

Original Research Article

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Effect of Plant Growth Regulators and their Methods of Application on Growth of *kharif* Onion (*Allium cepa* L.) cv Agrifound Dark Red

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ABSTRACT

An experiment was conducted in field of the nursery, Department of Horticulture, College of Agriculture, Rajmata Vijayaraje Scindia Krishi Vishwa Vidyalaya, Indore during *Kharif* season, 2017-2018 to see the effect of plant growth regulators and their methods of application on growth of *kharif* onion (*Allium cepa* L.) cv. Agrifound Dark Red during 2017 and result have shown significant differences among the treatments. The maximum (48.03) plant height and number of leaves, leaf length (41.53 cm), leaf width (1.63 cm), leaf area (428.53 cm²), pseudostem length (9.03 cm), fresh weight of plant (57.43g), dry weight of plant (12.11 g) were recorded under T₃ (GA₃ @ 100 ppm-foliar spray) at 80 days after transplanting (DAT). The maximum polar diameter (5.77 cm) and equatorial diameter (5.91cm) of onion bulb were also exhibited in the treatment T₃ (GA₃ @ 100ppm-foliar spray) and minimum neck thickness (1.18 cm) was recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray).

Keywords

PGRs, DAT,
Growth parameters,
Diameter,
Pseudostem length,
Neck thick

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Introduction

Onion is one of the most important bulbous vegetable crops grown all over the world. Onion (*Allium cepa* L.) belongs to the family *Amaryllidaceae* (*Alliaceae*) and locally known as Pyaj. It is an old world crop and it was domesticated in Iran and Pakistan i.e. Central Asia. The onion crop is an export oriented crop earning valuable foreign exchange for the

country. The demand for onion is worldwide. Onions are found in most of the markets of the world throughout the year and can be grown under wide range of Agro-climatic conditions. Irrespective of price, the demand remains almost constant in the market as it is primarily, used as seasoning for a wide variety of dishes in many homes. The crop export is done mainly to Malaysia, Singapore, Philippines, Indonesia, Gulf countries and Pakistan. Onion accounts for 70 percent of our

total foreign exchange earnings from the export of fresh vegetables. India is next to China in area and production of onion. Among the different states Maharashtra is leading state in terms of area and production. Other major onion states are Gujarat, Karnataka, Odisha, Uttar Pradesh, Andhra Pradesh, Tamil Nadu and Rajasthan. The area of onion is 1270.4 thousand hectare, total production is 21563.9 thousand metric tonnes and productivity is about 17.0 metric tonnes per hectare in India (Anonymous, 2017a). The area of onion production in Madhya Pradesh is 118.20 thousand hectares. Total production is 2848.0 thousand metric tonnes and productivity is about 24.09 metric tonnes per hectare (Anonymous, 2017b). Onion accounts for 310650.09 lakhs foreign exchange earnings from the export to different countries (Anonymous, 2017c). Government of India has declared onion as an essential commodity.

The pungency in onion is due to sulphur-bearing compound which is present in very small quantity (about 0.005%) in the form of volatile oil allyl propyl disulphides. The colour of the outer skin of onion bulbs is due to quercetin. It is consumed as a vegetable and condiment. The green leaves, immature and mature bulbs are eaten raw or used in vegetable preparations. It is an indispensable item in every kitchen and used to enhance flavour of different recipes. Onion has many medicinal values and used for preparation of various Homeopathic, Unani and Ayurvedic medicines. Phenolic compounds can offer significant anti-mellitus atherogenic protection by inhibiting the oxidation of low density lipoproteins (LDLs) (Scalbert *et al.*, 2005).

