of GA₃ (200 ppm) or the highest level of Stimufol fertilizer significantly scored the heaviest fresh and dry weights of flower, with the exception of dry weight of flower in the second season as medium concentration of GA₃ (100 ppm) surpassed the other concentrations. However, the heaviest fresh and dry weights of flower were obtained by using the combined treatment between GA₃ at 200 ppm and Stimufol fertilizer at 6 g/L in both seasons.

Table(4): Effect of GA₃ and Stimufol fertilizer treatments on length and diameter of flower stalk of *Hemerocallis* aurantiaca plants (2005/2006 and 2006/2007 seasons).

P	arameters		Flower	stalk leng	gth(cm)			Flower s	talk diam	eter (cm)	
				A ₃				G	A ₃		
Treatments		0.0	50 ppm	100 ppm	200 ppm	Mean	0.0	50 ppm	100 ppm	200 ppm	Mean
		Para translat Section and American		<u></u>		First s	eason				
	0.0	118.2	121.6	125.9	129.3	123.8	1.38	1.43	1.41	1.39	1.40
Stimufol fertilizer	2g/L	120.5	123.4	128.3	132.5	126.2	1.42	1.45	1.42	1.43	1.43
Simuloi lerinizer	4g/L	123.9	126.2	132.1	136.0	130.0	1.56	1.63	1.52	1.42	1.53
	6g/L	125.0	128.9	131.8	138.4	131.0	1.63	1.76	1.68	1.57	1.66
Mean		121.9	125.0	129.5	134.1		1.50	1.57	1.51	1.45	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer treat Interaction (GA ₃ X fertil			A. 43	7.32 6.21 12.41					N.S 0.12 0.15		
						Second	season				
	0.0	115.9	118.9	122.6	126.0	120.9	1.42	1.47	1.45	1.42	1.44
Stimufol fertilizer	2g/L	118.3	120.6	125.7	129.3	123.5	1.51	1.59	1.49	1.47	1.52
OMEGAVE IN PRINCE	4g/L	120.6	125.9	127.9	133.7	127.0	1.68	1.73	1.56	1.50	1.62
	6g/L	122.0	125.3	128.2	134.8	127.6	1.72	1.82	1.63	1.61	1.70
Mean		119.2	122.7	126.1	131.0		1.58	1.65	1.53	1.50	
LSD at 5 % for GA3 treatments Stimufol fertilizer treat Interaction (GA3 X fertili				6.11 5.26 9.87	,				N.S 0.16 0.19		-

Table(5): Effect of GA₃ and Stimufol fertilizer treatments on fresh and dry weights of flower of Hemerocallis aurantiaca plants (2005/2006 and 2006/2007 seasons).

Pa	rameters		Fresh w	eight of fl	ower (g)			Dry we	eight of flo	ower (g)	
				A ₃				G	A3		
Treatments		0.0	50 ppm	100 ppm	200 ppm	Mean	0.0	50 ppm	100 ppm	200 ppm	Mean
						First s	eason			· · · · · · · · · · · · · · · · · · ·	
	0.0	68.2	71.3	73.6	75.4	72.1	8.11	8.41	8.53	8.59	8.41
Stimufol fertilizer	2g/L	71.9	73.5	75.3	76.9	74.4	8.48	8.59	8.65	8.68	8.60
Simuloi letinizer	4g/L	73.7	76.8	77.1	78.7	76.6	8.62	8.90	8.78	8.81	8.77
	6g/L	75.2	75.9	78.5	81.6	77.8	8.64	8.85	8.87	9.00	8.84
Mean		72.3	74.4	76.1	78.2		8.46	8.68	8.70	8.77	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer treatmenter action (GA ₃ X fertilizer treatments)				3.26 3.64 5.28					N.S 0.31 0.52		
						Second	season				
	0.0	71.8	73.2	76.0	78.4	74.9	8.68	8.85	9.12	9.00	8.91
Stimufol fertilizer	2g/L	73.6	75.6	81.9	81.2	78.1	8.97	9.07	9.66	9.33	9.26
Stimuloi lei tilizei	4g/L	76.9	78.3	80.7	83.4	79.8	9.15	9.31	9.36	9.50	9.33
	6g/L	77.3	78.1	82.3	85.2	80.7	9.12	9.39	9.38	9.62	9.38
Mean		74.9	76.3	80.2	82.1		8.98	9.16	9.38	9.36	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer treatm Interaction (GA ₃ X fertiliz				4.25 4.87 6.09					N.S 0.37 0.61	-	

