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The Pharma Innovation



ISSN (E): 2277-7695 ISSN (P): 2349-8242 NAAS Rating: 5.23 TPI 2022; 11(9): 2350-2354 © 2022 TPI www.thepharmajournal.com

Received: 08-06-2022 Accepted: 12-07-2022

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Standardization of the pre-cooling method for extending the shelf life of tuberose florets stored at 10 $^\circ\mathrm{C}$

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Abstract

Investigation was carried out to evaluate the effect of different pre-cooling methods to extend shelf life of tuberose florets. The harvested flowers were treated with different pre-cooling methods (Pre-cooling at 4 °C, hydro cooling at 4 °C and by use of gel ice packs), followed by treatment with sucrose (10% and 4%) and boric acid (4% and 2%). Analysis was done in terms of physical and chemical characteristics of flowers. Among the three pre-cooling methods, flowers treated with 4% boric and pre-cooled at 4°C showed better shelf life up to 4 days with least PLW (22.02%), respiration rate (94.43 ml CO₂ kg⁻¹ h⁻¹), witnessed minimum wilting (30.37%), recorded least polyphenol enzyme activity without adversely affecting physico-chemical qualities even up to 4 days of storage. Main objective is to study the response of pre-cooling methods and chemicals (sucrose and boric acid) on extending the marketability of tuberose flowers.

Keywords: Polianthes tuberosa, boric acid, respiration rate, polyphenol enzyme activity

1. Introduction

Tuberose (*Polianthes tuberosa* L.) belongs to the family Asperagaceae native to Mexico. It is being grown in most of the tropical and sub-tropical countries of world (Asif *et al.*, 2001)^[1]. Tuberose is an important commercial cut as well as loose flower crop due to its pleasant fragrance, longer vase-life of spikes, higher returns and wider adaptability to varied climate and soil. They are valued much by the aesthetic world for their beauty and fragrance. Tuberose represents sensuality and is used in aromatherapy for its ability to open the heart and calm the nerves, restoring joy, peace and harmony.

Postharvest performance is worse in tuberose which has been shipped to distant markets. The quality of tuberose flower is affected by various pre and post-harvest factors such as temperature, relative humidity, frequency of irrigation, picking time, nutrition and handling practices Benschop (1993)^[2]. The major problem during marketing of tuberose loose flowers is the short shelf life and harvested flowers loss its fresh weight up to 40%, when it is kept overnight due to its high rate of respiration and higher metabolic activity (Reddy 2016)^[14]. Lack of standardized packaging method is the main reason which deteriorates the quality of loose flowers.

Proper post-harvest treatments can greatly extend the shelf life of tuberose and routinely be carried out with flowers intended for storage and transport. The different post-harvest technologies available are pre cooling, use of chemical preservatives and different packaging materials. Among these pre-cooling is considered to be critical process in enhancing the post-harvest longevity of the flowers. (Brosnan and Sun 2001) ^[3]. Pre cooling methods like room cooling, hydro cooling and gel ice pack which helps to remove the field heat, reduce load on refrigerated storage. Several treatments like chemicals, sugar and germicide also increases longevity of tuberose florets (Talukdar and Barooah 2011) ^[19]. Further, storage of the flowers at low temperatures enhances the post-harvest life. Adequate packaging protects the produce from physical, physiological and pathological deterioration during transport and marketing which helps in extending shelf life by retaining their attractiveness. Use of ethylene absorbent in packages helps in delaying senescence of florets by absorbing ethylene produced inside packages. Treating flower buds with novel chemicals before packaging and transport act as barrier for respiration, moisture loss, to extend the shelf life with better retention of colour and turgidity of petals.

2. Materials and Methods

2.1 Materials

The present investigation was undertaken in the Department of Post-Harvest Technology, University of Horticultural Sciences, Bagalkot, India during the year 2017-18. Prajwal, a commercially important tuberose cultivar grown in this region was used for the study. This cultivar is a single type, having more fragrance, extensively cultivated and used as loose flower. Flower buds were procured from historical place, Badami, near Bagalkot. The unopened flower buds at pin hole stage which were ready to open in subsequent days were harvested in the early morning and they were brought to the laboratory within 2 hours. Three different methods of precooling and two pulsing chemicals were used in the study to evaluate physiological, biochemical and physical characteristics of tuberose.

2.1.1 Pre cooling methods

- 1. Buds were pre cooled by placing them at 4 $^\circ\mathrm{C}$ for about one hour
- 2. Hydro cooling done by using cold distilled water of temperature 4°C was noted by using thermometer and buds were pre-cooled for about 5 min.
- 3. Buds were placed over the gel ice packs (25% weight of buds).

