

BIBLIOGRAPHY

JOEL S. CATONES. OCTOBER 2008. Effect of Gibberellic Acid GA₃ on the Growth and Flowering of milflores (*Hydrangea Macropylla*). Benguet State University

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ABSTRACT

The study was conducted at Lamtang, Puguis, La Trinidad, Benguet from November 2007 to March 2008 which aimed to compare the effect of GA₃ concentration on the vegetative growth and flowering of potted milflores plants; to determine the effect of different gibberellic acid GA₃ on the growth and flowering of milflores, and to determine the best concentration of gibberellic acid GA₃ that will enhance vegetative growth with better flower quality.

The earliest to develop flower buds observed were on milflores plant treated at three month and the longest flower bud to develop was one month from treatment. Plants sprayed with 50 ppm GA₃ were the earliest to develop flower buds while untreated plants required longer days to develop flower buds.

Milflores plants treated with smaller amount of GA₃ concentration was the earliest to develop flower followed by the higher amount of GA₃ concentration. Untreated plants show delayed flower development and flowering.

The highest number of leaves counted at 75% anthesis was produced by the plant treated at two months. Plant sprayed with 100 ppm GA₃ has the highest number of leaves counted followed by 50 ppm.

Milflores plants treated at two month produce the longest stem measured at 75% anthesis followed by one month and three month. Plant sprayed with 250 ppm GA₃ the longest stem measured followed by 100 ppm.

The widest flower diameter was produced by 250 ppm followed by 100 ppm GA₃ concentration at two month. The smallest flower diameter measured was observed in untreated plants.

The thickest stems were measured on milflores plant treated with 250 ppm and followed by plants sprayed with 50 and 100 ppm GA₃. Two month produce the thickest stems followed by three month and one month from treatment.

Plats treated at one month produce the largest number of leaves of lateral stems followed by three month and two month. GA₃ have a slight effect on the lateral stems of milflores plants.

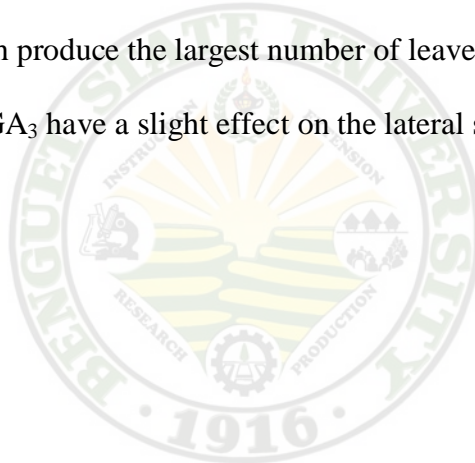


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INTRODUCTION

Nature of the Study

Hydrangea is a genus of shrub of the Saxifrage Family consisting of about 80 species. One of these species is the milflores (*hydrangea macrophylla*). Hydrangea is a strong deciduous shrub having a height and width of 6 feet. Leaves are broadly ovate, 4-6 inches long and 2.75-4 inches wide with coarse serrations, yellowish to fresh green in color. Flowers are in large bracts globose cyme of the hortensia or mop-head type with abundant, mostly sterile, individual florets, 4-5 petal linked sepal per flower (Anon, 2002).

This group consists of hardy and tender shrubs and woody climbers. They are mostly deciduous plants, though a few of the tender species are evergreen. They are native of the Himalayas, North and South America, and central and eastern Asia. These flowering shrubs have different flower forms from the large globes of the “mopheads” to the discs of the “lacecaps” to the thick cones of the oakleaf and panicle hydrangea. They come in an array of colors from pure white to brilliant crimson, pale lilac to intensive azure. Some varieties produce blossoms with two-toned colors, while some flowers have contrasting eyes, and some even is speckled or striped with another color. There are two kinds of florets in the flower heads. The fertile or perfect florets are small and inconspicuous. They bear the male and female parts and are usually found in the center of the cluster. The flower heads or mopheads hydrangeas consist almost entirely of sterile florets. Besides their lovely flowers, some hydrangeas are valued for their attractive foliage or bark. The size of the plants ranges from dwarf (about 1 ft high) to large bushes with stems over 10 ft high. There are also climbing hydrangeas whose aerial can bring stem up to 80 ft high. *H. aspera* subsp. *Sargentia* (Sargent Hydrangea) is an attractive medium to large-sized



shrub with hairy shoots and large, plush leaves. The large, flat flower head are produced in mid-to late summer and are bluish with white ray florets. This variety is excellent for a sheltered shrub border or woodland, but need shade and wind protection. H. “Ayesha” is a deciduous shrub that has shiny green leaves and flattened, fragrant flower heads consisting of cup-shape florets, resembling those of the lilac. They are grayish lilac or pink, depending upon the soil, eventually turning a greenish-blue to turquoise color. H. macrophylla ‘Alltona’ is a mophead hydrangea with flower heads growing up to a foot in diameter. This deciduous plant forms a small shrub with large, rose colored florets. This variety grows best in shade. H. quercifolia (Oakleaf Hydrangea) is a medium-sized, deciduous shrub valued for its splendid autumn colors. The large, deeply lobed leaves resemble those of the oak, thus the common name. In late summer, conical heads of large, white, sterile florets are produced. H. serratifolia is a tender, climbing hydrangea that grows best against a wall in sun or shade. It has small, leathery leaves and columnar panicles, up to 5 inches long, of creamy flowers. (Anon. undated)

Importance of the Study

Due to the economic potential and aesthetic value of potted milflores as bedding plants for outdoor landscaping and cut flowers, it is important therefore to find the best way of enhancing growth with quality flower of milflores with the use of gibberellic acid.