Onions are grown in three seasons, rabi, kharif and late kharif. For maintaining steady supply in the market, kharif crop of onion plays a major role. The production of *kharif* onion has several advantages i.e. increases total production per annum and fulfils the demand

of fresh onion in the market. *Kharif* onion provides high price as compared to *Rabi* season onion. The excessive vegetative growth is a problem in *kharif* onion. The plant height goes up to one meter and neck of the plant become thick, while, the bulb remains small. This is due to poor translocation of assimilates from leaves to bulbs. This translocation of food materials or for altering source to sink relationship is changed by application of plant growth regulators. The positive effect of plant growth regulators on horticultural crops have been shown by many workers (Lal *et al.*, 2013, Lal and Das, 2017, Jain *et al.*, 2017, Tameshwar *et al.*, 2017). The vegetative growth of *kharif* onion as represented by plant height, number of leaves per plant, fresh and dry weight of plant, increased to optimum level using GA₃ and NAA. CCC is very effective in inducing hardening of seedlings and increased growth of root and shoot. TIBA is antiauxins which produced male sterility, and reduce the incidence of *Fusarium* wilt. Therefore, the present investigation “Effect of plant growth regulators and their application methods on growth of *kharif* onion (*Allium cepa* L.) cv. Agrifound Dark Red” was carried out.

Materials and Methods

An experiment was conducted in field of the nursery, Department of Horticulture, College of Agriculture, Rajmata Vijayaraje Scindia Krishi Vishwa Vidyalaya, Indore during *Kharif* season, 2017-2018 to see the effect of plant growth regulators and their methods of application on growth of *kharif* onion (*Allium cepa* L.) cv. Agrifound Dark Red during 2017 and result have shown significant differences among the treatments. Plant height and number of leaves, leaf length, leaf width, leaf area, pseudostem length, fresh weight of plant, dry weight of plant were recorded on 20, 40, 60 and 80 DAT. Polar and

equatorial diameter, and neck thickness of bulb was recorded from vernier calliper at harvest. Analysis of variance (ANOVA) was used to test for differences among the treatments.

Results and Discussion

Plant Height

The plant height and number of leaves per plant increased significantly with the increasing crop growth period. At 20 days after transplantation (DAT), the maximum (17.67cm) plant height was recorded in T₃ (GA₃ @ 100 ppm-foliar spray), followed by T₆ (NAA @ 100 ppm-foliar spray) (16.47 cm), While, the minimum (13.87 cm) plant height was observed under control. After 40 DAT, the maximum (23.96cm) plant height was recorded in T₃ (GA₃ @ 100ppm-foliar spray), followed by T₆ (NAA @ 100ppm-foliar spray) (23.57 cm), While, the minimum (21.93 cm) plant height was found in control. In case of 60 DAT, treatment T₃ (GA₃ @ 100 ppm-foliar spray), T₆ (NAA @ 100 ppm-foliar spray) and T₂ (GA₃ @ 100 ppm- seedling dip.) were observed significantly higher (30.10 cm), (30.00 cm) and (29.27 cm) plant height, respectively. However, lowest plant height (25.70 cm) was recorded under control.

At 80 DAT, significantly maximum (48.03 and 47.93 cm) plant height, were observed under treatment T₃ (GA₃ @ 100ppm-foliar spray) and T₆ (NAA @ 100ppm-foliar spray), respectively and which were at par with each other. However, it was recorded lowest (42.07 cm) in control. Similarly, number of leaves per plant of onion increased significantly with the increasing crop growth period and T₃ (GA₃ @ 100ppm-foliar spray) was found to be the best treatments for maximum number of leaves at all stages of observations.

Maximum plant height was observed under treatment T₃ (GA₃ @ 100ppm-foliar spray). However, it was recorded lowest in treatment

