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# **Original Research Article**

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# Effect of GA<sub>3</sub> on Reproductive Growth and Cormel Production of Gladiolus

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## ABSTRACT

# Keywords

Cormel, Gladiolus, Gibbrellic Acid, Reproductive traits

#### **Article Info**

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Gladiolus (family Iridaceae) is one of the most important bulbous flower crops due to its long attractive spike with variety of colours, prolonged vase life and its ability to withstand long distance transportation. Four different varieties of Gladiolus (V1-American Beauty, V2-Pacifica, and V3- Summer Pearl and V4-Single Jester) with uniform size of corms were taken for assessing the effect of four concentrations of GA<sub>3</sub> (25, 50, 75 and 100 ppm) on cormel production. The experiment was conducted in Factorial Randomized Block Design. Un-treated control was also maintained. The observations were recorded on various vegetative growths, cormel production and floral attributes. Spikes were harvested when basal florets showed colour. In this investigation GA<sub>3</sub> 100 ppm was found most effective for enhancing reproductive growth Number of florets per spike (16.10) time taken for Colour breaking (89.53 days) were observed at 50 ppm. Maximum floret size (13.16cm) and earlier 50% flowering (84.81days) were recorded at 25 ppm GA<sub>3</sub>. Treatment with GA<sub>3</sub> showed delayed 50% heading in (101.66 days) at 100 ppm. The results revealed that var. Summer Pearl recorded earlier colour break (89.53 days), earlier 50% flowering (84.80 days) and maximum cormel production (16.43) maximum number of florets per spike (16.10) and maximum florets size (13.16 cm) were exhibited by var. American Beauty.

## Introduction

Floriculture has now emerged as an important venture in the world. It is available in a variety of colours and inflorescence having attractive florets. Gladiolus is an important bulbous flower grown throughout the world. It has originated in tropical and southern Africa. It belongs to family Iridaceae and is extensively grown for cut flower production apart from being used in herbaceous border, bedding, and pot culture. Commercial

cultivation of gladiolus has gained popularity due to its high export potential and prevalence of congenial growing condition in various parts of the country. Now a day in India cultivation of gladiolus is the focus of large floriculture industries because of its popularity as cut flower and long keeping quality. In India, West Bengal, Maharashtra, U.P., Himachal Pradesh, Haryana and Andhra Pradesh are the major gladiolus growing states. It has added advantage that flower stalk can be packed in bulb stage, when the

lower florets just show their colours. Therefore, its cultivation as a cut flower can be taken up in far off areas provided they are well connected with big cities. Moreover during winter the climate in the Indian plain is ideal for cultivation of gladiolus thus providing a comparative for export to European countries where sub-zero temperature and snow cover during winter is a limitation for successful cultivation of gladiolus. Jharkhand with its diverse agroclimatic conditions have great potentialities of flower cultivation which will help the small and marginal farmers generating remunerative self-employment and also for earning foreign exchange through export floricultural produce.

Growth regulators have been found to influence the growth and flowering of gladiolus.GA3 has many regulatory effects on plant development. It stimulates synthesis of hydrolytic enzymes for digestion endosperms reserves. GA<sub>3</sub> stimulate both cell division as well as cell enlargement. It can move freely along the stem in either acropetal or basipetal direction. The role of gibberellins complicated biologically, both biochemically and even today is not fully understood (Arora et al., 1992). Singh et al., (2008) reported that gibberellic acid treatment cause to increase in longevity of gladiolus cut flowers; Mutuee et al., (2001) concluded that gibberellins increase fresh weight of cut flowers; Weiss in 2000 reported that gibberellic acid could be a promoter for starting extension, coloration of flower and some genes of essential enzymes anthocyanin pathway such as chalkon chalkon synthetase. isomerase, dihydroflavinol reductase. This plant growth regulator is produced in anthers, which are in developmental stage and then it is transferred into corolla where it enhances physiological and biological processes such as petal growth and pigment production.

The plateau region of Jharkhand offers an excellent climatic condition for cultivation of gladiolus, but little work has been conducted on the standardization of technology for gladiolus growing in this region. For successful production of any bulbous crop, growth regulators play an important role. Hence, considering the above points an experiment entitled "Effect of GA3 on reproductive growth and cormel production of gladiolus" was undertaken.

## **Materials and Methods**

# **Experimental site**

The site is located in the experimental area of the department of horticulture, under the faculty of Agriculture, Birsa Agricultural University, Kanke, Ranchi, Jharkhand during Kharif season of 2010.