Objectives of the Study

1. To determine the effect of gibberellic acid (GA_3) on the growth and flowering of milflores.
2. To determine the best concentration of gibberellic acid (GA_3) that will enhance vegetative growth with better flower quality.



3. The study was conducted to determine the best stage of gibberellic acid (GA_3) application and to;

4. Determine the interaction effect on various stage of gibberellic acid (GA_3) application and GA_3 concentration.

Place and Time of the Study

The experiment was conducted at the Ornamental Horticulture Research Area, Benguet State University, Benguet August 2007 to January 2008.



REVIEW OF RELATED LITERATURE

Gibberellic acid GA₃ belongs to a group of plant hormones that promote plant growth. It is one of the plant growth regulators that can provide a wide range of plant response such as breaking dormancy, incubation of flowering, parthenocarpic fruit set and sex expression (Devlin 1977).

As Devlin (1977) reported, gibberellins are naturally occurring hormones that have been isolated from higher plant tissues. They act similarly as indole acetic acid (IAA), another kind of hormone to promote cell elongation, induce parthenocarpy, promote cambial activity, and issue new RPA and protein synthesis.

Gibberellins are known to alter the growth of plants, since their applications has been found to cause rosette plants to elongation in a manner suggestive to bolting. Investigations have been made on effects of GA₃ on the floral induction. Their influence on the growth and flowering of some crops has been investigated in greenhouse and in the field for the past years. Studies revealed earlier opening of flowers in those plants treated with GA₃ and reduction on the number of days required for the formation of flower buds (Leopold and Kriedman, 1964)

Plants treated with gibberellins generally grow to three times taller than the untreated plants. Promotion of flowering particularly in biennials and long day plants is one of the most striking effects of gibberellins. Biennials commonly do not produce long stem, and flower only if they are exposed to low temperature. However, when treated with gibberellins, they flower without cold temperature (Grealach and Adams, 1962).

Findings revealed that gibberellins variety influence the growth and flowering productivity of Shasta daisy. Lower GA₃ concentrations of 600 ppm at repeated application



improve quality and hastened flowering. The treated plants appeared healthy and having standard flower stem length. Similarly, GA₃ at flowering concentrations with repeated applications affected the highest number of flowering per plant, but plants treated with the highest concentration had slight decrease in floral production (Dulay, 1980).

Mendoza (1993) stated that gibberellins acid treatment can induce flowering of some periodically sensitive and some cold requiring plants. This means that gibberellins can replace the cold requirements of some plants. Subsequently, it has also been shown that the induction of flowering may bring about a natural size in the endogenous gibberellins content within the plant.

Application of GA₃ as foliar spray to stem species at 10- 100 ppm hasten flowering by ten days to four weeks in petunia, stock, larkspur, English daisy, china aster, and gerbera (Weaver 1972).

Weir (1974) stated that gibberellins applied to long day rosette plant produce flowers under short to long day. Likewise gibberellins inhibited flower initiation in many other plants especially woody plants. It has been reported that flowering in Azalea was hastened from 25 to 59 days when sprayed with GA₃ in comparison to plant which received cold treatment for 30 days prior to forcing (Barba and Pokorny, 1961).

As Nickell (1986) stated, GA₃ along with kinetin also are known to promote flowering and prevent flower drop in tomato at high temperature. Devlin (1977) also reported that GA₃ are used to increase the number of grapes in the cluster. Other commercial uses the stimulation of flower bud formation and fruit set apples and pears and the improvement in size; color and quality of the fruit of many plants.

Caluya and Imlan (1959) reported that GA₃ induces the rapid elongation of plants in general. Even as early as one week after first application, the treated plants were already taller than



the untreated plants. Also, it was found that the more concentrated the growth regulator, the higher was the weekly height increasement in the stem length.

Application of 1,250 ppm GA₃ at three leaf-stage and four-leaf stage gladiolus bolted earlier, produce bigger sized corns, loner flower spikes at flowering with more and bigger flowers (Albis, 1995).

The earliness in flowering could be attributed to he exogenous application of GA₃ that promote flowering by increasing the mitotic cell division in the apex, thus, rendering it more responsible to flower stimulus. Zeevart (1962) farther stated that flowering could be enhanced by increasing the GA₃ level of the plant through nitrogenous application. Gibberellins cause or promote flowering by either facilitating the formation of flowering hormones in the leaves or the expressions in the growing bud.

Gibberellins appear to affect almost all plant organs from root to flower, fruit and seed development. Shein and Jackson 1972 reported that gibberellic acid applied to decapitated stems, petioles or to lands promote growth of buds in the axile of primary leaves.

Galimba (1993) reported that three application of 1000 ppm GA₃ at weekly intervals significantly promote earlier lower development, improve flower quality, produce bigger sized bloom at harvesting and produce taller plants with longer floral stems of anthurium cv. Kansako.

It was also reported that gibberellic acid treatment greatly affected the flowering of Mr. Lincoln roses (Mendoza, 1993). Gibberellic acid at 500 ppm with two applications significantly enhanced cutflower production, increase yield, higher number of leaves per plant at anthesis, longer flower buds and blooms higher leaf area and leaf index at flowering and promote longer vase life.