2.2 Methodology

2.2.1 Sucrose and boric acid

Sucrose solution of 4% and 10% were prepared by dissolving 100 gm and 40 gm in 1000 ml of distilled water, respectively. Boric acid of 2% and 4% was prepared by weighing 2 gm and 4 g separately and dissolved in 100 ml of warm water and volume made up to 1000 ml.

The pre-cooled buds were dipped in solution for about 15 minutes and removed to air dry under ceiling fan. The control buds were dipped in distilled water. The treated as well as untreated buds were placed in baskets and kept under storage 10 °C for further observations.

2.3 Physiological loss in weight (%)

Flowers from each treatment were taken to record the physiological loss in weight (PLW). The weight of the flowers was recorded using precision electronic weighing balance (Make: Sartorius Weighing Technology GmbH, Gottingen, Germany, GE812) before storage as initial weight. On subsequent dates of observation during storage, the flowers were weighed and recorded as final weight on every 2 days intervals and the cumulative PLW was calculated with the following formula and expressed as per cent physiological loss in weight:

2.4 Respiration rate (ml CO₂ kg⁻¹h⁻¹)

The respiration rate of the flowers packed with different gas composition was measured after taking them out of the polyethylene pouch during their storage.

The tuberose flowers of known volume were enclosed in a hermetic container for specified time and head space gas concentration of CO_2 was measured by piercing the probe of an auto oxygen/carbon dioxide analyzer (Make: Quantek, Model: 902D Dual track) into the container through the septa

fixed on the lid of container and direct reading was noted down from the instrument screen. The respiration rate was by using formula:

Respiration rate (mlco2kg^{-l}h^{-l}) = $\frac{\% \text{ CO}_{2x} \text{ Head space}}{100 \text{ x Flower weight (kg) x Enclosing time (hr)}}$

2.5 Wilting (%)

The wilting or fading of florets were recorded on visual basis, count the number of wilted florets and expressed in terms of percentage. The days taken for 50% wilting were noted which ends the shelf life of florets.

2.6 Shelf life (No. of days)

Time taken for development of necrotic symptoms was recorded and shelf life was determined as number of days taken from placing of flower buds till wilting or fading of petals.

2.11 Experimental design and data analysis

The experiment was carried out with 11 treatments and the experiment was repeated 3 times and pooled data was subjected to statistical analysis. Flowers were arranged in Complete Randomised Design. Randomly selected fruits were taken to analyse physiological loss in weight, respiration rate, wilting and shelf life. Statistical analyses were performed using Web Agri Stat Package (WASP) Version 2. Significant differences among means at P = 0.05 were determined by post hoc tests using Duncan's multiple range test.

3. Results and discussion

3.1 Physiological loss in weight (PLW) (%)

Physiological loss in weight (PLW) refers to the loss in weight of the fresh produce due to physiological processes like transpiration, respiration etc. In the study, result found that irrespective of pre cooling method used, the PLW of flowers increased from 2nd to 4th (22.62%-40.43%) day of storage at 10 °C. On 4th day of storage, minimum PLW of florets (29.56%) was recorded in T₄ and maximum in untreated (control) *i.e.* (59.66%) without pre-cooling. Increase in PLW with increased period of storage due to various physiological activities like transpiration and respiration. which leads to reduction in water content and organic compounds which are used as substrate during respiration of produce, increase in PLW leads to decline in fresh weight of the flowers, similar results are noticed by Sharma (1981)^[16] in Rosa damascena. This may be attributed to the fact that the pre cooling removes field heat and this low temperature slows down the rate of respiration and evaporation (Brosnan and Sun, 2001) ^[3].

Among various treatments, minimum PLW was recorded in the florets (buds) pre-cooled at 4 °C+4% Boric acid. Precooling and boric acid helps in retaining higher relative water content and lowest rates of PLW, slows down the respiration rate from the harvested cut flowers, accompanied by increased membrane integrity of florets as reported by Mukhopadhyay, 1980 ^[10] in jasmine and Mujumder *et al.*, 2014 ^[9] in tuberose and Kumar and Bhattacharjee (2002) ^[7] in rose. The highest PLW was recorded in case of buds without pre cooling due to residual field heat trapped in the buds leading to rapid loss of moisture and weight.

3.2 Respiration rate (ml CO₂ kg⁻¹ hr⁻¹)

Respiration is a central process in living cells that mediates

the release of energy through the oxidative breakdown of carbon compounds (Starch, sugars and organic acids) and the formation of carbon skeletons necessary for maintenance and synthetic reactions after harvest (Wills *et al.*, 1998) ^[20]. The respiration rate of flower is an excellent indicator of the metabolic activity of the tissue and thus is a useful guide to the potential shelf life of flowers.

