



RESEARCH PAPER

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Influence of aluminium sulfate and copper sulfate on some characteristic in *Rosa hybrida*

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Abstract

Flowers play a vital role in angiosperm reproduction; they are often pigmented and or perfumed to attract pollinators. However, despite its irreplaceable ecological role, the flowers are energetically expensive to maintain beyond their useful life, and therefore have a limited life-span that is usually taken away after pollination; causing senescence syndrome. Aluminum sulfate can decrease cut rose petal acidity and cause fixation of anthocyanin pigments and increase cut rose flowers vase. The role of aluminum sulfate to increase the vase life of cut flowers is not limited to lowering the pH of vase solution. The experiment was conducted at the research laboratory of education complex of zahedan (in iran). Laboratory lighting was provided by fluorescent lamps. The field experiment was laid out in randomized complete block design with factorial design with four replications. Analysis of variance showed that the effect of aluminium sulfate and copper sulfate on all characteristic was significant.

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Introduction

Flowers play a vital role in angiosperm reproduction; they are often pigmented and or perfumed to attract pollinators. However, despite its irreplaceable ecological role, the flowers are energetically expensive to maintain beyond their useful life, and therefore have a limited life-span that is usually taken away after pollination; causing senescence syndrome. Senescence of flower is a complex process, so often researchers mainly concentrate on changes occurring during petal senescence. Petals provide an excellent model system for the study of fundamental aspects of senescence (Rogers, 2006; Desai *et al.*, 2012).

Senescence is a highly regulated final event of flower development that bears hallmarks of programmed cell death (PCD), resulting in colour changes, petal wilting, abscission of whole flower and flower parts with various physiological, biochemical and ultrastructural changes (Voleti *et al.*, 2000; Wagstaff *et al.*, 2003; Jones *et al.*, 2005; Tripathi & Tuteja, 2007; Seo *et al.*, 2009; Ichimura, 2010; Shahri, 2011). Roses are one of the most important cut flowers in the world (Şirin, 2011) and extremely perishable (Figueroa *et al.*, 2005) as well as other cut flowers such as *Eustoma grandiflorum* (Hojjati *et al.*, 2007; Farokhzad *et al.*, 2005), *Gerbera jamesonii* (Nair *et al.*, 2003). Short postharvest vase life is one of the most important problems in cut flowers (Zamani *et al.*, 2011).

So consider to maintaining postharvest quality of cut flowers is critical for preventing offlower post harvest losses. Senescence which is the main factor affecting on flower quality can be induced by several pre and post-harvest factors e.g., water stress (Sankat and Mujaffar, 1994), amount of carbohydrates (Coorts, 1973; Ketsa, 1989), microorganisms (Van Doorn and Witte, 1991), ethylene effects (Wan & Miller, 2003) as documented in carnation and roses (Mayak and Halevy, 1980; Halevy and Mayak, 1981; Quesada and Valpuesta, 2000) and *Lisianthus* (Farokhzad *et al.*, 2005; Hojjat *et al.*, 2007) and cultivar differences, season, development stage at harvest and cultivated

conditions (Doel and Wilkins, 1999). Application of some germicides has been suggested to prevent rapid proliferation of microorganisms and to decrease the longevity of cut flowers. Cut flower species respond to germicides variously.

$Al_2(SO_4)_3$ has been recommended for maintaining the vase life of several cut flowers (Liao *et al.*, 2001) and is used as an antimicrobial compound in commercial preservative solutions (Ichimura *et al.*, 2006). Aluminum sulfate acidifies vase solution, diminishes bacterial proliferation and enhances water uptake (Tjeerd and Jaap, 2003; Hassanpour Asil *et al.*, 2004). Roses and can be caused by physiological occlusion due to plant itself, microorganisms or air embolism (Van Doorn *et al.*, 1989).

Also aluminum sulfate can decrease cut rose petal acidity and cause fixation of anthocyanin pigments and increase cut rose flower's vase life (Put Henriette *et al.*, 1992; Tjeerd and Jaap, 2003; Hassanpour Asil *et al.*, 2004). The role of aluminum sulfate to increase the vase life of cut flowers is not limited to lowering the pH of vase solution. Its effect is based at least in part, on its action as an antimicrobial agent in the solution (Liao *et al.*, 2001).