Hermano and Ladilad (1980) found that the rate of growth effected by 400 ppm, four weeks from treatment. However, plant treated with the highest concentration (1000 ppm at GA₃ finally showed the fastest rate of growth in the later part of the vegetative stage).

In 1979, Kim started that some change on chemical compounds of plants occurs when treated with gibberellins. These were an increase in water, crude fat and protein content and decrease in glucose, fructose, galactose, and xylose content of winter cultivated Chinese cabbage, the changes in chemical compounds could lead to he accumulation of gibberellins for the synthesis of florigen, a hypothetical flowering hormone which cause early floral evocations.

In general gibberellins are group of plant hormones that promote plant growth. They are used for encouraging plant growth, hastening germination, and encouraging growth in odd weather increasing the size of some fruits, increasing the yield of some cops, and breaking dormancy of various seeds. The response of gibberellins varies with the plant specie (Galingan, 1997).

MATERIALS AND METHODS



Material

The materials to be used in the experiment will be rooted milflores (*Hydrangea Macrophylla*) cuttings; gibberellic acid; polyethylene black plastic bags (5x5 inch) filled with the potting mix of garden soil, compost (1:1); watering tools; garden tools; and identifying tags.

Methods

The experiment will be laid out in a complete randomized design (CRD) in factorial arrangement with three replication. Factor A will be the stage of application and factor B will be the gibberellic acid concentrations. There will be five samples per treatment replicated three times. The treatment will be as follows.

Factor A (Stage of Application)

S1 -1 month from planting

S2 -2 month from planting

S3 -3 month from planting

Factor B (Gibberellic Acid Concentration)

C1 – 0 ppm/ control

C2 – 50 ppm

C3 – 100 ppm

C4 – 250 ppm

C5 - 500 ppm

Planting of Milflores



Milflores cutting will be rooted first in a nursery and then transplanted in 5x5 inches polyethylene black plastic bags with 1 part garden soil and 1 part compost and .5 part chicken dung.

Gibberellic Acid Application

The various concentration of gibberellic acid will be applied once on the shoot of milflores plants 15 days and 30 days from planting as specified in the treatment.

Other Cultural Management Practice

All other cultural management practice in milflores production such as watering, and control of pest (insects, disease and weeds) will be strictly employed as the need arise.

Data to be Gathered;

The data to be gathered will be as follows:

1. Days from transplanting to flower bud formation(0.5 cm bud size)

-The number of days from transplanting to (0.5 cm bud size) flower bud formation was counted.

2. Number of days from flower bud formation to 75% anthesis.

-The number of days from flower bud formation to 75% flower anthesis was counted

3. Average number of leaves per plant at 75% anthesis.

-The number of leaves per plant was counted at 75% anthesis at flowering.

4. Final height at flowering (cm)



- The height was taken by measuring the plant from base to the top at flowering (75% anthesis)

5. Flower diameter at full bloom stage (cm)

- The size of the flower was measured at flowering stage (75% anthesis).

6. Stem diameter (cm)

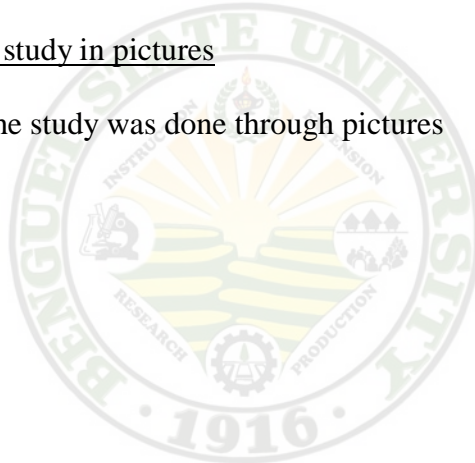
-This was obtained by measuring the diameter of the center of the plant at flowering (75% anthesis) with the use of vernier caliper.

7. Number of lateral stems at flowering

-This was counted by counting the lateral stems at flowering stage (75%)

