



## Research Article

# Optimization of stage of gibberellin spray in Cape gooseberry (*Physalis peruviana* L.) for the improvement of yield and fruit quality under subtropical condition

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## Abstract

The current experiment was conducted to optimize the actual time of gibberellin (GA<sub>3</sub>) spray in cape-gooseberry (*Physalis peruviana* L.) to improve growth, yield, and fruit quality traits. Experimental observations revealed that physiological growth (leaf relative water content, specific leaf weight) was improved significantly by three repeated applications of GA<sub>3</sub> @ 100 ppm each at vegetative, anthesis, and again at the fruit setting stage. GA<sub>3</sub> spray @ 25 ppm each at all three growth stages had the longest flowering period; although the harvest span was calculated maximum in three repeated GA<sub>3</sub> sprays @ 100 ppm each. Three applications of GA<sub>3</sub> @ 25 ppm each also increased the fruit yield at its maximum level (31.79 t ha<sup>-1</sup>). Three foliar spray of GA<sub>3</sub> at any concentration increased the TSS significantly over the respective control; however, the acidity was reduced at the maximum level when GA<sub>3</sub> was applied @ 50-100 ppm each at all three growth stages. But reducing and non-reducing sugar content was maximum under GA<sub>3</sub> spray @ 100 ppm each during all three growth stages (8.33 and 2.70%, respectively) with statistically at par results in three applications of GA<sub>3</sub> @ 25-75 ppm each. Hence, it can be concluded that GA<sub>3</sub> application @ 25 ppm each at vegetative, anthesis, and again at fruit setting stage is optimum to improve the yield of better quality cape gooseberry fruit significantly.

**Keywords** cape gooseberry, fruit quality, gibberellins, physiological growth, yield

## Introduction

Cape gooseberry (*Physalis peruviana* L.) is one of the most important nutritious fruit crops of the Solanaceae family. It is a good source of vitamin A (2380 IU 100 g<sup>-1</sup>), pectin (0.9%), minerals (calcium, phosphorus), and fibers which provide considerable health benefits. Various bioactive compounds (with anilides and phenolics) are also reported to be present in cape gooseberry. Keeping these nutritional benefits, the demand for this crop has increased many folds during the last two decades. To meet this demand, the area and production of the crop have increased significantly in the country. Although, it is a crop of tropical climate, its wide adaptability to different soil and climatic conditions, makes it suitable crop to grow in the wide subtropical region of the country too. Hence, growers from different patches of subtropics are now started to cultivate cape gooseberry commercially. But the key problem of growing this crop under subtropical climate is the low yield potentiality (only 400 – 500 g plant<sup>-1</sup> as compared to 700-900 g plant<sup>-1</sup> in

Received: 07 November 2022  
Accepted: 08 December 2022  
Online: 09 December 2022

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Emer Life Sci Res (2022) 8(2): 206-213

E-ISSN: 2395-6658  
P-ISSN: 2395-664X

DOI: <https://doi.org/10.31783/elsr.2022.82206213>



leading cape gooseberry producing countries) with poor keeping quality. This is mainly because of poor soil and nutrient management as well as a lack of technical knowledge among the growers. A large number of well-established low-cost production technologies have been reported throughout the world to increase the quality as well as production of different crops. Application of recommended dose of nutrients, adaptation of high-density planting system, use of micro-irrigation, application of plant growth regulators (PGRs) and biofertilizers play a significant role to improve the production as well as keeping quality of different fruit crops [1-6]. Among them, the foliar application of PGRs has emerged as an effective technique to stimulate fruit growth and quality. Getting higher yield with better fruit quality by the application of PGRs has also been reported earlier in different fruit crops [7-9].