II-5.Duration of flower on plant and flower vase life "days":

Data in Table (6) indicate that all tested GA₃ concentrations slightly increased duration of flower on plant and flower vase life, especially the high concentration in both seasons. Likewise, the three studied levels of Stimufol fertilizer resulted in slightly increments in these parameters. Meanwhile, 6g/L Stimufol fertilizer-sprayed plants showed to be the most effective level to induce the highest duration of flower on plant and flower vase life as compared with other levels or control in both seasons. Generally, all combinations between GA₃ and Stimufol fertilizer treatments prolonged the duration of flower on plant and flower vase life "days" in both seasons as compared with control. However, the highest values of flower duration on plant and flower vase life were scored by using the combined treatment between GA₃ at 200 ppm and Stimufol fertilizer at 6 g/L when compared with control and other combinations in both seasons.

The principal role of GA₃ upon flowering stimulation in many plants has been recommended to be manly due to its effect upon increasing the length of flower stalk (bolting), the process that strictly must proceeded flowering appearance process (Devlin and Witham, 1983). Moreover, the effects of GA₃ on flower growth measurements may be due to the fact that gibberellins play a role in flowering, probably it is further elaborated into florigen by the plant. Hence, gibberellins can not be the same substance as florigen but at least it may act as its precursor. The propounder of (Florigen concept) florigen but made up of two substances, namely gibberellins and anthesins. The latter are considered to be nitrogen rich compounds (Macleod and Millar, 1962).

The aforementioned results of flowering growth measurements i.e., flowering start, number of flowers, length and diameter of flower stalk, fresh and dry weights of flower, duration of flower on plant and flower vase life are in conformity with those obtained by Dantuluri et al., (2002) and Tiwari and Singh (2002) on Lilium maculatum, Wankhede et al., (2002) on Polianthes tuberosa, Gomaa (2003) on Dahlia pinnata and Youssef (2004) who mentioned that treated Strelitzia reginae plants with GA₃ at 100 or 200 ppm increased number of flowers/plant, length and diameter of flower, diameter of flower (cm), fresh weight of flower/plant, duration of flower on plant and vase life of flower as well as inducing early flowering.

Table(6): Effect of GA₃ and Stimufol fertilizer treatments on duration of flower on plant and flower vase life of *Hemerocallis aurantiaca* plants (2005/2006 and 2006/2007 seasons).

	Parameters	D	uration of	flower on	plant(day	ys)		Flowe	r vase life	(days)	
			(A ₃				G	A ₃		
Treatments		0.0	50 ppm	100 ppm	200 ppm	Mean	0.0	50 ppm	100 ppm	200 ppm	Mean
						First s	eason				
	0.0	11.2	11.7	11.9	12.3	11.8	6.81	7.03	7.80	7.73	7.34
Stimufol fertilizer	2g/L	11.9	12.3	12.3	13.2	12.4	6.92	7.61	8.21	8.01	7.69
Stimuloi lertilizer	4g/L	12.8	13.8	12.9	13.7	13.3	7.23	7.92	8.63	8.93	8.18
	6g/L	13.2	13.7	13.8	14.2	13.7	7.96	8.30	8.92	9.21	8.60
Mean		12.3	12.9	12.7	13.4		7.23	7.72	8.39	8.47	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer tre Interaction (GA ₃ X fert	1			0.93 1.21 1.34	· · · · · · · · · · · · · · · · · · ·				0.83 0.76 1.14		
						Second	season				
	0.0	11.6	12.0	12.9	12.7	12.3	6.98	7.14	7.95	8.03	7.53
Stimufol fertilizer	2g/L	12.2	12.3	13.0	12.9	12.6	7.23	7.63	8.17	8.26	7.82
Stimulor fer thizer	4g/L	13.9	13.6	13.6	13.5	13.7	7.99	8.02	8.73	8.90	8.41
	6g/L	13.7	13.9	14.1	14.3	14.0	7.83	8.19	9.10	9.36	8.62
Mean		12.9	13.0	13.4	13.4		7.51	7.75	8.49	8.64	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer treation (GA ₃ X fert				0.48 0.96 1.21					0.86 0.82 1.19		

