

Original Research Article

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Effect of Holding Solutions on the Water Relations in Vase Life of Cut Carnation cv. Kiro

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ABSTRACT

Keywords

Carnation, Holding solution, Water uptake, Transpirational loss of water, Water balance, Fresh weight change

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An investigation was carried out to study the effect of different holding solution combinations on vase life of cut carnation cv. Kiro. The flowers were kept under common holding solution (sucrose 4% + 8-HQS 200 ppm) along with GA₃ at 25 ppm recorded significantly highest water uptake on 6th day (33.83 g flower⁻¹) the highest transpirational loss of water (32.16 g flower⁻¹) and highest fresh weight change of flower (152.38 g flower⁻¹) which have extended the vase life period of carnation flower cv. Kiro.

Introduction

Carnation (*Dianthus caryophyllus* L.) is an important cut flower in the world. Carnation is a climatic flower that is highly sensitive to ethylene (Pun *et al.*, 1999). Due to high perishability, cut flowers are vulnerable to large post-harvest losses upto 50 per cent of the farm value (Singh *et al.*, 2007). Carnations are more susceptible to mechanical and physical damages and microbial infections by diseases and pests during and after harvest. Floral preservatives affect the quality of cut flowers by extending the vase life, increasing flower size and maintaining the colour of leaves and petals.

The vase life of cut flowers and foliage is often shortened by vascular occlusions that constrict vase solutions supply, reduction in stem conductivity is typically caused by blockage of cut stem ends and xylem conduits by microbes, physiological plugging and water columns in xylem vessels by cavitations and air emboli.

Cut flower and foliage longevity can be greatly affected by chemical composition of the vase solution. A broad range of biocides has been suggested to prevent the proliferation of microorganisms in vase solutions. However, their assumed antimicrobial action may be confounded by their other

physicochemical effects (Edrisi *et al.*, 2012). Water relations plays a critical role in the post-harvest life of cut flowers, water imbalance within the cut flower resulting in wilting, one of the major causes for termination of vase life (Halevy and Mayak, 1981).

Materials and Methods

The experiment was held in laboratory of Floriculture and Landscape Architecture, College of Horticulture, Dr. Y.S.R. Horticultural University, Venkataramannagudem, West Godavari dist (A.P), during year 2017-18. Experiment was laid out in completely randomised design. Sucrose 4% + 8-HQS 200 ppm was used as common holding solution. There are 11 treatments, T₁: holding solution + GA₃ 25 ppm, T₂: holding solution + GA₃ 50 ppm, T₃: holding solution + BA 25 ppm, T₄: holding solution + BA 50 ppm, T₅: holding solution + Al₂(SO₄)₃ 150 ppm, T₆: holding solution + Al₂(SO₄)₃ 300 ppm, T₇: holding solution + STS 0.25 mM, T₈: holding solution + STS 0.50 mM, T₉: holding solution + Salicylic acid 25 ppm, T₁₀: holding solution + Salicylic acid 50 ppm, T₁₁: control (only holding solution). All the treatments are replicated thrice at 25 ± 2°C ambient room temperature, 45-55 per cent relative humidity RH and 40 W cool white florescent tubes to maintain 12 hours photoperiod. In each conical flask, 6 flowers were placed in each 500 ml conical flasks 300 ml of holding solution. Observations were recorded changes on water uptake, transpirational loss of water, water balance, fresh weight change and vase life.

Results and Discussion

Carnation cut flowers cv. Kiro kept under different holding solution differed significantly on water uptake, among all the treatments the flowers kept in (T₁) holding solution (sucrose 4% + 8-HQS 200 ppm)

along with GA₃ at 25 ppm recorded the highest water uptake (30.28 g flower⁻¹) from 2nd day to 10th day (28.67 g flower⁻¹) of vase life period while the lowest water uptake (19.33 g flower⁻¹) was recorded by control (T₁₁) only holding solution though the vase life period (Table 1). Improved water uptake might be due to the combined effect of holding solution (sucrose 4% + 8-HQS 200 ppm) with GA₃. Sucrose is a source of energy and good respiratory substrate for the maintenance of osmotic potential in flowers and improved the ability of the tissue to absorb water, hence maintain turgidity. Similar results were also reported by Lol *et al.*, (1990) in gladiolus, Reddy and Singh (1996) and Bhaskar *et al.*, (1999) in tuberose.