# **Meteorological condition**

The climate of Ranchi is sub humid tropical. The geographical situation of the place is between 23.17° latitude and 85°19 E longitude and at an altitude of 625m above the mean sea level. The average annual rainfall is 1300-1400 mm and most of which is received between months of June to August. The climate is sub divided into distinct season i.e. summer, rainy and winter. Data on weather conditions were collected from meteorological observatory of the university throughout the experimental period.

The distribution and occurrence of the different aspect of have been illustrated in Figure 1. The mean daily temperature of the area is about 19.50° C, the region is characterized by moderate summer and cold winter with lowest January temperature (coldest month) dropping down to 3.5° C and highest May temperature (hottest month) rising upto 39.2° c. The mean humidity is about 64 percent in the area.

## **Experimental detail**

#### **Treatments and varieties**

The present investigation was carried out with four different levels of gibberellic acid (GA<sub>3</sub>) applied on the corms of four different varieties of Gladiolus (V1-American Beauty, V2-Pacifica, and V3- Summer Pearl and V4-Single Jester). Varieties of gladiolus were procured from department of horticulture, B.A.U., Kanke, Ranchi. The treatment combinations used in this experiment have been given in Table 1.

## Layout, plan and experimental design

The field experiment was laid out in Factorial Randomized Block Design (FRBD) with 20 treatments and three replications. There were three sub-plots, each having a size of 5mx 5m. All the treatments were accommodated in sub- plot randomly and each sub-plot consisted of 200 plants at a distance of 50cm x 20cm. A space of 50 cm was left between two sub plots.

## **Fertilizers**

NPK was applied in two split doses, as basal application doses @ 10:20:20g/ Sqm and @5:10:10/5g/Sqm in top dressing.

FYM applied as soil application @Uniform dose of 5 Kg/sqm.

# Preparation of experimental plot

The plot was cleaned by removing weeds, stubble and was thoroughly ploughed till the soil was brought to fine tilth. The plot was leveled and laid out according to plan. The well decomposed farm yard manure at the rate of 5kg/sqm was added to each plot and was incorporated in soil. Soil sampling was done randomly from different plots as per the soil

sampling techniques. The composite soil sample was prepared by mixing these samples and the composite soil sample was subjected to analysis. NPK were applied inform of urea, and MOP respectively SSP 10:20:20g/sqm. FYM was applied at the rate 5kg/sqm. For this 125 kg was applied in each plot. The corms of uniform size from all were selected; covering varieties removed gently without damaging the corms. The corms were soaked in GA3 for at least 24 hrs and dried properly under the shade and were sown in each plot according to layout and plan. Plants were irrigated as needed or whenever necessary throughout the cropping season by flooding method. Weeding and hoeing was done regularly starting from 15 days onwards after corms planting and whenever necessary throughout growing period. Regular Spraying of the plants with 0.2 percent Bavistin at an interval of 15-20 days was done to avoid the incidence of diseases. Spikes were harvested when the basal bud was just showing color. Spikes were cut just about where the stalk emerged from the sheathing leaf cluster. Corms were dug out after 45 days from flowering when the leaves started yellowing. They were dug out carefully so that no corms and cormels were lost in the process. The aerial parts, corms and cormels were separated and counted. The corms were dipped in 0.2 percent Bavistin solution for half an hr. and then air dried in shade and kept in Kraft paper bags in open crates for storage.

## **Recording of observation**

Following observation on growth, yield and quality of flower, corms and cormels were recorded at proper time during experimentation.

Number of florets per spike Floret size (cm) Colour break of first florets (days) Days to 50% flowering Cormel production Vase life

# Statistical analysis

Date obtained from several attributes was analyzed statistically by the procedure described by Gomez and Gomez, 1984.

#### **Results and Discussion**

The result with logical interpretation as influenced by different treatment has been presented in chapter with the help of appropriate table.