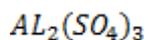
More study is necessary to determine the effect of aluminum sulfate on vase life of cut flowers, specially cut roses as one of the most important cut flowers in the world. Van Meetereu *et al.* (2001) suggested that it must be used a combination of calcium chloride, sodium carbonate and copper sulfate solution as a basic standard for the preservative solution. Motivation and aims of the study are Influence of aluminium sulfate and copper sulfate on some characteristic in rosa hybrid.

Material and methods

Location of experiment

The experiment was conducted at the research laboratory of education complex of zahedan (in iran). Temperature lab were 25 C°. Laboratory lighting was provided by fluorescent lamps and humid 70%.

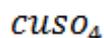
Product of Aluminium sulfate and Copper sulfate



$$Al_2(SO_4)_3 = 1l \times \frac{75mg}{l} \times \frac{1g}{1000mg} \times \frac{1molAL}{27grAL} \times \frac{1molAl_2(so_4)_3}{2molAl} \times \frac{342.174}{1molAl_2(so_4)_3} = 1/850gr$$

$$Al_2(SO_4)_3 = 1l \times \frac{150mg}{l} \times \frac{1g}{1000mg} \times \frac{1molAL}{27grAL} \times \frac{1molAl_2(so_4)_3}{2molAl} \times \frac{342.174}{1molAl_2(so_4)_3} = 3/7gr$$

$$Al_2(SO_4)_3 = 1l \times \frac{225mg}{l} \times \frac{1g}{1000mg} \times \frac{1molAL}{27grAL} \times \frac{1molAl_2(so_4)_3}{2molAl} \times \frac{342.174}{1molAl_2(so_4)_3} = 5/55gr$$



$$CuSO_4 = 1L \times \frac{5mg}{l} \times \frac{1g}{1000mg} \times \frac{1molcu}{63.5grcu} \times \frac{1molcuso}{1molcu^{+2}} \times \frac{249.62}{1mrcuso} = \frac{248.1}{6350} = 0.019$$

$$CuSO_4 = 1L \times \frac{10mg}{l} \times \frac{1g}{1000mg} \times \frac{1molcu}{63.5grcu} \times \frac{1molcuso}{1molcu^{+2}} \times \frac{249.62}{1mrcuso} = \frac{248.1}{6350} = 0.038gr$$

$$CuSO_4 = 1L \times \frac{15mg}{l} \times \frac{1g}{1000mg} \times \frac{1molcu}{63.5grcu} \times \frac{1molcuso}{1molcu^{+2}} \times \frac{249.62}{1mrcuso} = \frac{248.1}{6350} = 0.057gr$$



Fig. 1. Product of Aluminium sulfate and Copper sulfate.

Field experiment

The field experiment was laid out in randomized complete block design with factorial design with four replications.

Treatments

Treatments consisted the main factor in seven levels consisted of days (2, 4, 6, 8, 10, 12 and 14 day) and sub factor involves of chemical compounds in four

levels: Aluminium sulfate (0, 75, 150 and 225(mg/l)) and copper sulfate (0, 1, 2 and 3 (gr/l)).



Fig. 2. Treatments of experiments.

Data collect

Data collected were subjected to statistical analysis by using a computer program MSTATC. Least Significant Difference test (LSD) at 5 % probability level was applied to compare the differences among treatments` means.

Results and discussion

Relative weight of flower

Analysis of variance showed that the effect of aluminium sulfate on relative weight of flower was

significant (Table 1). The maximum of relative weight of flower (82.89) of 150(mg/l) was obtained (Table 2). Analysis of variance showed that the effect of copper sulfate on relative weight of flower was significant

(Table 1). The maximum of relative weight of flower (83.92) of treatments 10 gr.l⁻¹ was obtained (Table 2). The minimum of relative weight of flower (13.86) of treatments control was obtained (Table 2).

Table 1. Anova analysis of the rosa hybrida affected by aluminium sulfate and copper sulfate.