Among different PGRs, gibberellic acid ( $GA_3$ ) plays a crucial role in plant growth and development. When  $GA_3$  was applied exogenously, it accelerated the rates of cell division and cell elongation at the sub-apical meristematic region resulting in improved physiological growth with better quality fruits [9]. Kumar et al., [10] reported that the foliar application of increased concentration of gibberellins had a significant impact on the improvement of the fruit quality of cape gooseberry. But the high cost of this hormone makes it difficult for fruit growers to use it in too large quantity. It enforces researchers worldwide to optimize the precise dose of  $GA_3$  for its application. Further, no literature is available regarding the exact plant growth stages to apply  $GA_3$  for enhancing yield with improved fruit quality. Hence, it is also necessary to optimize the time of  $GA_3$  spray on cape gooseberry plants to enhance yield potentiality with improvement in keeping quality. Therefore, the current experiment was formulated to study the impact of different concentrations of  $GA_3$  spray (at different growth stages) on the growth, yield, and fruit quality of cape gooseberry, particularly in the subtropical region.

## Methodology

### Materials

The local cultivar of cape gooseberry (*Physalis peruviana* L.) was used as experimental material. Healthy cape gooseberry seedlings with uniform growth and free from any pest and disease injuries were selected for transplanting in the main experimental plot.

### Methods

$GA_3$  was sprayed on the experimental plants at 25, 50, 75, and 100 ppm only at the vegetative, anthesis, and fruit setting stages and also all three growth stages together. Double distilled water was sprayed on the control plants. The experiment was carried out for two successive growing seasons.

### Physiological and reproductive growth

Specific leaf weight was also estimated manually at the fruiting stage [11]. The duration for planting to the anthesis of 1<sup>st</sup> flower and the anthesis of 1<sup>st</sup> flower to 1<sup>st</sup> fruit setting were also recorded manually. Thereafter, the total flowering span was calculated by recording the date of anthesis of 1<sup>st</sup> flower to the last one. Similarly, the duration of fruit set to maturity was recorded manually. Further, the harvesting span was calculated by counting the duration between the harvesting of 1<sup>st</sup> fruit and to last one.

### Yield and fruit quality traits

Yield per plant was measured manually and yield hectare<sup>-1</sup> was calculated following the formula:

$$\text{Yield ha}^{-1} = \text{Yield plant}^{-1} \times \text{No. of plants accommodated in one hectare (37037)}.$$

Total soluble solids of the ripped fruits were measured through a hand refractometer (Make: Atago, Tokyo, Japan). Titratable acidity was estimated through the titration method [12]. Reducing sugar content was determined by the method described by Lane and Eynone [13], while the total carotenoid content of the fruit was estimated following the protocol described by Roy [14].

### Statistical analysis

The entire experiment was laid on a completely randomized block design (CRBD) having three replications. Pool data for two successive seasons were analyzed by statistical analysis software (SAS 9.3; SAS Institute, Cary, NC, USA) and means were differentiated through Duncan's multiple range test (DMRT).

## Results and Discussion

### Physiological and reproductive growth

The specific leaf weight of the cape gooseberry plant was influenced significantly in all the GA<sub>3</sub> treatments (Figure 1). It was recorded maximum when GA<sub>3</sub> was applied at all three growth stages (32.25 mg cm<sup>-2</sup>) and increased gradually with the increasing concentration of GA<sub>3</sub> with maximum at 100 ppm concentration (39.00% higher than the control). However, the interaction showed that three repeated applications of GA<sub>3</sub> @ 100 ppm each at vegetative, anthesis, and again at fruiting stage had the highest specific leaf weight (43.30% higher than the control).