The 8-HQS has germicidal and chelating properties might have reduced the stem blockage and maintained the water conductivity in accordance with Reddy *et al.*, (1995) in tuberose. GA₃ treated flowers had continuous water uptake this might be due to hydrolysis of polysaccharides and starch into glucose and fructose which decreased the water potential in stem and flower. These results are agreed with Nowak and Mynett (1985) in Lilium and Karimi (2007) in lily.

The transpirational loss of water was significantly highest (28.46 g flower⁻¹) from 2nd day to 10th day (27.86 g flower⁻¹) with flowers kept in (T₁) holding solution (sucrose 4% + 8-HQS 200 ppm) along with GA₃ at 25 ppm, while the lowest transpirational loss of water (18.53 g flower⁻¹) throughout the vase life period was recorded by control (T₁₁).

When the amount of transpiration exceeds absorption, a water deficit and wilting develops. Gibberellic acid increased the water uptake and reduced the transpiration. Phytohormones have been implicated in the regulation of flower senescence (Halevy and Mayak, 1981).

Table.1 Effect of different holding solutions on water uptake (g/flower) and transpirational loss of water (g/flower) during vase life of cut carnation cv. Kiro

Treatments	Time period (days)									
	Water uptake (g/flower)					Transpirational loss of water (g/flower)				
	2	4	6	8	10	2	4	6	8	10
T ₁ -Holding solution + GA ₃ @ 25ppm	30.28	32.19	33.83	31.74	28.67	28.46	29.77	32.16	29.12	27.86
T ₂ -Holding solution + GA ₃ @ 50ppm	28.62	30.82	31.56	29.93	27.89	27.52	28.65	31.06	27.86	25.42
T ₃ -Holding solution + BA @ 25ppm	26.95	28.91	29.55	27.84	25.49	25.64	27.53	28.42	26.73	24.86
T ₄ -Holding solution + BA @ 50ppm	25.21	27.85	28.73	26.67	24.38	24.59	25.94	27.13	24.91	22.74
T ₅ - Holding solution + Al ₂ (SO ₄) ₃ @ 150ppm	24.94	26.95	27.78	25.81	23.6	23.90	25.83	27.10	24.89	22.92
T ₆ - Holding solution + Al ₂ (SO ₄) ₃ @ 300ppm	25.15	27.13	28.31	26.17	23.94	24.25	26.74	27.62	25.65	23.19
T ₇ - Holding solution + STS @ 0.25 mM	23.44	25.72	26.29	24.45	22.53	22.76	24.72	25.46	23.63	21.85
T ₈ - Holding solution + STS @ 0.50 mM	24.58	26.81	27.42	25.62	23.49	23.62	25.53	26.37	24.56	22.64
T ₉ :Holding solution + Salicylic acid @ 25ppm	20.92	23.14	23.86	22.29	18.04	19.87	20.67	22.71	19.82	17.56
T ₁₀ : Holding solution + Salicylic acid @ 50ppm	22.68	24.53	24.94	23.47	18.89	20.87	22.42	23.54	21.45	18.20
T ₁₁ : Control (only holding solution)	19.33	20.95	18.22	15.07	-	18.53	20.07	17.80	14.47	-
Mean	24.72	26.81	27.31	26.27	23.81	23.63	25.26	26.30	23.91	20.65
SE d	0.625	0.569	0.610	0.628	0.494	0.553	0.517	0.723	0.534	0.451
C.D at 5%	1.306	1.187	1.274	1.310	1.031	1.155	1.079	1.509	1.114	0.941

Holding solution- Distilled water + sucrose 4% + 8-HQS 200 ppm

*Significant at (P≤0.05)

Table.2 Effect of different holding solutions on water balance (g/flower) and fresh weight change (g/flower) during vase life of cut carnation cv. Kiro