T₁ (control). Plant height is a genetically controlled character but several studies have indicated that the plant height can be either increased or decreased by the application of synthetic plant growth regulators. The increase in plant height by foliar spray of GA₃ 100ppm and NAA 100 ppm might be due to rapid increase in cell division and cell elongation in the meristemic region. However significant reduction in these characters can be seen in the growth retardant treatments such as TIBA and cycocel in all the stages of growth. The mechanism of reduction in such traits due to application of growth retardants appears to be due to slowing down of cell division and reduction in cell expansion. It has been suggested that, TIBA and cycocel are anti-gibberellin dwarfing agents, leading to a deficiency of gibberellin in the plant and reduce the growth by blocking the conversion of geranyl pyrophosphate to copalyl pyrophosphate which is the first step of gibberellin synthesis. Thus, reduction in plant height is due to retardation of transverse cell division particularly in cambium which is the zone of meristematic activity at the base of the internodes. These results are in close conformity with those of Suheela *et al.*, (2005), Islam *et al.*, (2007), Bose *et al.*, (2009), Rashid (2010), Patel *et al.*, (2010a), Patel *et al.*, (2010b), Ouzounidou *et al.*, (2011), Govind *et al.*, (2015), Shashi Kumar and Shashidhar (2016) and Thakur *et al.*, (2018).

Maximum number of leaves plant⁻¹ was observed under treatment T₃ (GA₃ @ 100ppm-foliar spray). In general, leaf is considered as an important functional unit of plant which contributes to yield. Probable reason may be due to the role of these materials in enhancing cell division activity, increasing of proline accumulation of plant and increasing of endogenous phyto hormones i.e. increasing promotion hormones (IAA, GA₃ and cytokinins) and reducing ABA content which found that bio-regulators make

a shift in hormonal balance characterized by increasing in endogenous phyto hormone in plant. Similar results were also obtained by Suheela *et al.*, (2005), Islam *et al.*, (2007), Bose *et al.*, (2009), Rashid (2010), Patel *et al.*, (2010a), Patel *et al.*, (2010b), Govind *et al.*, (2015) and Thakur *et al.*, (2018).

Leaf length (cm)

The leaf length and width of various treatments of onion is given in Table 2 at 20, 40, 60 and 80 days after transplantation. The maximum (15.60 cm) leaf length was registered in T₃ (GA₃ @ 100ppm-foliar spray) as compared to control (T₁) (11.97 cm) at 20 DAT. At 40 DAT, significantly maximum (17.56 cm) leaf length was registered in T₃ (GA₃ @ 100ppm-foliar spray). However, minimum (15.31 cm) leaf length was observed in treatment T₁ (Control). The maximum leaf length (23.53 cm and 41.53 cm) was registered in T₃ (GA₃ @ 100ppm-foliar spray) at 60 DAT and 80 DAT, respectively. However, minimum leaf length (19.17 cm and 34.13 cm) was observed in control.

Leaf width (cm)

The data clearly indicated that the leaf width of onion plants responded significantly to various treatments of plant growth regulators and methods of application. At 20 days after transplanting, the significantly maximum leaf width (0.40 cm) was recorded under the treatment T₃ (GA₃ @ 100ppm-foliar spray), followed by T₆ (NAA @ 100ppm-foliar spray) (0.37 cm) as compared to other genotypes, while, minimum leaf width (0.30 cm) was recorded under control.

The maximum leaf width (1.38 and 1.48 cm) was recorded under the treatment T₃ (GA₃ @ 100ppm-foliar spray) at 40 DAT and 60 DAT, respectively. While, minimum leaf width (1.14 cm and 1.23 cm) was recorded under control at 40 DAT and 60 DAT, respectively. At 80

days after transplantation, the significantly maximum (1.63 cm) leaf width was recorded under the treatment T₃ (GA₃ @ 100ppm-foliar spray), followed by T₆ (NAA @ 100ppm-foliar spray) (1.60 cm) as compared to other treatments. While, minimum leaf width (1.37 cm) was recorded under control. Maximum leaf length and leaf width were recorded in T₃ (GA₃ @ 100ppm-foliar spray).

The increase in plant height by foliar spray of GA₃ 100ppm and NAA 100ppm might be due to rapid increase in cell division and cell elongation in the meristemic region. The foliar spray of GA₃ and NAA might be responsible for rapid increase in cell division and cell elongation in the meristemic region. These findings are in agreement with the findings of Nandekar and Sawarkar (1992), Patel *et al.*, (2010a) and Patel *et al.*, (2010b) they reported that increase in leaf length with the foliar spray of GA₃ and NAA. Singh *et al.*, (1995) and Islam *et al.*, (2007) also supported the leaf length and width increased with these treatments.