8. Documentation of the study in pictures

- Documentation of the study was done through pictures



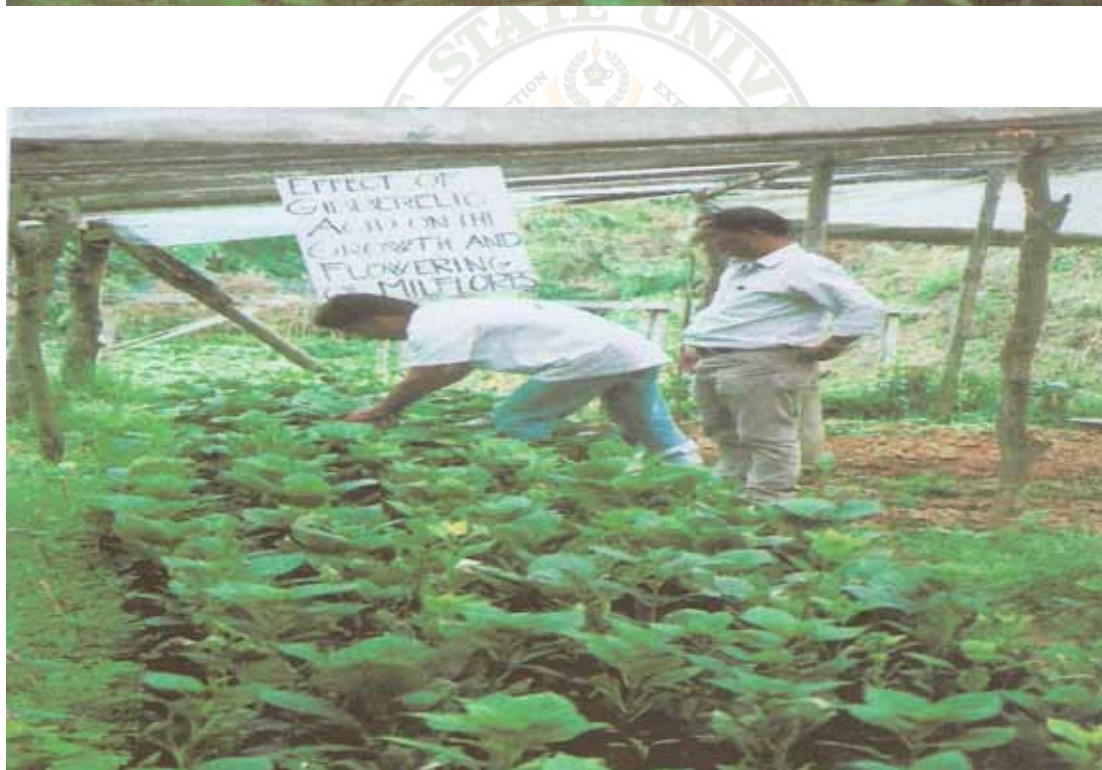


Plate 1. Gibberellic acid application



Plate 2. Gibberellic acid application



Plate 3. Days from flower bud formation to 75% anthesis.





Plate 4. Over view of potted milflores plants treated with gibberellic acid (GA_3) at flowering.



RESULTS AND DISCUSSION

1. Days from transplanting to flower bud formation (0.5 cm bud size)

Effect on the Stage of Application. Table 1 show that the effect of the stage of application had significant differences. Plant sprayed after three months from planting has the earliest bud to develop, followed by two months to form flower buds. One month was the last to develop flower buds from the time of application.

Weir (1974) stated that gibberellins applied to long day roseate plants produce flowers under short to long day. Like wise gibberellins inhibited flower initiation in many other plants especially woody plants.

Effect of GA₃ concentration. Result show that a significant difference was obtained on the number of days from transplanting to flower bud formation as affected by the different concentration of GA₃ applied on milflores. The untreated or control required the longest number of days to initiate flower buds from the time of transplanting. Plant sprayed with 50 ppm, was the earliest to develop flower buds, and followed by 100 ppm. Plant sprayed with the highest concentration of 500ppm has developed flower buds late among the treated plants.

Interaction Effect. A significant difference was obtained between the different GA₃concentration on the stage of application from transplanting to flower bud formation. Untreated plants showed significantly delayed of flower bud formation with 160.586 days; while the earliest to initiate flower buds were the plant sprayed with 50 ppm with 144.400 days. It was observe further, that as the amount of GA₃ concentration increases the number of days also increases.



Table 1. Days from transplanting to flower bud formation (0.5 cm)

ENTRY	MEAN
<u>Stage of Application</u>	
1 month	160.586a
2 month	157.040b
3 month	126.577c
<u>Gibberellic Acid Concentration</u>	
C1 – 0 ppm/ control	154.556a
C2 – 50 ppm	144.400c
C3 – 100 ppm	145.206c
C4 – 250 ppm	148.308b
C5 - 500 ppm	147.896

2. Number of Days from Flower Bud Formation to 75% Anthesis

Effect on the Stage of Application. Result shows no significant effect on the stage of application from flower bud formation to 75% anthesis stage as shown in table 2. However, two month from flower bud formation has reaches its 75% anthesis earlier followed by one month. Three month required longer number of days to reach 75% anthesis. Application of GA₃as foliar spray to stem species at 10-100 ppm hasten flowering by ten days to four weeks (Weaver, 1972).

Effect of GA₃ Concentration. Result showed significant differences on the number of days from flower bud formation to 75% anthesis as affected by the different GA₃ concentration. A delayed in flower development were observed in plants which were not sprayed with GA₃ concentration or the control. Plant treated with 100 ppm GA₃ concentration has the earliest to



reach its 75% anthesis stage, while plant sprayed with 500 ppm GA₃ concentration has reaches its 75% anthesis late.

Interaction Effect. Significant interactions were observed between the stage of application from flower bud formation to 75% anthesis and the plant sprayed with different GA₃ concentration. Plant sprayed with 100 ppm GA₃ concentration that is treated three month after flower bud formation has the earliest to reaches its 75% anthesis while control at three month required longer number of days to bloom at 75% anthesis.

Studies revealed earlier opening of flower in that plant treated with GA₃ and reduction on the number of days required for the formation of flower buds (Leopold and Kriedman, 1964).

Table 2. Number of days from flower bud formation t0 75% anthesis

ENTRY	MEAN
<u>Stage of Application</u>	
1 month	61.487a
2 month	59.391a
3 month	62.215a
<u>Gibberellic Acid Concentration</u>	
C1 – 0 ppm/ control	68.300a
C2 – 50 ppm	57.467b
C3 – 100 ppm	56.056b
C4 – 250 ppm	61.000b
C5 - 500 ppm	62.333ab



3. Average Number of Leaves per Plant

Effect on the Stage of Application. Table 3 shows that the number of leaves per plant had significant differences. Plant sprayed after two month has the highest number of leaves with 15.556 leaves per plant, followed by one month after spraying with 14.327 leaves per plant. Three month from spaying had the least number of leaves with 13.313 per plant at 75% anthesis.