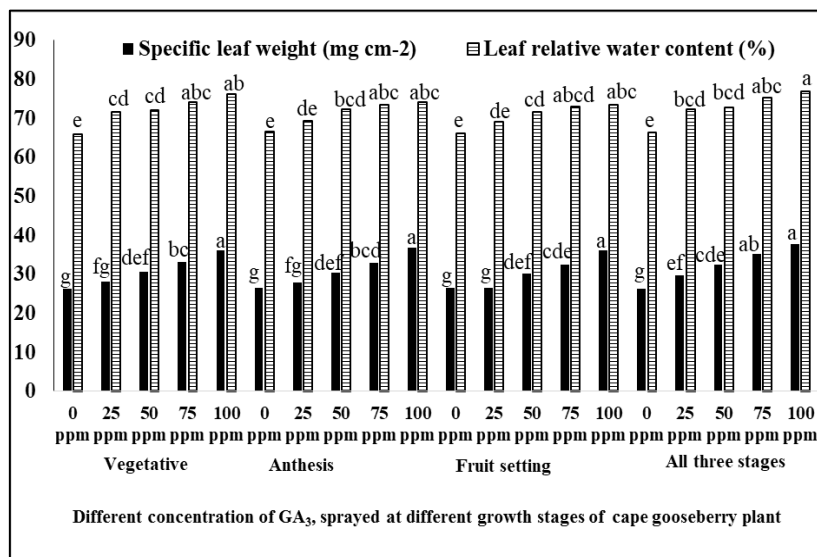


Figure 1. Effect of foliar application of GA<sub>3</sub> on specific leaf weight and leaf relative water content of cape gooseberry (*Physalis peruviana* L.) plants

Similarly, 100 ppm GA<sub>3</sub> spray each at all three growth stages had maximum leaf relative water content too (76.87%) which was statistically at par with three applications @ 75 ppm each (75.26%) (Figure 1). Increased concentration of GA<sub>3</sub> within the plant system helps to synthesize several amino acids. These amino acids are subsequently incorporated in proteins and nucleic acid and ultimately provide the framework for chloroplast and other photosynthetic elements resulting in improved physiological functions within the plant system [15]. Further, regardless of concentration, foliar application of GA<sub>3</sub> repeatedly for three times during vegetative, anthesis, and again at fruit setting stage had its long-lasting effect resulting in higher physiological growth as compared to a single application. Duration from planting to anthesis of 1<sup>st</sup> flower indicates a significant variation among the treatments (Figure 2). A single application of GA<sub>3</sub> @ 25 ppm during the vegetative stage helped to induce flowering in cape gooseberry at the earliest with par result in its single application (25 ppm) at the fruit setting stage and its application (25 ppm) at all three growth stages.

Similarly, the longest flowering span was calculated in the treatment comprising three repeated applications of 25 ppm GA<sub>3</sub> each at vegetative, anthesis, and again at fruiting stage (18.00 extra days as compared to respective control) which was statistically at par with 25 ppm GA<sub>3</sub> spray only at vegetative stage. However, the smallest flowering duration was recorded in three repeated sprays of GA<sub>3</sub> at all three growth stages @ 100 ppm each (Figure 2).

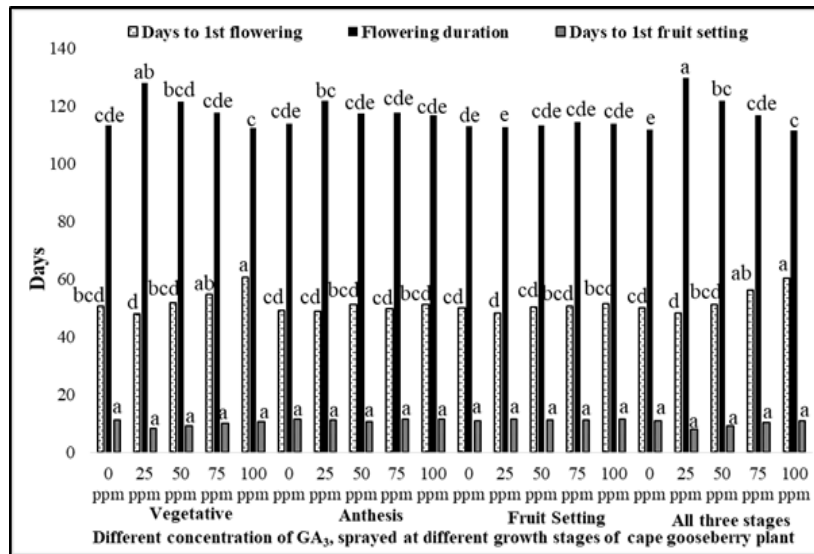


Figure 2. Effect of foliar application of GA<sub>3</sub> on flowering and fruiting behaviour of cape gooseberry (*Physalis peruviana* L.) plants