Leaf area (cm²)

The leaf area and Pseudostem length was recorded and has been presented in table 3. At 20 days after transplantation, the significantly maximum (15.49, 15.48, 14.82, 14.27, 13.99 and 13.63 cm²) leaf area were recorded in T₃ (GA₃ @ 100ppm-foliar spray), T₆ (NAA @ 100ppm-foliar spray), T₂ (GA₃ @ 100ppm-seedling dip.), T₅ (NAA @ 100ppm-seedling dip.), T₄ (GA₃ @ 100ppm-dropping method) and T₇ (NAA @ 100ppm-dropping methods), respectively and which were at par with each other and minimum (10.22 cm²) was observed under control. The maximum leaf area (120.68 cm² and 186.53 cm²) was recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray) at 40 DAT and 60 DAT, respectively whereas minimum leaf area (88.81 cm² and 144.99 cm²) was recorded under control at 40 DAT and 60 DAT, respectively.

Table.1 Effect of plant growth regulators and methods of application on plant height (cm) and number of leave per plant at 20, 40, 60 and 80 DAT

Treatments	Plant height (cm)				Number of leaves per plant			
	20DAT	40DAT	60DAT	80DAT	20DAT	40DAT	60DAT	80DAT
Control	13.87	21.93	25.7	42.07	2.53	4.67	4.8	5.67
GA ₃ @ 100ppm- seedling dip.	16.2	23.37	29.27	46.16	2.93	5.2	5.63	6.47
GA ₃ @ 100ppm-foliar spray	17.67	23.96	30.10	48.03	3.13	5.33	6.00	6.47
GA ₃ @ 100ppm-dropping methods	15.93	23.17	29.03	45.89	2.80	5.10	5.53	6.40
NAA @ 100ppm- seedling dip.	16.02	23.23	29.03	46.13	2.93	5.13	5.53	6.40
NAA @ 100ppm-foliar spray	16.47	23.57	30.00	47.93	3.00	5.27	5.63	6.47
NAA @ 100ppm-dropping methods	15.60	23.00	28.67	45.80	2.80	5.07	5.47	6.30
CCC @ 100ppm- seedling dip.	15.13	22.77	28.33	44.20	2.73	4.93	5.40	6.27
CCC @ 100ppm-foliar spray	15.27	22.83	28.47	45.73	2.80	5.03	5.43	6.30
CCC @ 100ppm-dropping methods	15.00	22.63	28.30	43.65	2.73	4.93	5.37	6.20
TIBA @ 100ppm- seedling dip.	14.33	22.30	27.57	42.97	2.73	4.73	5.20	6.00
TIBA @ 100ppm-foliar spray	14.50	22.40	27.73	43.31	2.73	4.87	5.33	6.13
TIBA @ 100ppm-dropping methods	14.20	22.03	25.90	42.93	2.67	4.67	5.00	6.00
S.Em±	0.33	0.24	0.35	0.33	0.08	0.06	0.05	0.07
C.D. (5%)	0.98	0.7	1.04	0.97	0.25	0.19	0.16	0.21

Table.2 Effect of plant growth regulators and methods of application on leaf length and width (cm) at 20, 40, 60 and 80 DAT