# Number of florets per spike

Regarding number of florets per spike has been presented in Table 2. From the perusal of data, it was observed that different levels of GA<sub>3</sub> had significantly affected the number of florets per spike and it was found that T4 level produced the maximum number of florets (14.55), followed by T2 (11.85) and T3 (11.54). The different varieties also showed significant influence on number of florets. V1 produced maximum number of florets per spike (12.62), followed by V4 (12.02). The interaction of VXT also had significant influence on number of florets per spike. The treatment combination V1T2 (16.10) recorded maximum number of florets per spike (16.10), followed by V3T4 (15.36) and V4T4 (15.00) was least number of floret per spike was recorded in V2T1 (8.97)

#### Florets size (cm)

The diameter of floret (cm) was recorded and presented in Table 3. Floret size of 12.53 cm was measured in T4 which was significantly maximum as compared to different levels of GA3 in various treatments. However, different varieties had no significant influence

on floret size. Though maximum floret size of 9.63 cm was recorded in V2 followed by V4 (9.52cm), As evident from the data, the interaction of VXT was found effect the floret size significantly. Maximum floret size of 13.16 cm was recorded under treatment combination V1T1 followed by V4T4 (12.53). The treatment combination V1T0 recorded the least floret size (6.66cm).

## Days taken to colour break of basal florets

The observations on number of days taken for colour break of basal florets have been presented in Table 4. From the perusal of data it was revealed that different levels of GA3 affected the character significantly comparison to control (T0) where GA3 was not used Earliest colour break was recorded with T3 (93.88 days), followed by T1 and T2 (98.80 days). The higher doses of GA3 took lesser number of days for colour break of basal floret as compared T0 in (130.12 days). Similarly, different varieties were found to have non-significant effect on days taken to colour break of basal florets. However, earliest colour break was recorded in V1 (103.04 days) followed by V3 (103.57 days). The interaction of VXT significantly influenced on days taken for colour break of basal floret. Earliest colour break was recorded with treatment combination V3T2 (89.53 days) followed by V2T3 (89.92 days), the maximum number of days was recorded treatment combination V3T0 under (132.80days)

# Days to 50% flowering

The data pertaining to days to 50% flowering have been presented in Table 5. From the perusal of data it was evident that minimum number of days to 50% flowering was recorded with T4 (95.73 days) which was significantly superior to all the levels of GA3 except T3 (97.55 days) and T2 (104.39 days)

which were at par with T4 (95.73 days). The varieties failed to produce any significant influence on days to 50% flowering. However, shortest duration was recorded with V3 (101.63 days) followed by V2 (103.40). As evident from data, the interaction of VXT was found to have non-significant influence on days to 50% flowering. However least number of days for 50% flowering was recorded by treatment combination V3T1 (84.81days), while maximum numbers of days were recorded in V4T0 (122.61 days)

# **Cormel production**

The numbers of cormels were recorded at harvest and have been presented in Table 6. The data revealed that number of cormels were significantly maximum in T4 (13.57) followed by T3 (9.54). Although, the effect of varieties on cormel production, was found to non-significant. However maximum cormel production was recorded in V3 (9.52). The result revealed that interaction VXT significantly increased production V3T4 recorded significantly maximum number of cormels (16.43), followed by V4T4 (13.13) in comparison to all other treatment combinations.

#### Vase Life

The data pertaining to vase life of cut flowers have been presented in 7. Perusal of data revealed that vase life of cut flowers was significantly influenced by GA<sub>3</sub>. With application of higher concentration of GA3 that is (100 ppm) cut flowers survived maximum number of days in the vase (11.68days). The varieties did not influence the vase-life significantly, however maximum vase-life was observed with the V3 (10.76days). The interaction VXT showed that GA3 in combination with different varieties significantly enhanced the vase-life. The treatment combination V4T4 recorded

the maximum vase-life of 14.6 days as compared to only 7.76 days in controls.

Numbers of florets per spike recorded at final stage revealed that this character was maximum at higher doses of GA<sub>3</sub>. Nilimesh and Ray Choudhary (1988) observed that number of florets per spike increased significantly with the use of GA<sub>3</sub>. Bose et al., (1980) reported that in gladiolus, the number of florets was least affected by growth regulators. Dua et al., (1984) reported increase in number of florets using higher concentrations of GA<sub>3</sub> as compared to lower concentrations of GA<sub>3</sub> on gladiolus. Ravidas et al., (1992) observed greatest plant growth and higher number of florets per spike with GA<sub>3</sub> treatment (50-100ppm); Karaguzel (1999), observed increased flower number by soaking of corms at 100ppm GA<sub>3</sub> for one hour. It was found maximum (16.10) at 50 ppm GA<sub>3</sub>. This phenomenon with respect to increase number of florets is in full agreement with Karaguzel (1999).

The data on days taken to colour break for basal floret showed significant effect (Table 4). Earlier colour break of basal floret (89.92 days) was observed in the Variety Pacifica treated with 75ppm GA<sub>3</sub>. It may be due to higher anthocyanin pigment biosynthesis. Gibberellin increases the activity of anthocyanin gene transcription and aggregate pigment in corolla.