Effect of GA₃ Concentration. There is significant effect of the different GA₃ concentration on the number of leaves produce per plant at 75% anthesis. Plant sprayed with 100 ppm GA₃ concentration obtained the highest number of leaves per plant at 75% anthesis with 15.90, followed by 250 ppm and 50 ppm GA₃ concentration having 14.176 and 14.704, respectively. The lowest number of leaves counted was obtained from plants of untreated or control and 500 ppm GA₃ concentration which had 13.761 and 13.444 leaves per plant at 75% anthesis.

Interaction Effect. A significant difference was obtained between the stage of application and the different GA₃ concentration in terms on the number of leaves per plant at 75% anthesis. Plant sprayed with 100 ppm GA₃ concentration after two month produces the highest number of leaves with 18.223 per plant at 75% anthesis. The months that produce the least number of leaves were from three month treated with 500 ppm GA₃ concentration with 10.223 leaves per plant at 75% anthesis.

Mendoza 19993, stated that gibberellic acid at 500 ppm with two applications significantly enhance cutflower production, increase yield, higher number of leaves per plant at anthesis, longer flower buds and blooms higher leaf area index at flowering and promote longer vase life.



Table 3. Average number of leaves per plant

ENTRY	MEAN
<u>Stage of Application</u>	
1 month	14.327ab
2 month	15.556a
3 month	13.313b
<u>Gibberellic Acid Concentration</u>	
C1 – 0 ppm/ control	13.761b
C2 – 50 ppm	14.704ab
C3 – 100 ppm	15.908a
C4 – 250 ppm	14.176b
C5 - 500 ppm	13.444b

4. Final Height at Flowering (cm)

Effect on the Stage of Application. Result show a significant differences were obtained on the height of plant at 75% anthesis. Two months from spraying had the longest stems with 58.281 cm, followed by one month from spraying with 49.942 cm per plant. Three month from spraying had the shortest stem measured with 44.998 cm at 75% anthesis.

Effect of GA₃ Concentration. The different GA₃ concentration shows a significant effect on the height of milflores plants at 75% anthesis. Plants sprayed with 250 and 100 ppm GA₃ concentration produce longer stem length at 75% anthesis which had 57.394 and 55.924 cm per



plant. The shortest stem measured was obtained from untreated plants with 39.482 per plant at 75% anthesis.

Interaction Effect. A significant difference was obtained from the stage of application with the different GA₃ concentration on the height of milflores plants at 75% anthesis. Result show that 250 ppm GA₃ concentration which was sprayed two month after treatment has the longest stem length with an average of 71.500 cm. on the other hand, untreated plants at two month had the shortest stem length measured with 37.890 cm. it was observed further, that all plants treated with GA₃ concentration except for the stage of application had a longer stems compared to those which were not treated.

Table 4. Final height at flowering

ENTRY	MEAN
<u>Stage of Application</u>	
1 month	49.942b
2 month	58.281a
3 month	44.998c
<u>Gibberellic Acid Concentration</u>	
C1 – 0 ppm/ control	39.482d
C2 – 50 ppm	52.983b
C3 – 100 ppm	55.924a
C4 – 250 ppm	57.394a
C5 - 500 ppm	49.583c



Caluya and Imlan (1959) reported that GA₃ induce the rapid elongation of plants in general. Even as early as one week after first application, the treated plants were already taller than the untreated

5. Flower Diameter

Effect on the Stage of Application. Result shows significant differences were observe on the flower diameter of milflores plants at 75% anthesis. Two months from spraying had the widest and biggest flower diameter measured with 21.897 cm. then followed by three months from spraying with 18.106 cm flower diameter measured. One month from spraying have the smallest flower that bloom measured with 16.300 cm at 75% anthesis.

Effect of GA₃ Concentration. The different GA₃ concentration shows a significant effect on the flower diameter of milflores plants at 75% anthesis. Plants sprayed with 250 and 100 ppm GA₃ concentration obtained the biggest flower diameter measured with 19.699 cm at 75 % anthesis. Plant treated with 500 and 50 ppm GA₃ concentration has light significant differences on flower diameter with 18.849 and 18.726 am at 75% anthesis. Untreated plant obtained the smallest flower diameter measured with 17. 053 cm per plant at 75% anthesis.

Interaction Effect. A significant difference were observed between the difference GA₃ concentration and the stage of application in terms on stem diameter measured on milflores plants at 75% anthesis. Plants treated with 250 ppm GA₃ concentration at two month produce the widest flower diameter measured with 23.250 cm at 75% anthesis. And smallest flower diameter measured was from untreated plants at one month with 14.500 cm per plant at 75% anthesis.