Treatments	Leaf length (cm)				Leaf width (cm)			
	20DAT	40DAT	60DAT	80DAT	20DAT	40DAT	60DAT	80DAT
Control	11.97	15.31	19.17	34.13	0.3	1.14	1.23	1.37
GA₃ @ 100ppm- seedling dip.	14.07	16.93	22.67	39.87	0.36	1.30	1.46	1.58
GA₃ @ 100ppm-foliar spray	15.60	17.56	23.53	41.53	0.40	1.38	1.48	1.63
GA₃ @ 100ppm-dropping methods	14.00	16.70	22.53	39.63	0.35	1.27	1.45	1.54
NAA @ 100ppm- seedling dip.	14.05	16.87	22.53	39.63	0.35	1.28	1.45	1.58
NAA @ 100ppm-foliar spray	14.57	16.97	23.50	41.17	0.37	1.31	1.48	1.60
NAA @ 100ppm-dropping methods	13.9	16.57	22.30	39.39	0.35	1.27	1.41	1.52
CCC @ 100ppm- seedling dip.	13.39	16.23	21.83	37.70	0.34	1.25	1.39	1.52
CCC @ 100ppm-foliar spray	13.83	16.37	21.97	39.27	0.34	1.25	1.40	1.52
CCC @ 100ppm-dropping methods	12.77	16.00	21.67	36.83	0.33	1.24	1.37	1.51
TIBA @ 100ppm- seedling dip.	12.36	15.70	20.93	36.10	0.32	1.20	1.36	1.49
TIBA @ 100ppm-foliar spray	12.40	15.70	21.23	36.70	0.32	1.21	1.36	1.51
TIBA @ 100ppm-dropping methods	12.13	15.53	19.83	35.57	0.30	1.19	1.34	1.47
S.Em±	0.3	0.31	0.33	0.39	0.005	0.027	0.008	0.008
C.D. (5%)	0.89	0.93	0.98	1.15	0.01	0.08	0.02	0.02

Table.3 Effect of plant growth regulators and methods of application on leaf area (cm²) and pseudostem length (cm) at 20, 40, 60 and 80 DAT

Treatments	Leaf area (cm ²)				Pseudostem length (cm)			
	20DAT	40DAT	60DAT	80DAT	20DAT	40DAT	60DAT	80DAT
Control	10.22	88.81	144.99	289.17	2.07	5.90	7.27	7.47
GA₃ @ 100ppm- seedling dip.	14.82	109.76	178.16	388.06	2.67	6.34	8.33	8.42
GA₃ @ 100ppm-foliar spray	15.49	120.68	186.53	428.53	2.91	6.58	8.86	9.03
GA₃ @ 100ppm-dropping methods	13.99	105.12	170.77	374.13	2.52	6.26	8.00	8.17
NAA @ 100ppm- seedling dip.	14.27	106.92	175.48	374.16	2.52	6.30	8.10	8.23
NAA @ 100ppm-foliar spray	15.48	116.5	183.11	417.91	2.67	6.48	8.63	8.86
NAA @ 100ppm-dropping methods	13.63	103.67	167.13	369.47	2.49	6.22	7.97	8.03
CCC @ 100ppm- seedling dip.	12.35	100.83	157.45	357.18	2.30	6.15	7.77	7.83
CCC @ 100ppm-foliar spray	12.45	101.81	164.6	367.29	2.48	6.19	7.83	7.95
CCC @ 100ppm-dropping methods	11.94	96.87	156.53	351.97	2.25	6.10	7.70	7.77
TIBA @ 100ppm- seedling dip.	11.56	94.08	152.58	342.59	2.10	6.00	7.47	7.59
TIBA @ 100ppm-foliar spray	11.56	94.19	156.22	349.07	2.15	6.07	7.63	7.70
TIBA @ 100ppm-dropping methods	10.27	91.93	150.21	329.55	2.10	5.95	7.40	7.53
S.Em±	0.67	3.87	3.3	6.5	0.05	0.01	0.05	0.05
C.D._{.5%} level	1.96	11.3	9.63	18.97	0.16	0.05	0.14	0.17

Table.4 Effect of plant growth regulators and methods of application on fresh and dry weight of plant (g) at 20, 40, 60 and 80 DAT