Further, the earliest fruit setting was recorded in three repeated applications of GA<sub>3</sub> @ 25 ppm each (3.00 days earlier than the respective control) (Figure 2). Reproductive growth of cape gooseberry plants in terms of precocity in flowering, flowering duration, and precocity in fruit setting has improved significantly in three repeated applications of GA<sub>3</sub> @ 25 ppm with at par result in 50 ppm spray. However, a concentration beyond 50 ppm was found ineffective for improving the reproductive growth of cape gooseberry plants. It might be due to the early completion of the vegetative phase in the plants, sprayed with either 25 or 50 ppm GA<sub>3</sub> as compared to its higher concentrations [9].

The duration of fruit set to maturity also varied significantly under different GA<sub>3</sub> treatments (Figure 3). A single application of GA<sub>3</sub> @ 25 ppm at vegetative stage had the least duration from fruit set to maturity (51 days). However, the duration from fruit set to maturity increased gradually with the increasing concentration of GA<sub>3</sub>, and it was recorded maximum when GA<sub>3</sub> was sprayed at all three growth stages @ 100 ppm each (63.33 days). Similarly, repeated application of GA<sub>3</sub> @ 100 ppm each at all three growth stages had the largest harvest span too (143.00 days) with at par duration in single spray @ 100 ppm at fruit setting stage only (142.33 days) and anthesis stage only (142.00 days) (Figure 3).

### Yield and fruit quality traits

Average fruit weight was also increased gradually with the increasing GA<sub>3</sub> concentration, irrespective of stages of application with the maximum in 100 ppm spray (35.10% larger than the control) (Table 1). Further, regardless of concentration, maximum fruit weight (9.94 g) was estimated in the plants that received three sprays of GA<sub>3</sub> (vegetative, anthesis, and fruiting stages) with par value in single GA<sub>3</sub> treatment at fruit setting stage (9.77 g). The interaction revealed that a single application of 100 ppm GA<sub>3</sub> at the fruit setting stage had the highest fruit weight (40.78% higher than the respective control) with per result in three repeated applications of 100 ppm GA<sub>3</sub> (38.48% higher than the respective control). Similarly, irrespective of concentration, yield ha<sup>-1</sup> was increased significantly when GA<sub>3</sub> was applied at all three growth stages (28.82 tonnes ha<sup>-1</sup>). A shape increase in yield was obtained with the increase of GA<sub>3</sub> concentration and it was obtained maximum in 100 ppm spray (28.34 tonnes ha<sup>-1</sup>) with non-significant variation in 75 and 50 ppm spray (28.09 and 28.01 tonnes ha<sup>-1</sup>, respectively). The interaction indicates that 25 ppm GA<sub>3</sub> applications each at three growth stages had maximum fruit yield ha<sup>-1</sup> (59.19% higher than the respective control) with par value in three repeated applications at either 50 or 75 ppm. The yield of any crop mainly depends on the physiological growth of the plant. The plants having improved physiological growth viz. relative water content of leaf, specific leaf weight would produce more photosynthetic