Application of 1,250 ppm GA₃, at three leaf-stage and four-leaf stage gladiolus bolted earlier, produce bigger size corms, longer flower spikes at flowering with more and bigger flowers (Albis, 1995)



Table 5. Stem diameter (cm)

ENTRY	MEAN
<u>Stage of Application</u>	
1 month	16.331c
2 month	21.897a
3 month	18.106b
<u>Gibberellic Acid Concentration</u>	
C1 – 0 ppm/ control	17.053c
C2 – 50 ppm	18.726b
C3 – 100 ppm	19.530ab
C4 – 250 ppm	19.699a
C5 - 500 ppm	18.849ab

6. Stem Diameter (cm)

Effect on the Stage of Application. Significant differences were observed on the stem diameter of milflores plants at 75% anthesis. Milflores plants sprayed with GA₃ at two month have the biggest stem diameter measured with 1.186cm, followed by three month from spraying with 1.131 cm per plant at 75% anthesis. One month from spraying GA₃ have the smallest stem diameter measured with 0.724 cm per plant at 75% anthesis. Further more, because of the smallness of the stems of milflores plants treated at one month, it needs a supporting material (stick) so that the flower will not touch the ground.

Effect of GA₃ Concentration. A significant effect were obtained from the different GA₃ concentration with the stem diameter of milflores plants. Plans treated with 250 ppm GA₃



concentration have the biggest stem diameter measured with 1.128 cm at 75% anthesis. A significant difference was measured on 500 and 100 ppm GA₃ concentration with 1.084 and 1.030 cm per plant at the time of flowering. Untreated plants obtained the smallest stem diameter that has been measured with 0.550 cm at 75% anthesis.

Table 6. Flower diameter (cm)

ENTRY	MEAN
<u>Stage of Application</u>	
1 month	0.724b
2 month	1.186a
3 month	1.131a
<u>Gibberellic Acid Concentration</u>	
C1 – 0 ppm/ control	0.844a
C2 – 50 ppm	1.084ab
C3 – 100 ppm	1.030ab
C4 – 250 ppm	1.128a
C5 - 500 ppm	0.981b

Interaction Effect. A significant effect was observed between the different GA₃ concentration and the stage of application in terms of stem diameter at 75% anthesis. Milflores plants treated with 100 ppm GA₃ concentration at two month have the largest stem diameter ,



while untreated plants at one month have the smallest stem diameter measured with 0.550 cm at 75% anthesis.

7. Number of Lateral Stems

Effect on the Stage of Application. A significant difference was obtained on the number of lateral stems with the effect of the stage of application at 75% anthesis. Plants treated at one month have the largest number of lateral stems counted with 3.0 lateral stems per plant, while the smallest number of lateral stems counted was two month from treatment with 2.200 lateral stems per plant at 75% anthesis.

Effect of GA₃ Concentration. Spraying GA₃ concentration on potted milflores plants did not significantly affect the number of lateral stems produced per plant at 75% anthesis.

Interaction Effect. Table 7 shows a comparable significant effect between the different GA₃ concentration and the stage of application on the potted milflores plants. Milflores plants treated with 50 ppm at one month and 500 ppm GA₃ concentration obtained the highest number of lateral stems with 3.333 per plant at 75% anthesis. While untreated plants at two month produce the least number of lateral stems at 75% anthesis with 1.667 per plant.

Hermano and Ladilad (1980) found that the rate of growth effected by 400 ppm, four weeks from treatment. However, plant treated with the highest concentration (1000 ppm) at GA₃ finally showed the fastest rate of growth in the later part of the vegetative stage.



Table 7. Number of lateral stems

ENTRY	MEAN
<u>Stage of Application</u>	
1 month	3.000a
2 month	2.200b
3 month	2.533ab
<u>Gibberellic Acid Concentration</u>	
C1 – 0 ppm/ control	2.222a
C2 – 50 ppm	2.556a
C3 – 100 ppm	2.667a
C4 – 250 ppm	3.000a
C5 - 500 ppm	2.444a



SUMMARY, CONCLUSION AND RECOMMENDATION

Summary

The study was conducted at Lamtang, Puguis, La Trinidad, Benguet from November 2007 to March 2008 which aimed to compare the effect of GA₃ concentration on the vegetative growth and flowering of potted milflores plants; to determine the effect of different gibberellic acid GA₃ on the growth and flowering of milflores, and to determine the best concentration of gibberellic acid GA₃ that will enhance vegetative growth with better flower quality.

The earliest to develop flower buds observed were on milflores plant treated at three month and the longest flower bud to develop was one month from treatment. Plants sprayed with 50 ppm GA₃ were the earliest to develop flower buds while untreated plants required longer days to develop flower buds.

Milflores plants treated with smaller amount of GA₃ concentration was the earliest to develop flower followed by the higher amount of GA₃ concentration. Untreated plants show delayed flower development and flowering.

The highest number of leaves counted at 75% anthesis was produced by the plant treated at two months. Plant sprayed with 100 ppm GA₃ has the highest number of leaves counted followed by 50 ppm.

Milflores plants treated at two month produce the longest stem measured at 75% anthesis followed by one month and three month. Plant sprayed with 250 ppm GA₃ the longest stem measured followed by 100 ppm.

The widest flower diameter was produced by 250 ppm followed by 100 ppm GA₃ concentration at two month. The smallest flower diameter measured was observed in untreated plants.



The thickest stems were measured on milflores plant treated with 250 ppm and followed by plants sprayed with 50 and 100 ppm GA₃. Two month produce the thickest stems followed by three month and one month from treatment.

Plats treated at one month produce the largest number of leaves of lateral stems followed by three month and two month. GA₃ have a slight effect on the lateral stems of milflores plants.

Conclusion

Based on the study conducted, and on the data gathered, potted milflores plants treated at two month shows to have responded better on the different GA₃ concentration based on the observed effect on its vegetative growth, and flower quality produced per plant.