In the study, the treatment T_4 (pre-cooled at 4 °C + 4% Boric acid) recorded minimum respiration rate (91.79 ml CO₂ kg⁻¹ hr⁻¹), however irrespective of treatments the respiration rate increased from 2nd to 4th (117.78 ml - 130.14 ml CO₂ kg⁻¹ hr⁻¹) day of storage. At the end of 4th day of storage the T4 recorded minimum respiration rate (94.43 ml CO₂ kg⁻¹ hr⁻¹) and maximum in control (167.56 ml CO₂ kg⁻¹ hr⁻¹) whereas other treatments showed intermediate results. This may be attributed to the rate of respiration slows down on pre cooling (Brosnan and Sun, 2001)^[3] further decrease in the respiration rate of the roses cv. 'Raktagandha' just after pre cooling treatment with cold storage at 4 °C for 24 h and ice cold water spray for 45 min as reported by (Palani kumar et al., 2000) ^[12]. Use of 4% boric acid showed minimum respiration rate, due to their role in inhibiting respiratory enzymes. Similar findings are also reported in tuberose (Mujumder et al., 2014) [9]

3.3 Wilting of florets

The most common and visibly apparent senescence symptom in flowers is the loss of turgidity of cell, resulting in wilting and death. Loss of weight will cause many perishable commodities to appear wilted or shrivelled under warm, dry conditions within few hours. Wilting occurs at the end of flower life due to increase in water loss by transpiration as reported by (Mayak *et al.*, 1974) ^[8]. The lesser wilting of florets (30.37%) was seen in the treatment T₄ (pre cooled at 4 °C treated with 4% boric acid), probably due to lesser rate of respiration and evaporation in pre-cooling (Bronsan and Sun, 2001) ^[3]. The results are in conformity with those of Palani kumar *et al.* (2000) ^[12] in 'Raktagandha' cut roses, pre cooled at 4 $^{\circ}$ C and in tuberose (Shil *et al.*, 2017) ^[17]. In addition to pre-cooling, boric acid could have improved water balance by increasing osmotic concentration and pressure potential of petal cells in tuberose, further boric acid also is reported to improve membrane stability and resistance against senescence related changes.

3.4 Shelf life

The term shelf life is commonly used in case of loose flowers and it refers to total life of a flower from the time of its harvesting (In the bud state) to the point of any of the following conditions like loss of colour, wilting, or loss of major aesthetic characteristics. In this investigation pre cooled tuberose showed better shelf life compared to control, this might be due to cooling is related to immediate decrease in whole metabolism of the flower which favours the extension of shelf life, maintaining its quality and reducing expenditures in the subsequent cold storage (Pellegrini and Belle, 2008)^[13]. Among different pre cooled methods used florets pre-cooled at 4 °C (T_1 , T_2 , T_3 and T_4) showed maximum shelf life of 4 days, followed by hydro cooling influenced by all physiological parameters similar findings in rose by (Palani kumar 2000) ^[12], hydro cooling may cause bud damage (Suisuwan 1994)^[18]. The action of ethylene can be slowed or stopped by anti-ethylene treatments this can be achieved by boric acid which delayed ethylene production as it is effective substitute for AOA, an inhibitor of ACC synthase activity and also add additional advantage in extending shelf life by improving water balance (Serrano et al., 2001)^[15]. It might be assumed that longest shelf life of flowers can be achieved by maintaining low temperature storage is helpful in conservation of flowers, because it reduces degradation of certain enzymes and ethylene production and slows the various process related to senescence (Nowak and Rudnick, 1990) [11].

	PLW (%)			
Treatments	Days of storage		Mean	
	2	4		
T_1 : $P_1 + 10\%$ sucrose	19.02	33.1	26.06	
$T_{2:} P_1 + 2\%$ boric acid	17.50	30.8	24.15	
$T_{3:} P_1 + 4\%$ sucrose	20.43	35.16	27.79	
$T_{4:} P_1 + 4\%$ boric acid	14.5	29.56	22.02	
T ₅ : $P_2 + 10\%$ sucrose	23.03	40.23	31.63	
$T_{6:} P_2 + 2\%$ boric acid	22.16	39.16	30.66	
$T_{7:} P_2 + 4\%$ sucrose	24.00	42.36	33.18	
$T_{8:} P_2 + 4\%$ boric acid	21.33	37.23	29.28	
$T_{9:} P_3 + 4\%$ sucrose	27.26	54.70	40.98	
$T_{10:} P_3 + 4\%$ boric acid	25.00	51.23	38.11	
T _{11:} Control	34.50	59.66	47.08	
Mean	22.62	40.43		
S.Em ±	0.36	0.38		
CD at (0.01)	1.01	1.07		