S.O.V	df	Relative weight of flower	Fresh weight of flower	Absorption of solution	Life of flower
Aluminium sulfate	3	311.76**	133.65**	0.23**	5.16*
Copper sulfate	3	695.51**	87.78**	0.47**	7.83**
Aluminium sulfate* Copper sulfate	9	115.65*	30.03**	0.26**	3.88*
Error	48	52.7	8.59	0.05	1.62
Cv	-	9.29	16.96	15.35	10.62

*, **, ns: significant at p<0.05 and p<0.01 and non-significant, respectively.

Fresh weight of flower

Analysis of variance showed that the effect of aluminium sulfate on fresh weight of flower was significant (Table 1). The maximum of fresh weight of flower (19.72) of 225(mg/l) was obtained (Table 2). Analysis of variance showed that the effect of copper sulfate on fresh weight of flower was significant (Table 1). The maximum of fresh weight of flower (18.77) of treatments 15 gr.l⁻¹ was obtained (Table 2). The minimum of fresh weight of flower (13.86) of treatments control was obtained (Table 2).

Analysis of variance showed that the effect of copper sulfate on absorption of solution was significant (Table 1). The maximum of absorption of solution (1.78) of treatments 15 gr.l⁻¹ was obtained (Table 2). The minimum of absorption of solution (11.37) of treatments control was obtained (Table 2).

Table 2. Comparison of different traits affected by aluminium sulfate and copper sulfate.

Treatment	Relative weight of flower	Fresh weight of flower	Absorption of solution	Life of flower
aluminium sulfate				
0 (mg/l)	72.26b	14.005b	1.41c	11.25b
75 (mg/l)	77.82a	15.69b	1.58ab	12.00ab
150 (mg/l)	82.89a	19.69a	1.68a	12.62a
225 (mg/l)	79.26a	19.72a	1.46bc	12.12ab
copper sulfate				
0 (gr/l)	68.60b	13.86b	1.42b	11.37b
5 (gr/l)	80.07a	17.64a	1.42b	11.75b
10 (gr/l)	83.92a	18.83a	1.52b	11.87b
15 (gr/l)	79.63a	18.77a	1.78a	13.00a

Any two means not sharing a common letter differ significantly from each other at 5% probability.

Life of flower

Analysis of variance showed that the effect of aluminium sulfate on life of flower was significant (Table 1). The maximum of life of flower (12.62) of 150 (mg/l) was obtained (Table 2). Analysis of variance showed that the effect of copper sulfate on life of flower was significant (Table 1). The maximum of life of flower (13.00) of treatments 15 gr.l⁻¹ was obtained (Table 2). The minimum of life of flower (11.37) of treatments control was obtained (Table 2).

Absorption of solution

Analysis of variance showed that the effect of aluminium sulfate on absorption of solution was significant (Table 1). The maximum of absorption of solution (1.68) of 150 (mg/l) was obtained (Table 2).

References

Desai R, Patel R, Mankad A. 2012. Petal senescence in cut Tagetes erecta L. flowers: Role of phenolics. International Journal of Science Environment and Technology. **1**, 485 – 490.

Ichimura K, Shimizu H, Goto R. 2009. Ethylene production by gynoeceium and receptacle is associated with sepal abscission in cut Delphinium flowers. Postharvest Biology and Technology. **52**, 267–272.

Jones ML, Chaffin GS, Eason JR, Clark DG. 2005. Ethylene sensitivity regulates proteolytic

activity and cysteine protease gene expression in petunia corollas. *Journal of Experimental Botany*. **56**, 2733-2744.

Rogers HJ. 2006. Programmed cell death in floral organs: How and why do flowers die. *Annals of Botany journal*. **97**, 309–315.

Shahri W, Tahir I, IslamST, Bhat MA. 2011. Physiological and biochemical changes associated with flower development and senescence in so far unexplored *Helleborus orientalis* Lam. cv. Olympicus. *Physiology and Molecular Biology of Plant*. **17**, 33–39.