Berman and Rajni (2004) working with gladiolus cultivar American Beauty observed that GA<sub>3</sub> showed significant influence on growth and flowering (colour break of basal floret); Goren *et al.*, (1990), working with gladiolus reported that GA<sub>3</sub> significantly increased the floral traits (earlier colour break of basal floret); Ramachandrudu and Thangam (2007), reported that GA<sub>3</sub> (150 ppm and 100 ppm) advanced the flowering (days to spike emergence and flowering).

**Table.1** Symbol of different treatments

Sl. No.	Symbol	Treatments (GA <sub>3</sub> PPM)
1.	T0V1	Control
2.	T1V1	25 ppm
3.	T2V1	50 ppm
4.	T3V1	75 ppm
5.	T4V1	100 ppm
6.	T0V2	Control
7.	T1V2	25 ppm
8.	T2V2	50 ppm
9.	T3V2	75 ppm
10.	T4V2	100 ppm
11.	T0V3	Control
12.	T1V3	25 ppm
13.	T2V3	50 ppm
14.	T3V3	75 ppm
15.	T4V3	100 ppm
16.	T0V4	Control
17.	T1V4	25 ppm
18.	T2V4	50 ppm
19.	T3V4	75 ppm
20.	T4V4	100 ppm

Table.2 Number of florets per spike of gladiolus varieties as affected by levels of GA<sub>3</sub>

	V1	V2	V3	V4	Mean
T0	10.13	10.90	11.50	11.40	10.98
<b>T1</b>	11.31	8.97	9.65	12.03	10.49
<b>T2</b>	16.10	12.73	9.08	9.50	11.85
T3	11.48	12.13	10.47	12.20	11.54
<b>T4</b>	14.10	13.76	15.36	15.00	14.55
Mean	12.62	11.70	11.20	12.02	
	VARIETIES	TREATMENT	VxT		
SEm	0.28	0.31	0.63		
CD5%	0.81	0.91	1.82		
CV5%	9.32				

WHERE, T0 = Control, T1 = 25ppm, T2 = 50 ppm, T3 = 75 ppm, T4 = 100ppm V1 = American Beauty, V2 = Pacifica, V3 = Summer Pearl, V4 = Single Jester

Table.3 Floret size (cm) of gladiolus varieties as affected by levels of GA<sub>3</sub>

	V1	V2	V3	V4	Mean
<b>T0</b>	6.66	8.40	8.70	9.60	8.36
T1	13.16	7.23	8.80	9.73	9.79
T2	9.53	12.50	6.90	8.60	9.38
T3	9.80	10.20	11.40	7.10	9.60
T4	8.16	9.83	10.07	12.53	12.53
Mean	9.46	9.63	9.18	9.52	
	varieties	treatment	vxt		
SEm	0.15	0.17	0.34		
CD 5%	N.S.	0.49	0.99		
CV 5%	6.38				

WHERE, T0 = Control, T1 = 25ppm, T2 = 50 ppm, T3 = 75 ppm, T4 = 100ppm V1 = American Beauty, V2 = Pacifica, V3 = Summer Pearl, V4 = Single Jester

**Table.4** Colour break (Days) of first basal floret of gladiolus varieties as affected by levels of  $GA_3$ 

	V1	V2	V3	V4	Mean
T0	127.80	129.90	132.80	129.82	130.12
T1	95.93	95.36	99.66	124.18	98.81
T2	99.66	110.72	89.53	95.33	98.81
T3	95.19	89.92	97.65	92.75	93.88
<b>T4</b>	96.53	108.53	98.16	95.75	99.75
Mean	103.04	106.90	103.57	107.57	
	varieties	treatment	vxt		
SEm	2.24	2.50	5.07		
CD5%	N.S.	7.18	14.36		
CV5%	8.20				

WHERE, T0 = Control, T1 = 25ppm, T2 = 50 ppm, T3 = 75 ppm, T4 = 100ppm V1 = American Beauty, V2 = Pacifica, V3 = Summer Pearl, V4 = Single Jester

**Table.5** Days to 50% flowering of gladiolus varieties as affected by GA<sub>3</sub>

	V1	V2	V3	V4	Mean
T0	114.30	116.00	107.25	122.61	115.40
T1	119.36	113.29	84.81	115.43	108.21
<b>T2</b>	96.79	88.33	119.50	113.00	104.39
T3	98.66	100.73	96.96	93.83	97.55
T4	99.26	98.73	99.63	85.30	95.73
Mean	105.67	103.40	101.63	106.03	
	varieties	treatment	vxt		
SEm	3.72	4.16	8.33		
CD 5%	NS	11.93	NS		
CV 5%	13.80				