Treatments	Fresh weight of plant (g)				Dry weight of plant (g)			
	20DAT	40DAT	60DAT	80DAT	20DAT	40DAT	60DAT	80DAT
Control	9.00	24.20	36.00	41.93	0.90	3.85	6.72	8.37
GA₃ @ 100ppm- seedling dip.	17.00	36.30	48.43	58.53	2.90	7.89	9.78	11.88
GA₃ @ 100ppm-foliar spray	18.20	43.27	50.07	60.27	3.60	8.98	10.32	12.11
GA₃ @ 100ppm-dropping methods	15.60	35.27	43.97	55.73	2.60	7.39	8.96	11.63
NAA @ 100ppm- seedling dip.	16.10	35.60	47.23	57.43	2.80	7.72	9.76	11.67
NAA @ 100ppm-foliar spray	17.40	40.30	48.53	59.00	3.50	7.99	10.07	12.01
NAA @ 100ppm-dropping methods	14.60	34.30	41.23	53.73	2.60	7.38	8.59	10.27
CCC @ 100ppm- seedling dip.	14.30	34.00	41.10	51.23	2.50	5.25	8.41	10.10
CCC @ 100ppm-foliar spray	14.50	34.23	41.13	51.80	2.60	6.38	8.43	10.13
CCC @ 100ppm-dropping methods	13.90	29.07	40.77	50.70	2.30	4.83	7.40	9.78
TIBA @ 100ppm- seedling dip.	13.50	24.67	38.83	46.33	1.80	4.44	7.00	8.85
TIBA @ 100ppm-foliar spray	13.50	28.97	39.77	48.70	2.00	4.66	7.10	9.13
TIBA @ 100ppm-dropping methods	12.60	24.23	37.70	43.93	1.30	4.34	6.90	8.51
S.Em±	2.47	2.39	2.46	1.88	0.19	0.31	0.54	0.29
C.D. (5%)	7.23	6.98	7.18	5.51	0.56	0.92	1.60	0.85

Table.5 Effect of plant growth regulators and methods of application diameter and neck thickness of bulb

Treatments	Diameter of bulb (cm)		Neck thickness of bulb (cm)
	Polar	Equatorial	
Control	5.15	5.32	1.40
GA₃ @ 100ppm- seedling dip.	5.73	5.78	1.23
GA₃ @ 100ppm-foliar spray	5.77	5.91	1.18
GA₃ @ 100ppm-dropping methods	5.61	5.71	1.25
NAA @ 100ppm- seedling dip.	5.69	5.77	1.23
NAA @ 100ppm-foliar spray	5.74	5.88	1.23
NAA @ 100ppm-dropping methods	5.58	5.66	1.27
CCC @ 100ppm- seedling dip.	5.51	5.62	1.29
CCC @ 100ppm-foliar spray	5.53	5.66	1.28
CCC @ 100ppm-dropping methods	5.45	5.59	1.30
TIBA @ 100ppm- seedling dip.	5.35	5.53	1.36
TIBA @ 100ppm-foliar spray	5.43	5.55	1.35
TIBA @ 100ppm-dropping methods	5.20	5.40	1.38
S.Em±	0.02	0.06	0.01
C.D. (5%)	0.06	0.18	0.03

Leaf area fairly gives a good idea of the photosynthetic capacity of the plant. In the present study, it has been observed that the application of plant growth regulators had profound influence on assimilatory surface area. In general, leaf area increased from 20 DAT to 80 DAT. The treatment T₃ (GA₃ @ 100ppm-foliar spray). The results of the present investigation are in accordance with the observations of Ganiger *et al.*, (2002).

Pseudostem length (cm)

The significantly maximum (2.91, 6.58 and 8.86 cm) pseudostem length were recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray), followed by T₆ (NAA @ 100ppm-foliar spray) (2.67, 6.48 and 8.0 cm) at 20, 40 and 60 DAT respectively whereas minimum (2.07, 5.90 and 7.27 cm) was found under control. At 80 DAT, the maximum pseudostem length (9.03 cm) were recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray) and minimum (7.47 cm) in control.

Foliar application of growth regulators recorded the significant difference with respect to pseudostem length of onion. The maximum pseudostem length were recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray). It may be due to the growth regulators, like GA₃ and NAA are involved in cell division, cell expansion, cell elongation and cell differentiation there by leading to enhanced pseudostem length.