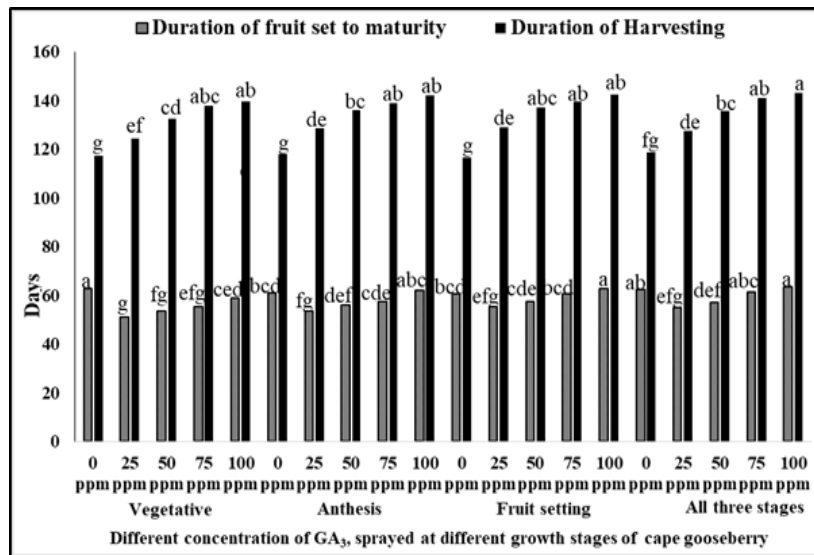


Figure 3. Effect of foliar application of GA<sub>3</sub> on duration of fruit set to maturity and harvesting span of cape gooseberry (*Physalis peruviana* L.) plants

Table 1. Effect of gibberellins (GA<sub>3</sub>) on yield and biochemical attributes of cape gooseberry (*Physalis peruviana* L.)