WHERE, T0 = Control, T1 = 25ppm, T2 = 50 ppm, T3 = 75 ppm, T4 = 100ppm V1 = American Beauty, V2 = Pacifica, V3 = Summer Pearl, V4 = Single Jester

Table.6 Cormel production of gladiolus varieties as affected by GA<sub>3</sub>

	V1	V2	V3	V4	Mean
<b>T0</b>	2.66	4.83	6.13	7.00	5.15
T1	7.40	4.73	8.13	11.60	7.96
T2	12.13	12.80	6.13	7.16	9.55
T3	8.80	8.46	10.80	8.12	9.04
T4	12.26	12.46	16.43	13.13	13.57
Mean	8.65	8.66	9.52	9.40	
	VARIETIES	TREATMENT	VxT		
SEm	0.43	0.48	0.97		
CD 5%	N.S	1.40	2.80		
CV 5%	18.60				

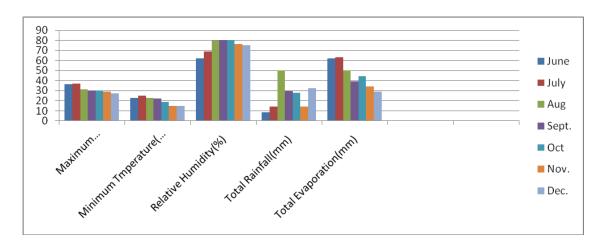
WHERE, T0 = Control, T1 = 25ppm, T2 = 50 ppm, T3 = 75 ppm, T4 = 100ppm V1 = American Beauty, V2 = Pacifica, V3 = Summer Pearl, V4 = Single Jester

**Table.7** Vase life (days) of gladiolus varieties as affected by levels of GA<sub>3</sub>

	V1	V2	V3	V4	Mean
<b>T0</b>	7.76	8.20	10.96	12.13	9.76
<b>T1</b>	7.70	8.26	9.00	14.06	9.75
<b>T2</b>	9.60	9.03	7.93	15.03	10.40
T3	8.80	11.36	14.40	7.50	10.51
<b>T4</b>	8.70	12.26	11.16	14.60	11.68
Mean	9.78	10.69	10.76	10.45	
	VARIETIES	TREATMENT	VxT		
SEm	0.27	0.30	0.61		
CD 5%	NS	0.87	1.70		
CV 5%	14.70				

WHERE, T0 = Control, T1 = 25ppm, T2 = 50 ppm, T3 = 75 ppm, T4 = 100ppm V1 = American Beauty, V2 = Pacifica, V3 = Summer Pearl, V4 = Single Jester

**Fig.1** Meteorological condition during the period of experimentation



Characteristics of florets in terms of diameter (Table 3) were improved significantly due to different treatment combination. The increase in diameter of the florets represents expansion of the cell of florets. Yousuf et al., (2006), observed that presoaking of gladioli corms for 24 hrs at different concentration (0-100ppm) in GA3 increased flower diameter. Chattar et al., (2006) conducted field trials with gladiolus cultivars, (Red Beauty, Jester, Summer Face) and four levels of GA<sub>3</sub> (0,100,200 and 300 ppm). They observed maximum floret size (13.01cm and 12.83cm); Umrao et al., (2006) obtained significant increase in diameter of the florets by treatment with GA<sub>3</sub> at 100 ppm; Berman and Rajni (2004) working with gladiolus cultivar American Beauty observed that GA<sub>3</sub> (75p pm) significantly influenced on floral diameter of gladiolus. Our findings are in full agreement with Yousuf et al., (2006).

Duration of 50% flowering as affected by different treatment combination was found to significant (Table 5). Earlier flowering was obtained in Summer Pearl at 50 ppm GA<sub>3</sub>, It may be due to increased level of exogenous gibberellins activity in plants as gibberellins enhance cell elongation and cell enlargement. (Salisbury and Ross, 1986); Ravidas et al., (1992) observed greatest plant growth and earliest flowering with GA3 treatment (50-100ppm); Prakash and Jha (1998) observed that GA<sub>3</sub> treatment at 150ppm improved all the floral traits including earlier flowering; Nilimesh and Roy Choudhary (1988), observed the decrease in days to flowering with GA<sub>3</sub> treatment.