Recommendation

It is therefore recommended the use of gibberellic acid (GA₃) as a growth regulator for milflores plants. And a further study regarding non milflores plants with a higher concentration as a consideration on its growth development should be conducted.



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APPENDICES

Appendix Table 1. Days from transplanting to flower bud formation (0.5 cm bud size)

TREATMENT	REPLICATION			TOTAL	MEAN
	I	II	III		
S1 C1	167	167	170	504.30	168.100
C2	152.25	155	159.75	467.00	155.667
C3	155.25	159	155	469.25	156.417
C4	160.5	163.67	162	486.17	162.057
C5	162	158.4	161.67	482.07	160.690
S2 C1	157.8	159.2	166.7	483.70	161.233
C2	151.8	151.8	158.2	461.80	153.933
C3	150.5	156.5	156	463.00	154.333
C4	159.8	156.8	156	472.60	157.533
C5	158	159.5	157	474.50	158.167
S3 C1	137	134	132	403.00	134.333
C2	122.4	123	125.4	370.80	123.600
C3	125.6	126.6	122.4	374.60	124.867
C4	126	124.4	125.6	376.00	125.333
C5	123	126.75	124.5	374.25	124.750
<hr/>					
Total					



Analysis of Variance

Source of Variation	Degrees of Freedom	Sum of Squares	mean of Squares	Computed F	TABULAR F	
					0.05	0.01
Factor A	2	10486.118	5243.059	788.19**	3.32	5.59
Factor B	4	574.500	143.625	21.59**	4.69	4.02
AB	8	64.873	8.109	1.22ns	2.27	5.17
Error	30	199.561	6.652			
Total	44	11325.052				

** = Highly Significant

Coefficient of variation= 40.49%

Ns = Not Significant



Appendix Table 2. Number of days from flower bud formation to 75% anthesis

TREATMENT	REPLICATION			TOTAL	MEAN
	I	II	III		
S ₁ C ₁	65	66	67	198.00	66.000
C ₂	55.8	50.33	56.5	162.63	54.467
C ₃	60	61.67	63.5	185.17	61.723
C ₄	61	60.5	63	184.50	61.500
C ₅	65	66.5	60.5	192.00	64.000
S ₂ C ₁	65	66.2	67.5	198.70	66.233
C ₂	55	57.75	57	162.75	56.583
C ₃	56.75	56.67	57.5	170.92	56.973
C ₄	57	59.5	58	174.50	58.167
C ₅	60	57.5	59.5	177.00	59.000
S ₃ C ₁	75.5	71.5	71	218.00	72.667
C ₂	60.66	62.66	61.5	184.82	61.607
C ₃	62	61.66	64.75	148.41	49.470
C ₄	62	64.4	63.6	190.00	63.333
C ₅	64	62	66	192.00	64.000
Total					



Analysis of Variance

Source of Variation	Degrees of Freedom	Sum of Squares	mean of Squares	Computed F	TABULAR F	
					0.05	0.01
Factor A	2	64.482	32.241	0.77ns	3.32	5.59
Factor B	4	827.954	206.989	1.94**	2.69	4.02
AB	8	427.155	53.394	1.27ns	2.27	5.17
Error	30	1257.256	41.909			
Total	44	2576.847				

** = Highly Significant

Coefficient of variation= 10.61%

Ns = Not Significant



Appendix Table 3. Average number of leaves per plant

TREATMENT	REPLICATION			TOTAL	MEAN
	I	II	III		
S ₁ C ₁	14	12	13	13.000	39.00
C ₂	11	17	13.25	13.750	41.25
C ₃	19	12	15	15.333	46.00
C ₄	14.33	13	14.5	13.943	41.83
C ₅	16	16.5	14.33	15.610	46.83
S ₂ C ₁	15.25	13	14.25	14.167	42.50
C ₂	15	16.67	15	15.557	46.67
C ₃	18	16.67	20	18.223	54.67
C ₄	16	16	14	15.333	46.00
C ₅	13	14.5	16	14.500	43.50
S ₃ C ₁	15.75	14	12.6	14.117	42.35
C ₂	16	14.67	13.75	14.807	44.42
C ₃	15.75	12.5	14.25	14.067	42.50
C ₄	14.25	13.6	11.9	13.250	39.75
C ₅	11	8.67	11	10.223	30.67
Total					



Analysis of Variance

Source of Variation	Degrees of Freedom	Sum of Squares	mean of Squares	Computed F	TABULAR F	
					0.05	0.01
Factor A	2	37.859	18.929	6.64**	3.32	5.59
Factor B	4	33.639	8.410	2.95*	2.69	4.02
AB	8	51.156	6.395	2.24ns	2.27	5.17
Error	30	85.557	2.852			
Total	44	208.211				

** = Highly Significant

Coefficient of variation= 11.73%

*= Significant

Ns = Not Significant



Appendix Table 4. Final height at flowering (cm)

TREATMENT	REPLICATION			TOTAL	MEAN
	I	II	III		
S ₁ C ₁	48	33.5	37.5	119.00	39.667
C ₂	49.3	55	53.75	158.05	52.683
C ₃	51.33	53	53.33	157.66	52.553
C ₄	52.33	53.75	52.67	158.75	52.917
C ₅	50.67	51	54	155.67	51.890
S ₂ C ₁	37.5	39	37.12	113.62	37.873
C ₂	63	64.67	59	186.67	62.223
C ₃	72	62.67	71.5	206.17	68.723
C ₄	74.5	71.5	68.5	214.50	71.500
C ₅	51.25	51.5	50.5	153.25	51.083
S ₃ C ₁	40.62	39.87	42.6	122.72	40.907
C ₂	48.1	42.66	41.37	132.13	44.043
C ₃	45.12	47.25	47.12	139.49	46.497
C ₄	47.5	47.7	48.1	143.30	47.767
C ₅	44.5	48.33	44.5	137.33	45.777
Total					