Treatment	Fruit weight (g)	Yield (t ha <sup>-1</sup> )	TSS (°B)	Titratable acidity (%)	Carotenoid content (µg g <sup>-1</sup> FW)	
<b>Stage of application (S)</b>						
Vegetative	9.08 <sup>b</sup>	26.66 <sup>b</sup>	15.28 <sup>a</sup>	0.77 <sup>a</sup>	49.12 <sup>b</sup>	
Anthesis	9.27 <sup>b</sup>	23.95 <sup>c</sup>	15.36 <sup>a</sup>	0.77 <sup>a</sup>	48.96 <sup>b</sup>	
Fruit setting	9.77 <sup>a</sup>	25.29 <sup>b</sup>	15.40 <sup>a</sup>	0.75 <sup>ab</sup>	50.36 <sup>b</sup>	
Vegetative, anthesis and fruit setting	9.94 <sup>a</sup>	28.82 <sup>a</sup>	15.41 <sup>a</sup>	0.74 <sup>b</sup>	51.80 <sup>a</sup>	
<b>Concentration of gibberellins (C)</b>						
0 ppm	7.72 <sup>d</sup>	19.73 <sup>c</sup>	13.95 <sup>a</sup>	0.91 <sup>a</sup>	40.97 <sup>c</sup>	
25 ppm	9.34 <sup>c</sup>	26.72 <sup>b</sup>	15.59 <sup>a</sup>	0.77 <sup>b</sup>	51.02 <sup>b</sup>	
50 ppm	9.93 <sup>b</sup>	28.01 <sup>a</sup>	15.71 <sup>a</sup>	0.73 <sup>c</sup>	52.03 <sup>ab</sup>	
75 ppm	10.16 <sup>ab</sup>	28.09 <sup>a</sup>	15.77 <sup>a</sup>	0.70 <sup>d</sup>	53.01 <sup>a</sup>	
100 ppm	10.43 <sup>a</sup>	28.34 <sup>a</sup>	15.79 <sup>a</sup>	0.68 <sup>e</sup>	53.28 <sup>a</sup>	
<b>S × C Interaction</b>						
Vegetative	0 ppm	7.61 <sup>h</sup>	19.44 <sup>g</sup>	0.93 <sup>a</sup>	14.67 <sup>d</sup>	40.95 <sup>d</sup>
	25 ppm	8.67 <sup>e</sup>	26.45 <sup>de</sup>	0.78 <sup>bc</sup>	20.01 <sup>c</sup>	49.13 <sup>c</sup>
	50 ppm	9.53 <sup>ef</sup>	28.97 <sup>bc</sup>	0.73 <sup>cde</sup>	21.58 <sup>abc</sup>	50.94 <sup>bc</sup>
	75 ppm	9.63 <sup>ef</sup>	28.86 <sup>bc</sup>	0.72 <sup>def</sup>	21.78 <sup>abc</sup>	52.11 <sup>abc</sup>
	100 ppm	9.94 <sup>cde</sup>	29.55 <sup>ab</sup>	0.70 <sup>efgh</sup>	22.71 <sup>ab</sup>	52.48 <sup>abc</sup>
Anthesis	0 ppm	7.80 <sup>h</sup>	19.64 <sup>g</sup>	0.90 <sup>a</sup>	15.72 <sup>d</sup>	40.58 <sup>d</sup>
	25 ppm	8.92 <sup>fg</sup>	23.24 <sup>f</sup>	0.79 <sup>b</sup>	19.81 <sup>c</sup>	49.39 <sup>c</sup>
	50 ppm	9.63 <sup>ef</sup>	25.31 <sup>ef</sup>	0.74 <sup>bcde</sup>	21.21 <sup>bc</sup>	50.78 <sup>bc</sup>
	75 ppm	9.81 <sup>de</sup>	23.43 <sup>efg</sup>	0.71 <sup>efg</sup>	22.08 <sup>abc</sup>	51.78 <sup>abc</sup>
	100 ppm	10.17 <sup>abcde</sup>	26.11 <sup>de</sup>	0.69 <sup>efgh</sup>	22.97 <sup>ab</sup>	52.28 <sup>abc</sup>
Fruit setting	0 ppm	7.70 <sup>h</sup>	19.85 <sup>g</sup>	0.91 <sup>a</sup>	15.51 <sup>d</sup>	41.25 <sup>d</sup>
	25 ppm	9.75 <sup>de</sup>	25.39 <sup>ef</sup>	0.77 <sup>bcd</sup>	20.20 <sup>c</sup>	51.39 <sup>abc</sup>
	50 ppm	10.10 <sup>abcde</sup>	26.06 <sup>de</sup>	0.73 <sup>cde</sup>	21.61 <sup>abc</sup>	52.11 <sup>abc</sup>
	75 ppm	10.48 <sup>abcd</sup>	27.29 <sup>cd</sup>	0.69 <sup>efgh</sup>	22.78 <sup>ab</sup>	53.45 <sup>ab</sup>
	100 ppm	10.84 <sup>a</sup>	27.83 <sup>cd</sup>	0.66 <sup>h</sup>	24.06 <sup>a</sup>	53.62 <sup>ab</sup>
Vegetative, anthesis and fruit setting	0 ppm	7.77 <sup>h</sup>	19.97 <sup>g</sup>	0.92 <sup>a</sup>	15.26 <sup>d</sup>	41.08 <sup>d</sup>
	25 ppm	10.02 <sup>bcde</sup>	31.79 <sup>a</sup>	0.73 <sup>cde</sup>	21.44 <sup>bc</sup>	54.16 <sup>ab</sup>
	50 ppm	10.43 <sup>abcd</sup>	31.71 <sup>a</sup>	0.72 <sup>efg</sup>	21.99 <sup>abc</sup>	54.28 <sup>ab</sup>
	75 ppm	10.67 <sup>abc</sup>	30.78 <sup>a</sup>	0.67 <sup>fgh</sup>	23.48 <sup>ab</sup>	54.71 <sup>a</sup>
	100 ppm	10.76 <sup>ab</sup>	29.86 <sup>ab</sup>	0.67 <sup>gh</sup>	23.67 <sup>ab</sup>	54.75 <sup>a</sup>

Value indicates mean of three replicates. Different letters in the same column indicate significant differences at P ≤ 0.05 (Duncan's Multiple Range Test).