The effect of GA<sub>3</sub> on cormel production was found to be significant (Table 6). Large number of cormels was obtained in variety Summer Pearl at 100ppm GA<sub>3</sub> (16.43). It may be due to exogenous gibberellic acid, which promotes extensive growth of intact plant (Salisbury and Ross1989), and thus increase

the number of cormels. The present finding are in close agreement with Bhattacharjee (1984), Singh et al., (1996) who recorded increased number of cormels per plant at 75ppm GA<sub>3</sub>. Berman and Rajni (2004) observed increase number of cormels per plant by Soaking of corms for 24 hrs; Yousuf et al., (2006) also observed that presoaking of gladioli corms for 24 hrs at concentration (0-100ppm) in GA<sub>3</sub> increased number of cormels per plant; Dua et al., (1984) found better corm multiplication when the plants were sprayed thrice with 100ppm GA<sub>3</sub>; Kumar and Singh (2008) observed application of GA<sub>3</sub> (150 ppm) resulted in maximum yield of corms per hectare; Umrao et al., (2006) obtained increase in number of cormels per plant by soaking of corms of cultivar Nova Lux in GA<sub>3</sub> (100 and 150 ppm. Ogale et al., (2000), reported importance of GA<sub>3</sub> for vegetative growth and cormel production of gladiolus.

The influence of GA<sub>3</sub> on vase life was found to be significant (Table 7), longest vase life (14.60 days) was obtained in variety Single Jester at 100ppm GA<sub>3</sub> (14.60 days). It may be gibberellic acid, it increases conservation of carbohydrates in stem and petal tissue and improves cell membrane thus delaying membrane degradation in cell (Kwack et al., 1997), Pal and Choudhary (1998) also reported maximum spike length at 20 ppm GA<sub>3</sub>, while 40 ppm GA<sub>3</sub> produced spikes having longest life in the field. Chattar et al., (2006) conducted field trials with gladiolus cultivars,(Red Beauty, Jester, Summer face.) with four different levels of GA<sub>3</sub> (0,100,200,and300 ppm). They observed maximum vase life (14.33 and 13.70) at 200 and 300 ppm GA<sub>3</sub>. Umrao et al., (2006) obtained maximum longevity and vase life at GA<sub>3</sub> 100 and 150ppm. Emami et al., (2011) observed maximum vase life at 75ppm GA<sub>3</sub>; Singh et al., (2008) observed that GA<sub>3</sub> treatment increased longevity of gladiolus cut flowers. Our findings are in full agreement

with Chattar *et al.*, (2006), who reported longest vase life in higher concentration of GA<sub>3</sub>.

Summary and Conclusions are as follows:

Total number of florets per spike with maximum 14.55 at T4 (100ppm) and the variety V1 (12.62) was found best while the treatment combination V1T2 (16.10) recorded maximum number of florets.

The diameter of the floret was recorded maximum (12.53cm) with T4 level of GA<sub>3</sub>. In case of varieties, maximum floret size (9.63cm) was recorded in V2.The interaction result V1T1 recorded maximum floret size of 13.16cm.

GA<sub>3</sub> levels T<sub>3</sub> (75ppm) recorded earliest colour break in 93.88 days. Minimum 103.04 days were taken for colour break of basal floret by V1. The interaction result V3T2 recorded 89.53 days for colour break

The duration of 50% flowering was observed with T4 (95.73days). By decreasing the levels of GA<sub>3</sub> 50% flowering duration prolong to 104.39 days in T2. The variety V3 recorded 101.63 days for 50% flowering. Interaction of VXT i.e V3T1 recorded least number of days (84.81) for 50% flowering.

The number of cormels was recorded maximum with T4 (13.57). Maximum cormel production was recorded in V3 (9.52).

The treatment combination V3T4 recorded maximum number of cormels 16.43.

Vase life of cut flower was recorded maximum (11.68days) with application of T4 (100ppm GA<sub>3</sub>). The maximum vase life 10.76 days was obtained by the variety V3. The treatment combination V4T4 recorded maximum vase life of 14.16 days

On the basis of above finding it can be safely inferred that GA<sub>3</sub> at all levels recorded better superior Reproductive growth, vield attributing features and floral traits as well as prolonged vase life and higher net returns in different cultivar of gladiolus. Though all the four varieties responded positively to GA<sub>3</sub> treatment at all levels but the variety American Beauty along with concentration treatment with 100 ppm GA<sub>3</sub> gave the best result i.e. floret size, and number of floret per spike and hence can be recommended for commercial cultivation.

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