The aforementioned results of Stimufol fertilizer are coincided with those obtained by Barman and Pal (1993) on *Polianthes tuberosa*, Mukherjee *et al.*, (1994) on gladiolus, Singh and Uma (1996) on *Polianthes tuberose* and Shahin (1998) on *Hemerocallis aurantiaca*, showed that spraying the plants with greenzit (foliar fertilizer containing macro and micro elements) at the rates of 1, 3 or 5cm³ per liter improved flowering start, number of flowers, length and diameter of flower stalk, fresh and dry weights of flower and flower vase life. Youssef (2004) stated that treating *Strelitzia reginae* plants with Stimufol fertilizer at 4 or 6 g/L enhanced flowering growth parameters i.e., number of days to start flowering "flowering date", length, diameter, fresh and dry weights of flowering stalk.

III-Chemical composition determinations:

Data in Tables (7&8) reveal that all tested GA₃ concentrations slightly increased leaf N, P, K and total carbohydrates contents with non-significant increments in most cases. Regarding Stimufol fertilizer treatments, data showed that leaf N, P, K and total carbohydrates contents of hemerocallis plants increased with increasing Stimufol fertilizer level. Hence, 6g/L Stimufol-sprayed plants gave the highest values in this concern.

Moreover, all combinations between GA₃ concentrations and Stimufol fertilizer levels statistically increased leaf N, P, K and total carbohydrates content in both seasons. However, the highest values of these parameters were recorded by using the combinations of Stimufol fertilizer at the highest level in both seasons.

The stimulated effect of Stimufol fertilizer may be due to the role of Stimufol on supplying the plants with their required nutrients for more carbohydrates and proteins production which are necessary for vegetative, flowering, bulbs growth and chemical composition (Marschner, 1995).

The aforementioned results of chemical composition measurements i.e., leaf N, P, K and total carbohydrates contents concerning Stimufol fertilizer are coincided with those obtained by Shahin (1998) on *Hemerocallis aurantiaca*, demonstrated that spraying the plants with greenzit (foliar fertilizer containing macro and micro elements) at the rates of 1, 3 or 5 cm³ per liter increased leaf N, P, K and total carbohydrates contents, Naglaa and Kandeel (2001), Ataa-Alla and Zaghloul (2002) on *Iris tingitana*, Youssef

Table(7): Effect of GA₃ and Stimufol fertilizer treatments on leaf nitrogen and phosphorus content (%) of Hemerocallis aurantiaca plants (2005/2006 and 2006/2007 seasons).

Pa	rameters			N%					P%			
	į			A ₃				G	A ₃			
Treatments		0.0	50 ppm	100 ppm	200 ppm	Mean	0.0	50 ppm	100 ppm	200 ppm	Mean	
						First s	eason			***************************************		
	0.0	2.63	2.84	3.02	3.26	2.94	0.123	0.132	0.143	0.143	0.135	
Stimufol fertilizer	2g/L	2.96	2.98	3.60	3.46	3.25	0.142	0.153	0.148	0.169	0.153	
Sumuloi ierunzer	4g/L	3.83	3.79	3.92	4.32	4.00	0.146	0.162	0.169	0.166	0.158	
	6g/L	3.92	4.23	4.52	4.27	4.24	0.151	0.167	0.173	0.183	0.169	
Mean		3.34	3.46	3.77	3.83		0.141	0.154	0.158	0.165		
LSD at 5 % for GA ₃ treatments Stimufol fertilizer treatm Interaction (GA ₃ X fertilize	1			N.S 0.83 0.92					N.S 0.018 0.023			
		Second season										
	0.0	2.90	3.24	3.41	3.38	3.23	0.136	0.142	0.152	0.152	0.146	
Stimufol fertilizer	2g/L	3.36	3.94	3.92	3.79	3.75	0.142	0.146	0.157	0.163	0.152	
Stimuloi ici tilizci	4g/L	3.97	3.87	4.36	4.11	4.01	0.167	0.153	0.169	0.179	0.167	
	6g/L	4.02	4.13	4.52	4.23	4.23	0.158	0.167	0.158	0.186	0.167	
Mean		3.60	3.80	4.05	3.90		0.151	0.152	0.159	0.170		
LSD at 5 % for GA3 treatments Stimufol fertilizer treatm Interaction (GA3 X fertilize				N.S 0.87 0.91					N.S 0.016 0.022			

Table(8): Effect of GA₃ and Stimufol fertilizer treatments on leaf potassium and total carbohydates content (%) of Hemerocallis aurantiaca plants (2005/2006 and 2006/2007 seasons).