Table 1: Effect of different pre-cooling methods on physiological loss in weight (%) in buds of tuberose cv. Prajwal stored at 10 °C

P1: Pre-cooled at 4 °C, P2: Hydro-cooled at 4 °C, P3: Gel ice pack

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Table 2: Effect of different pre-cooling methods on respiration rate (ml CO2kg/hr) in buds of tuberose cv. Prajwal stored at 10°C

	Respiration (mlCO2kg/hr) Days of storage			
Treatments			Mean	
	2	4		
$T_{1:} P_1 + 10\%$ sucrose	90.33	119.33	104.83	
T_2 : $P_1 + 2\%$ boric acid	95.2	113.93	104.56	
$T_{3:} P_1 + 4\%$ sucrose	110.23	120.16	115.19	
$T_{4:} P_1 + 4\%$ boric acid	89.16	94.43	91.79	
$T_5: P_2 + 10\%$ sucrose	123.93	134.43	129.18	
$T_{6:} P_2 + 2\%$ boric acid	117.433	126.26	121.84	
$T_7: P_2 + 4\%$ sucrose	126.2	135.2	130.7	
$T_{8:} P_2 + 4\%$ boric acid	113.23	122.43	117.83	
T _{9:} P ₃ + 4% sucrose	142.36	153.33	147.84	
$T_{10:} P_3 + 4\%$ boric acid	134.36	144.5	139.43	
T _{11:} Control	154.16	167.56	160.86	
Mean	117.78	130.14		
S.Em ±	0.43	0.33		
CD at (0.01)	1.2	0.78		

P1: Pre-cooled at 4 °C, P2: Hydro-cooled at 4 °C, P3: Gel ice pack

Table 3: Effect of different pre-cooling methods on wilting (%) in buds of tuberose cv. Prajwal stored at 10°C

	Wiltin	Mean	
Treatments	Days of		
	2	4	
T_1 : $P_1 + 10\%$ sucrose	32.41	43.5	37.41
T_2 : $P_1 + 2\%$ boric acid	26.33	41.33	33.83
$T_3: P_1 + 4\%$ sucrose	29.41	42.41	36.45
T_4 : P ₁ + 4% boric acid	22.25	38.5	30.37
$T_{5:} P_2 + 10\%$ sucrose	38.25	46.5	42.37
$T_{6:} P_2 + 2\%$ boric acid	36.25	45.28	40.76
$T_{7:} P_2 + 4\%$ sucrose	39.166	48.08	43.62
$T_{8:} P_2 + 4\%$ boric acid	34.58	44.25	39.41
$T_{9:} P_3 + 4\%$ sucrose	42.36	52.25	47.30
$T_{10:} P_3 + 4\%$ boric acid	41.25	50.25	45.75
T ₁₁ : Control	43.25	60.25	51.75
Mean	35.04	46.60	
S.Em ±	0.27	0.26	
CD at (0.01)	0.78	0.74	

P1: Pre-cooled at 4 °C, P2: Hydro-cooled at 4 °C, P3: Gel ice pack

Table 4: Effect of different pre-cooling methods on shelf life, fragrance score and freshness index of tuberose cv. Prajwal stored at 10 °C

Treatments	Shelf life
T_1 : $P_1 + 10\%$ sucrose	3.66
T_2 : P_1 + 2% boric acid	3.33
$T_3: P_1 + 4\%$ sucrose	3.00
$T_4: P_1 + 4\%$ boric acid	4.00
$T_5: P_2 + 10\%$ sucrose	3.00
$T_6: P_2 + 2\%$ boric acid	2.66
$T_7: P_2 + 4\%$ sucrose	3.00
$T_8: P_2 + 4\%$ boric acid	3.00
$T_9: P_3 + 4\%$ sucrose	2.00
T_{10} : $P_3 + 4\%$ boric acid	2.00
T ₁₁ : Control	1.00
Mean	2.78
S.Em ±	0.35
CD at (0.01)	0.98

P1: Pre-cooled at 4°C, P2: Hydro-cooled at 4°C, P3: Gel ice pack

Conclusion

On the basis of results obtained in the present investigation on different post-harvest treatments used for extending shelf life of tuberose florets during storage and transport, it can be concluded that, pre-cooling at 4 $^{\circ}$ C proved is the method followed by hydro cooling and gel ice pack and with respect vase life solutions, sucrose and boric acid were found

effective in maintaining freshness of florets by lowering the rate of weight loss, respiration, and wilting thus helping in extending the shelf life of tuberose florets.

Acknowledgments

We are thankful to the Department of Post-Harvest Technology, College of Horticulture, University of Horticultural Sciences, Bagalkot, Karnataka, India for providing the laboratory facilities and technical support.

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