Fresh weight of plant (g)

At 20 DAT, the fresh weight of plants increased significantly by the different treatments at all the growth stages. The significantly maximum (18.2g) fresh weight of plant was recorded in the treatment T₃ (GA₃ @ 100ppm-foliar spray), followed by T₆ (NAA @ 100ppm-foliar spray) (17.4 g) as compared to other treatments. However, the

treatment T₁ (Control) was exhibited minimum fresh weight of plant (9.0 g).

At 40 DAT, the significantly maximum (43.27, 40.30 and 36.30g) fresh weights of plant were recorded in the treatment T₃ (GA₃ @ 100ppm-foliar spray), T₆ (NAA @ 100ppm-foliar spray) and T₂ (GA₃ @ 100ppm- seedling dip.), respectively and which were at par with each other. However, the treatment T₁ (Control) was exhibited minimum fresh weight of plant (24.20 g). The significantly maximum (50.07g) and (60.27g) fresh weights of plant were recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray), followed by T₆ (NAA @ 100ppm-foliar spray) (48.53 and 59.0g), T₂ (GA₃ @ 100ppm- seedling dip.) (48.43 and 58.53g), T₅ (NAA @ 100ppm- seedling dip.) (47.23 and 57.43g) and T₄ (GA₃ @ 100ppm-dropping method) (43.97 and 55.73g) at 60 and 80 DAT, respectively and which were at par with each other whereas minimum (36.0 and 41.93g at 60 and 80 DAT, respectively) was found under control.

Dry weight of plant (g)

The average dry weight of plant of different treatments is given in Table 4. Dry weight of plant was recorded at 20, 40, 60 and 80 days after transplanting. As regards to 20 DAT, the dry weight of plants increased significantly by the different treatments at all the growth stages. The significantly maximum (3.6 and 3.5 g) dry weight of plant were recorded in the treatment T₃ (GA₃ @ 100ppm-foliar spray) and T₆ (NAA @ 100ppm-foliar spray), respectively and which were at par with each other. However, the treatment T₁ (control) was exhibited minimum dry weight of plant (0.9 g).

At 40 DAT, the significantly maximum (8.98g) dry weight of plant was recorded in the treatment T₃ (GA₃ @ 100ppm-foliar spray)

followed by T₆ (NAA @ 100ppm-foliar spray) (7.99 g) as compared to other treatments. However, the treatment T₁ (Control) was exhibited minimum dry weight of plant (3.85 g). The significantly maximum (10.32 g and 12.11 g) dry weight of plant were recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray), followed by T₆ (NAA @ 100ppm-foliar spray) (10.07 and 12.01 g), T₂ (GA₃ @ 100ppm-seedling dip.) (9.78 and 11.88 g), T₅ (NAA @ 100ppm- seedling dip.) (9.76 and 11.67 g) and T₄ (GA₃ @ 100ppm-dropping method) (8.96 and 11.63 g) at 60 and 80 DAT, respectively and which were at par with each other. Therefore, it was observed minimum (6.72 and 8.37 g at 60 and 80 DAT, respectively) in treatment T₁ (control).

Foliar application of growth regulators recorded the significant difference with respect to fresh weight of onion plant. In general, fresh weight of plant increased from 20 DAT to 80 DAT. The significantly maximum fresh weight of plant was recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray). It may be due to the role of these materials on enhancing cell division activity, increasing of proline accumulation of plant and increasing of endogenous phytohormones i.e. increasing promotion hormones (IAA, GA₃ and cytokinins) and reducing ABA content which found that bio-regulators make a shift in hormonal balance characterized by increasing in endogenous phytohormone in plant. Results of the present investigation were also in confirmatory with the findings of Ledesma *et al.*, (2000), Islam *et al.*, (2007) and Ouzounidou *et al.*, (2011).