Pa	rameters			K%				Total	carbohyd	rates%	
			G	A3				G	A3		
Treatments		0.0	50 ppm	100 ppm	200 ppm	Mean	0.0	50 ppm	100 ppm	200 ppm	Mean
						First s	eason				
	0.0	1.32	1.46	1.49	1.45	1.43	9.2	9.8	10.2	10.6	10.0
Stimufol fertilizer	2g/L	1.41	1.49	1.57	1.52	1.50	9.8	10.7	10.9	11.2	10.7
Summon termizer	4g/L	1.53	1.58	1.62	1.64	1.60	10.6	10.3	11.6	11.9	11.1
·	6g/L	1.56	1.62	1.70	1.61	1.62	10.9	11.6	12.2	12.6	11.8
Mean		1.46	1.54	1.60	1.56		10.1	. 10.6	11.2	11.6	
LSD at 5 % for GA ₃ treatments Stimufol fertilizer treatn Interaction (GA ₃ X fertiliz				N.S 0.16 0.19					1.46 1.63 1.82		
						Second	season				
	0.0	1.39	1.43	1.48	1.47	1.44	9.8	10.3	10.9	10.5	10.4
Stimufol fertilizer	2g/L	1.47	1.48	1.53	1.50	1.50	10.3	10.8	11.6	10.8	10.9
Stimuloi ici tinzci	4g/L	1.46	1.53	1.59	1.52	1.53	10.7	11.6	12.2	11.3	11.5
	6g/L	1.52	1.57	1.63	1.58	1.58	11.2	11.9	12.9	12.8	12.2
Mean		1.46	1.50	1.56	1.52		10.5	11.2	11.9	11.4	
LSD at 5 % for GA3 treatments Stimufol fertilizer treatn				N.S 0.10					1.37 1.59		
Interaction (GA ₃ X fertilize	er)			0.18					1.78		

(2004) on *Strelitzia reginae*, El-Sayed (2004) on *Iris tingitana*, Abou-El-Ella (2007) on *Acanthus mollis* and Youssef and Goma (2007) showed that spraying *Iris tingitana* plants with Stimufol fertilizer at 4 or 6 g/L significantly increased leaf N, P, K and total carbohydrates contents.

The aforementioned results of GA₃ contaminant with those obtained by Tawila (2000) on *Polianthes tuberose*, Dantuluri *et al.*, (2002) on *Lilium maculatum*, Wankhede *et al.*, (2002) on *Polianthes tuberose*, Tiwari and Singh (2002) on *Lilium maculatum*, Gomaa (2003) on *Dahlia pinnata*, Salama (2003) on *Strelitzia reginae*, Youssef and Goma (2007) on *Iris tingitana* and Abou El-Ella (2007) reported that spraying *Acanthus mollis* plants with GA₃ at 100 or 200 ppm increased leaf N,P,K and total carbohydrates contents.

IV-Anatomical studies:-

IV -1. Regarding cross section of leaf:

As shown in Table (9) and Figure (1) different applied treatments enhanced in mostly characters of leaf compared with the control, especially leaf blade thickness, number of vascular bundle in blade. Also, it could be noticed that increases of vascular bundle length, thickness of phloem and xylem in vascular bundle,etc.

Table(9): Effect of GA₃ and Stimufol fertilizer treatments on histological features (micron) in leaf of Hemerocallis aurantiaca plants during 2007 season.