Treatments	Time period (days)									
	Water balance (g/flower)					Fresh weight change (g/flower)				
	2	4	6	8	10	2	4	6	8	10
T ₁ -Holding solution + GA ₃ @ 25ppm	8.96	7.52	7.07	6.75	6.28	129.20	137.32	152.38	138.80	122.56
T ₂ -Holding solution + GA ₃ @ 50ppm	8.16	7.02	6.83	6.14	5.92	121.52	130.50	141.26	126.71	115.80
T ₃ -Holding solution + BA @ 25ppm	7.69	6.94	6.15	5.73	5.34	120.75	126.63	130.81	122.49	114.65
T ₄ -Holding solution + BA @ 50ppm	7.27	6.07	5.75	5.17	4.85	118.47	123.46	126.22	119.75	109.24
T ₅ - Holding solution + Al ₂ (SO ₄) ₃ @ 150ppm	6.80	5.85	5.00	4.35	4.06	115.01	117.47	120.48	115.34	112.23
T ₆ - Holding solution + Al ₂ (SO ₄) ₃ @ 300ppm	6.84	5.94	5.10	4.72	4.23	115.62	118.78	123.56	116.45	111.80
T ₇ - Holding solution + STS @ 0.25 mM	5.89	5.05	4.45	3.80	3.14	112.35	114.66	117.81	109.86	98.58
T ₈ - Holding solution + STS @ 0.50 mM	6.75	5.74	4.90	4.22	3.95	114.71	116.50	119.65	114.26	108.71
T ₉ :Holding solution + Salicylic acid @ 25ppm	5.12	4.79	3.98	3.23	2.49	107.35	109.82	104.81	99.45	92.37
T ₁₀ : Holding solution + Salicylic acid @ 50ppm	5.68	5.11	4.21	3.97	3.14	110.33	112.74	109.40	103.92	96.58
T ₁₁ : Control (only holding solution)	4.59	4.13	3.09	2.47	-	103.90	105.60	101.22	90.56	-
Mean	6.70	5.83	5.13	4.595	3.94	115.35	119.37	124.25	114.93	98.84
SE d	0.138	0.123	0.106	0.068	0.064	1.999	1.864	1.96	2.450	2.314
C.D at 5%	0.288	0.259	0.221	0.143	0.134	4.172	3.889	4.089	5.112	4.828

Holding solution- Distilled water + sucrose 4% + 8-HQS 200 ppm

*Significant at (P≤0.05)

The water balance differed significantly among the treatments, among all the treatments, flowers kept in (T₁) holding solution (sucrose 4% + 8-HQS 200 ppm) along with GA₃ at 25 ppm recorded the highest water balance (8.96 g flower⁻¹) through the vase life period while the lowest water balance (4.59 g flower⁻¹) was recorded by control (T₁₁) only holding solution (Table 2). Improved water balance might be due to the combined effect of holding solution (sucrose 4% + 8-HQS 200 ppm) with GA₃. Water balance is a major factor influencing the quality and longevity of cut flowers. The increased water balance is due to increased water uptake and with decreased loss of water. The physiological efficiency of GA₃ could maintain the positive water balance. Gibberellic acid further promoted dry matter (starch) hydrolysis into reducing sugars in the stem and flower heads, leading to enhanced vase life and quality of cut flowers via improved water balance. 8-HQS plays an important role in improving the water balance of cut freesia by preventing the growth of microorganisms in xylem and thus maintained water uptake by flower stems (Kwon and Kim, 2000). Sucrose helps in improving the water balance of cut flowers by affecting the osmotic potential of cut flowers and the water holding capacity of the tissues allowing less water to be transpired (Halevy *et al.*, 1978).

The fresh weight was significantly highest (129.20 g flower⁻¹) on 2nd day to 10th day (122.56 g flower⁻¹) with flowers kept in (T₁) holding solution (sucrose 4% + 8-HQS 200 ppm) along with GA₃ at 25 ppm while the lowest fresh weight change (103.90 g flower⁻¹) was recorded by control (T₁₁) only holding solution (Table 2). The gain in fresh weight change was due to the synergistic effect of holding solution (sucrose 4% + 8-HQS 200 ppm) along with GA₃ might be due to GA₃ increased water relations in the floral tissue by inducing stomatal closure in the leaves

might have led to an increased metabolic activity without loss of quality. The combined effect of sucrose and 8-HQS might have improved the water retention of the flower contributed high fresh weight change. Similar results were reported by Jitendra kumar and Daljeet Singh (2004), Sagar *et al.*, (2005) in tuberose.

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