assimilates which in turn improve the yield of the crop. In addition, increased accumulation of carbohydrates directly increased the fruit weight which ultimately improves the crop yield. Earlier findings of Singh et al., [9] in cape gooseberry also confirmed similar results. The total soluble solid (TSS) of ripped cape gooseberry fruits has improved significantly in all GA<sub>3</sub>-treated plants (Table 1). Application of GA<sub>3</sub> at either concentration but at all three growth stages increased the TSS content significantly as compared to the respective control. The improvement of TSS in three applications of GA<sub>3</sub> might be due to the higher accumulation of sugars and other soluble compounds, particularly through protein hydrolysis and ascorbic acid oxidation [16]. However, the reverse pattern was observed for titratable acidity and it was reduced significantly when GA<sub>3</sub> was applied at all three growth stages (0.74%). The lowest acidity in three repeated applications of GA<sub>3</sub> might be due to the dilution effect and increased conversion of fruit acidity to sugar and other solids as observed by Kaur and Kaur [17] in cape gooseberry. The amount of reducing along with non-reducing sugar content in ripped cape gooseberry fruits was not varied significantly with different stages of application (Figure 4).

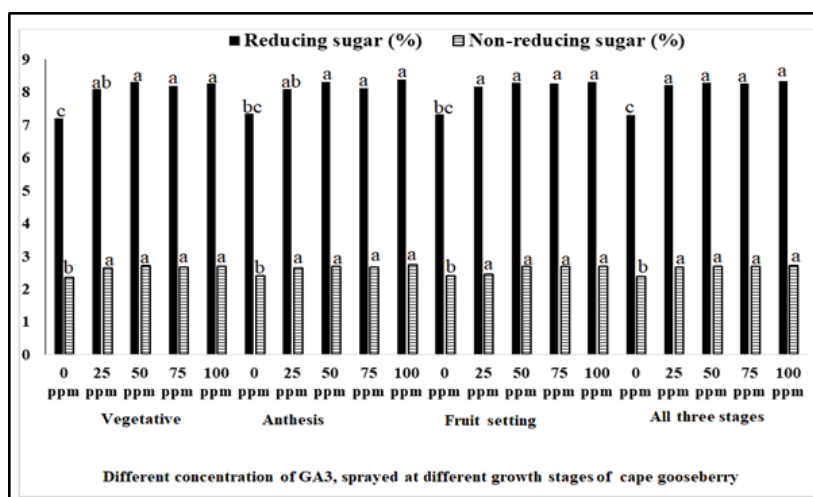


Figure 4. Effect of foliar application of GA<sub>3</sub> on sugar content of Ripped cape gooseberry (*Physalis peruviana* L.) fruits

However, reducing as well as non-reducing sugar content was estimated maximum in the plants that received three repeated applications of GA<sub>3</sub> (8.07 and 2.62, respectively). The interaction table revealed that the reducing sugar content was estimated maximum under a single GA<sub>3</sub> spray @ 100 ppm at anthesis (8.38%) which was statistically at par with its application during all three growth stages @ 25-100 ppm (8.21 - 8.33%). A similar pattern was also observed for non-reducing sugar content in ripped cape gooseberry fruits. Increased accumulation of reducing and non-reducing sugars by GA<sub>3</sub> spray might be due to the faster conversion of starch and acids into simple sugars [18]. Further application of GA<sub>3</sub> @ 25-100 ppm each at all three growth stages also helped to increase the carotenoid content to its maximum level in the fruit (54.16 – 54.75  $\mu\text{g g}^{-1}$  FW) with a statistically non-significant variation. Present research work confirms that repeated application of GA<sub>3</sub> improved the physiological growth of cape gooseberry significantly with improved reproductive behavior, resulting in maximum yield.

Fruit quality traits in terms of TSS, sugar, and carotenoid content of ripped cape gooseberry fruit was also improved significantly in three repeated application of GA<sub>3</sub>. Among four different concentrations, improvement in yield and quality attributes was statistically at par in 25- 100 ppm concentration. Hence, it can be concluded that the application of GA<sub>3</sub> @ 25 ppm each at vegetative, anthesis, and again at fruit setting stage is optimum to improve the yield of better quality cape gooseberry fruit significantly under subtropical conditions.



## Acknowledgements

Financial support from PG Research Contingency, Bihar Agricultural University, BAU, Sabour, Bhagalpur, India, is thankfully acknowledged

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