Treatments	Control	GA3 at 2	200 ppm
Characters (micron)		Stimufol at 4g/L	Stimufol at 6g/L
Thickness of blade.	706.50	638.10	663.00
Upper epidermal cuticle thickness.	11.70	15.75	14.40
Lower epidermal cuticle thickness.	7.65	11.25	15.30
Upper epidermal thickness.	54.00	58.80	62.55
Lower epidermal thickness.	22.85	29.25	33.75
Length of vascular bundle.	387.45	398.02	408.99
Thickness of upper fibers in vascular bundle	45.00	51.30	60.30
Thickness of phloem in vascular bundle	67.50	78.30	98.77
Thickness of xylem in vascular bundle.	146.92	164.70	187.80
Thickness of lower fibers in vascular bundle	102.60	103.05	105.97
number of vessels in the vascular bundle.	7.50	8.00	9.50
Thickness of widest xylem vessel in vascular bundle	41.40	43.65	37.80

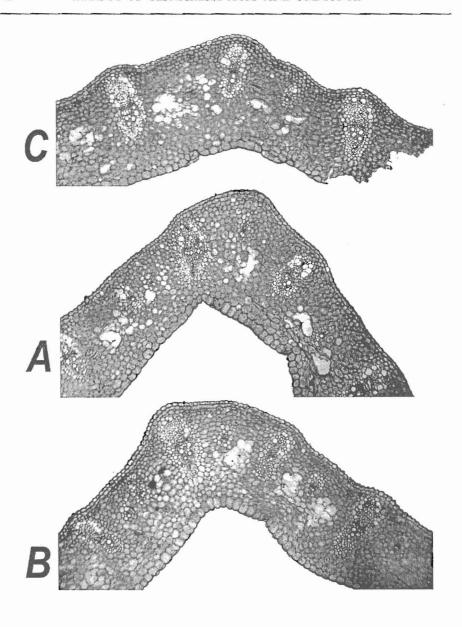


Fig. (1): Shows the effect of GA_3 and Stimufol fertilizer on histological features in leaf of Hemerocallis aurantiaca plants (100 x).

C-control A- GA_3 at 200 ppm + Stimufol fertilizer at 4g/L. B- GA_3 at 200 ppm + Stimufol fertilizer at 6g/L.

IV -2. Concerning cross section of flower stalk:

As shown in Table (10) and Figure (2) different applied treatments improved in mostly characters of flower stalk anatomy compared with the control, specially epidermal cell layer thickness, number and thickness of cortex and pith, number of vascular bundles and thickness of xylem and phloem.

Such enhancement attributed to GA3 may be due to the fact that GA3 stimulates the flower bud formation and enhance on cell division and elongation. The Stimufol fertilizers supply the plants with macro and micro elements to increase and improve the histological parameters.

In general, the stimulatory effects of the applied treatments upon the anatomy features of treated plants could be attributed to effects of GA₃ and Stimufol fertilizer in supplying the plants with their requirements of nutrients for more carbohydrates and proteins production which are necessary for vegetative, flowering, bulbs growth and chemical composition.

The same trend was found by Rak and Nowak (1989) on snapdragon cuttings; El- Shourbagy *et al.*, (1995) on flax; Marschner (1995) on higher plant; Shou-Fang *et al.*, (2003) on Chinese chestnut and Naor *et al.*, (2004) on *Iris tingitana*.

Conclusively, in order to produce early flowered *Hemerocallis aurantiaca* plants with higher flowers number, longer flower stalk and heaviest flower fresh and dry weights, it is preferable to spray the plants with the combined treatments between GA3 at 200ppm and Stimufol fertilizer at 4 or 6g/L.

Table (10): Effect of GA_3 and Stimufol fertilizer treatments on the histological features (micron) of the flower stalk of *Hemerocallis aurantiaca* plants during 2007 season.

Treatments	Control	GA ₃ at 200 ppm	
Characters (micron)		Stimufol at 4g/L	Stimufol at 6g/L
Flower stalk diameter.	4209.79	5145.30	6174.36
Epidermal cuticle thickness.	11.25	13.30	16.20
Epidermal cell thickness.	20.70	21.60	25.20
Cortex layers thickness.	288.00	387.00	403.20
Thickness of fibers in vascular bundle	147.60	148.50	163.80
number of vascular bundles.	86.00	91.00	117.00
Length of large vascular bundle.	515.70	1319.76	1375.57
Thickness of phloem in vascular bundle	129.15	141.75	153.90
Thickness of xylem in vascular bundle.	140.23	154.20	148.65
number of vessels in xylem bundle.	18.00	19.50	23.00
Thickness of widest xylem vessel in vascular bundle	45.00	69.30	69.75
Parenchymatous pith thickness.	640.00	1350.00	1728.00