Seo S, Kang SW, Shim IS, KimW, Fujihara S. 2009. Effects of various chemical agents and early ethylene production on floral senescence of *Hibiscus syriacus* L. *Plant Growth Regulation*. **57**, 251–258.

Tripathi SK, Tuteja N. 2007. Integrated signaling in flower senescence. *Plant signaling and Behavior. Journal of Experimental Botany*. **2**, 437-446.

Voleti SR, Singh V, Arora A, Singh N, Kushwaha SR. 2000. Physiology of flower senescence in floriculture crops. In A. Hemantaranjan (Ed.). *Advances in Plant Physiology*. **12**, 423-439.

Wagstaff C, Leverentz MK, Griffiths G, Thomas B, Chanasut U, Stead AD. 2002. Cysteine protease gene expression and proteolytic activity during senescence of *Alstroemeria* petals. *Journal of Experimental Botany*. **53**, 233-240.

Doel JM, Wilkins HF. 1999. *Floriculture: principles and species*. Prentice Hall, inc New Jersey. *Physiology and Molecular Biology of Plant*. **17**, 33–39.

Farokhzad A, Khalighi A, Mostofi Y, Naderi R. 2005. Role of Ethanol in the Vase Life and Ethylene Production in Cut *Lisianthus* (*Eustoma grandiflorum* Mariachii. cv. Blue) Flowers. *Advances in Plant Physiology*. **1**, 309-312.

Hojjati Y, Khalighi A, Farokhzad AR. 2007. Chemical Treatments of *Eustoma* Cut Flower Cultivars for Enhanced Vase Life. *Journal of Agriculture and Social Sciences*. **3(3)**, 75-78.

Halevy AH, Mayak S. 1981. Senescence and postharvest physiology of cut flowers. Part 2. *Hortic. Annals of biological reaserach*. **3**, 59-143.

Ketsa S. 1989. Vase life characteristics of inflorescences of *dendrobium* Pompadour. *Journal of Experimental Botany*. **64**, 611–643.

Nair SA, Singh V, Sharma TV. 2003. Effect of chemical preservatives on enhancing vase-life of gerbera flowers. *Advances in Plant Physiology*. **41**, 56-58.

Vandoorn WG, WitteYE. 1991. The mode of action of bacteria in the vascular occlusion of cut rose flowers. *Acta Horticulture journal*. **298**, 165-176.

Wu M, Lorenzo Z, Saltveit ME, Reid MS. 1992. Alcohol and carnation senescence. *Journal of Experimental Botany*. **27**, 136-138.

Zamani S, Kazemi M, Aran M. 2011. Postharvest life of cut rose flowers as affected by salicylic acid and glutamin. *World applied science Journal*. **12(9)**, 1621-1624.

Şirin U. 2011. Effects of different nutrient solution formulations on yield and cut flower quality of gerbera (*Gerbera jamesonii*) grown in soilless culture system. *African Journal of Agricultural Research*. **6(21)**, 4910-4919.

Ichimura K, Ueyama S. 1998. Effect of temperature and application of aluminum sulfate on the postharvest life of cut rose flowers. *Bulletin of the National Research Institute of Vegetables Ornamental Plants and Tea. World applied science Journal* **13**, 51- 60.

Liao LJ, Lin YH, Huang KL, Chen WSh, Cheng YM. 2000. Postharvest life of cut rose flowers as affected by silver thiosulfate and sucrose. African Journal of Agricultural Research. **41**, 299 – 303.

Put Henriette MC, Clerkx Anke CM, Boekestein A. 1992. Aluminum sulphate restricts migration of *Bacillus subtilis* in xylem of cut roses: a scanning electron microscope study. Scientia Horticulture journal. **51**, 261- 274.

Van Doorn WG, Schurer K, De Witte Y. 1989. Role of endogenous bacteria in vascular blockage of cut rose. Journal of Plant Physiology. **134**, 375-381.

Hassanpour Asil M, Hatamzadeh A, Nakhai F. 2004. Study on the effect of temperature and various chemical treatments to increase vase life of cut rose flower “Baccara”. Research Journal of Guilan Agriculture Faculty. **1(4)**, 121- 129.