The amount of total dry matter produced is an indication of the overall efficiency of utilization of resources and better interception of light even if the dry matter production in general is the indication of the efficiency of genotypes. The enhanced productivity of crop through approaches is chiefly achieved by coordinating plant processes to synthesize

maximum dry matter and partitioning of the major quantum of this increased dry matter into effective yield contributing factors. Poor translocation of assimilates to the reproductive parts (bulb) is the major constraint in onion.

This can be overcome by the application of growth regulators, which can improve canopy structure and increase the productivity through manipulation of source-sink relationship. In the present study, it was observed that partitioning of total dry matter in leaf and bulb parts varied significantly due to the growth regulator treatments. The amount of dry weight of plant produced is an indication of the overall efficiency of the utilization of resources and better light interception. The data pertaining to total dry weight per plant indicated that, it increased from 20 DAT to 80 DAT. The increase in dry weight of plant up to 80 DAT may be due to higher rate of CO₂ fixation and RUBP Carboxylase activity in the early stage of crop growth.

The application of growth regulators significantly improved dry weight of plant and was recorded as maximum in treatment T₃ (GA₃ @ 100ppm-foliar spray). Similarly, Nirmal *et al.*, (1994), Ledesma *et al.*, (2000), Ganiger *et al.*, (2002), Suheela *et al.*, (2005) and Ouzounidou *et al.*, (2011) also reported significant variation in dry weight of plant.

Diameter of bulb (cm)

The polar diameter of bulb increased significantly due to different treatments. Significantly maximum (5.77, 5.74 and 5.73cm) polar diameter of onion bulb were exhibited in the treatment T₃ (GA₃ @ 100ppm-foliar spray), T₆ (NAA @ 100ppm-foliar spray) and T₂ (GA₃ @ 100ppm- seedling dip.), respectively and which were at par with each other. However, the minimum polar diameter of bulb was observed in Control (5.15 cm).

The equatorial diameter of bulb increased significantly due to different PGRs. Significantly maximum equatorial diameter of onion bulb (5.91, 5.88, 5.78 and 5.77cm) were exhibited in the treatment T₃ (GA₃ @ 100ppm-foliar spray), T₆ (NAA @ 100ppm-foliar spray), T₂ (GA₃ @ 100ppm- seedling dip.) and T₅ (NAA @ 100ppm- seedling dip.), respectively and the minimum equatorial diameter of bulb was observed in control (5.32 cm).

It could be noticed that, all treated plants resulted in the highest polar and equatorial diameter of bulb comparing with untreated control. It can be concluded that, spraying onion plant with (GA₃ @ 100ppm-foliar spray), (NAA @ 100ppm-foliar spray) and (GA₃ @ 100ppm- seedling dip.) resulted in rapid cell division and elongation leading to bigger bulb formation. Results was also in confirmatory with the findings of Tomar *et al.*, (1988), Shakhda and Gajipara (1998), Tiwari *et al.*, (2001), Islam *et al.*, (2007), Bose *et al.*, (2009), Rashid(2010) and Patel *et al.*, (2010a).

Neck thickness of bulb (cm)

The neck thickness of bulb was significantly influenced by PGR. The minimum neck thickness (1.18 cm) was recorded in treatment T₃ (GA₃ @ 100ppm-foliar spray) as compared to control (1.40 cm).

The significantly lower neck thickness was noticed in the treatment GA₃ @ 100ppm-foliar spray.

The higher neck thickness was noticed in control. The thickness of the stem (neck) is an important parameter for storage of bulb. Hence, more the thickness of the neck more will be the rotting due to more fungous infection. The results of the present investigation are in accordance with the observations of Islam *et al.*, (2007), Bose *et*

al., (2009) and Govind *et al.*, (2015).

Foliar application of GA₃ 100ppm (T₃) was recorded significant maximum growth parameters (plant height, number of leaves plant⁻¹, leaf length, leaf width, leaf area, and pseudostem length), fresh and dry weight of plant, polar and equatorial diameter of bulb and neck thickness.

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