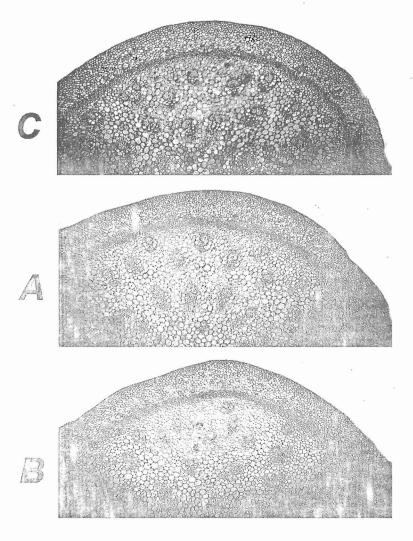


Fig. (2): Shows the effect of GA₃ and Stimufol fertilizer on histological features in flower stalk of Hemerocallis aurantiaca plants (100 x).

C-control A- GA_3 at 200 ppm + Stimufol fertilizer at 4g/L. B- GA_3 at 200 ppm + Stimufol fertilizer at 6g/L.

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تأثير الجبريللين وسماد الأستيموفول علي نمو وإزهار نبات الهميروكالس. فاتن حسن محمود إسماعيل*, أحمد سعيد محمد يوسف** *قسم النبات الزراعي-**قسم البساتين - كلية الزراعة ـجامعة بنها

أجريت هذه التجربة بمزرعة الزينة بقسم البساتين ومعمل التشريح بقسم النبات بكلية الزراعة بمشتهر - جامعة بنها وذلك خلال موسمي 2006 و 2007 لدراسة تأثير 16 معاملة ممثلة في التفاعل بين أربعة تركيزات من الجبريللين (صفر, 50، 100 و 200 جزء في المليون) وأربع تركيزات من سماد الاستيموفول (صفر, 2, 4, 6 جم/ لتر) على النمو الخضري، الزهري، القياسات التشريحية والمحتوى الكيماوي لنبات الهميروكالس.

وقد أوضحت النتائج المتحصل عليها أن جميع تركيزات الجبريللين قد حسنت من قراءات النمو الخضري (عدد الأوراق, الوزن الطازج والجاف للأوراق,عدد الخلفات) و قياسات الإزهار (موعد الإزهار أعدد الإزهار أبيات علول وقطر الحامل الزهري الوزن الطازج والجاف للأزهار وطول مدة بقاء الأزهار علي النبات وأيضا في الفازة) والقياسات التشريحية وقياسات المحتوى الكيماوي (الكربوهيدرات الكلية , النتروجين, الفسفور, البوتاسيوم) وخاصة التركيزات المتوسطة والمرتفعة. كما وجد أن جميع مستويات سماد الأستيموفول قد حسنت من جميع القراءات المدروسة وخاصة التركيزات المتوسطة والمرتفعة. كذلك وجد أن جميع التفاعلات بين الجبريللين وسماد الأستيموفول قد حسنت جميع القراءات المدروسة وخاصة المعاملات المختلطة بين الجبريللين وسماد الأستيموفول قد حسنت جميع القراءات المدروسة وخاصة المعاملات المختلطة بين الجبريللين

أما بالنسبة للدراسات التشريحية فقد أظهرت اختلافات واضحة بين القعاملات المختلفة حيث أن معظم معاملات التفاعل بين الجبريللين وسماد الأستيموفول أعطت أفضل القياسات بالنسبة للقطاعات العرضية لنصل الورقة والحامل الزهري مقارنة بالكنترول. و كانت أفضاً المعاملات هي المعاملات المختلطة بين الجبريللين بتركيز 200 جزء في المليون وسماد الأستيموفول بتركيز 6 جم/ لتر.

ومن خلال النتائج المتحصل عليها يمكن التوصية بمعاملة نباتات الهمير وكالس بالمعاملة المختلطة من الجبريللين بتركيز 200 جزء في المليون وسماد الأستيموفول بتركيز 6 جم/ لتر لكي يتم الحصول علي أفضل نمو خضري و زهري مع أعلي جودة.