Analysis of Variance

Source of Variation	Degrees of Freedom	Sum of Squares	mean of Squares	Computed F	TABULAR F	
					0.05	0.01
Factor A	2	1352.029	676.014	73.60**	3.32	5.59
Factor B	4	1833.406	458.352	49.90**	2.69	4.02
AB	8	951.595	111.949	12.95**	2.27	5.17
Error	30	275.543	9.185			
Total	44	4412.573				

** = Highly Significant

Coefficient of variation= 5.93%



Appendix Table 5. Flower diameter (cm)

TREATMENT	REPLICATION			TOTAL	MEAN
	I	II	III		
S ₁ C ₁	14	13.5	16	43.50	14.500
C ₂	15.1	16.83	14.25	46.18	15.393
C ₃	17.66	16	18	51.66	17.220
C ₄	18.33	16	18.5	52.83	17.610
C ₅	17.67	17.5	15.33	50.50	16.833
S ₂ C ₁	20.13	20	19.38	59.51	19.837
C ₂	21.5	22.67	22.17	66.34	22.113
C ₃	22.65	22.83	22.5	67.98	22.660
C ₄	23.75	22.5	23.5	69.75	23.250
C ₅	21.25	21.62	22	64.87	21.623
S ₃ C ₁	17.75	16.12	132	50.47	16.823
C ₂	18.7	19	125.4	56.01	18.670
C ₃	18.88	19.25	122.4	56.13	18.710
C ₄	18.31	17	125.6	54.71	18.237
C ₅	18.63	17.17	124.5	54.27	18.090
Total					



Analysis of Variance

Source of Variation	Degrees of Freedom	Sum of Squares	mean of Squares	Computed F	TABULAR F	
					0.05	0.01
Factor A	2	243.930	121.965	143.52**	3.32	5.59
Factor B	4	39.560	9.890	11.64**	4.69	4.02
AB	8	8.540	1.067	1.26ns	2.27	5.17
Error	30	25.494	0.850			
Total	44	317.523				

** = Highly Significant

Coefficient of variation= 4.91%

Ns = Not Significant



Appendix Table 6. Stem diameter (cm)

TREATMENT	REPLICATION			TOTAL	MEAN
	I	II	III		
S ₁ C ₁	.60	.45	.60	1.65	0.550
C ₂	.66	.70	.94	2.30	0.767
C ₃	.73	.50	.70	1.93	0.643
C ₄	.90	.70	.85	2.45	0.817
C ₅	.83	1.0	.70	2.53	0.843
S ₂ C ₁	.64	1.20	1.05	2.89	0.963
C ₂	1.50	1.47	1.40	4.37	1.457
C ₃	1.40	1.33	1.30	4.03	1.343
C ₄	1.35	1.30	1.15	3.80	1.267
C ₅	.70	.90	1.1	2.70	0.900
S ₃ C ₁	.875	1.10	1.08	3.06	1.020
C ₂	1.20	.97	.92	3.09	1.030
C ₃	1.01	1.15	1.15	3.31	1.103
C ₄	1.20	1.28	1.42	3.90	1.300
C ₅	1.20	1.10	1.30	3.60	1.200
Total					



Analysis of Variance

Source of Variation	Degrees of Freedom	Sum of Squares	mean of Squares	Computed F	TABULAR F	
					0.05	0.01
Factor A	2	1.909	0.955	50.12**	3.32	5.59
Factor B	4	0.432	0.108	5.67**	4.69	4.02
AB	8	0.630	0.079	4.13*	2.27	5.17
Error	30	0.571	0.019			
Total	44	3.543				

** = Highly Significant

Coefficient of variation= 13.62%

* = Significant



Appendix Table 7. Number of lateral stems

TREATMENT	REPLICATION			TOTAL	MEAN
	I	II	III		
S ₁ C ₁	3	2	3	8	2.667
C ₂	4	3	3	10	3.333
C ₃	3	2	4	9	3.000
C ₄	2	4	3	9	3.000
C ₅	3	3	3	9	3.000
S ₂ C ₁	1	2	2	5	1.667
C ₂	2	2	2	6	2.000
C ₃	3	2	3	8	2.667
C ₄	2	3	3	8	2.667
C ₅	1	3	2	6	2.000
S ₃ C ₁	1	3	3	7	2.333
C ₂	2	2	3	7	2.333
C ₃	3	2	2	7	2.333
C ₄	3	3	4	10	3.333
C ₅	2	2	3	7	2.333
Total					



Analysis of Variance

Source of Variation	Degrees of Freedom	Sum of Squares	mean of Squares	Computed F	TABULAR F	
					0.05	0.01
Factor A	2	4.844	2.422	4.95*	3.32	5.59
Factor B	4	2.978	0.744	1.52ns	4.69	4.02
AB	8	2.489	0.311	0.64ns	2.27	5.17
Error	30	14.667	0.489			
Total	44	24.978				

* = Significant

Coefficient of variation= 27.12%

Ns = Not Significant

