Effect of Salicylic Acid on the Growth, Metabolic Activities and Oil Content of Basil and Marjoram

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ABSTRACT

The response of sweet basil (Ocimum basilicum L.) and marjoram (Majorana hortensis) plants to foliar application of salicylic acid (SA) at 10⁻⁵, 10⁻⁴ and 10⁻³ M was determined in pot experiments conducted during 2004 - 2005. SA increased plant height, number of (branches, nodes & leaves) per plant, leaf area, fresh and dry weight of herbs, total carbohydrates, crude protein, total amino acids, free proline, photosynthetic pigments as well as microelement content and uptake up to 10^{-4} M relative to un-treated controls and decreased thereafter in both basil and marjoram. All SA treatments enhanced putrescine, spermidine as well as total polyamines contents, while reduced the level of spermine in both plants. Oil percentage and yield per plant for three cuttings also increased about two fold on a fresh weight basis with SA application at 10^{-4} M in case of basil and 10⁻³ M in marjoram relative to un-treated controls. GC/MS revealed that common components of Ocimum basilicum essential oil under all treatments were linalool (46.63 - 43.32%), methyl eugenol (13.83 - 5.68%), 1, 8 - cineol (13.20 -4.43%), eugenol (12.64-7.16%) and α -cadinol (9.59 - 4.46%). SA at 10⁻⁴ M increased the production of top quantity and quality of basil oil to the fragrance and food industries by increasing the percentage of eugenol and antioxidant activity in the herb. On the other hand, the marjoram essential oil contains cis-sabinene hydrate(37.50 - 14.27%), terpinen- 4-ol (24.33 -13.99%), p-cymene (18.21 - 2.29%), sabinene (17.69 - 4.11%), γ -terpinene (10.64 - 4.77) in addition to α -terpineol (5.52 -3.96%), trans-sabinene hydrate (5.45 - 8.19%), α-terpinene (2.41 - 0.00%) and β-carvophyllene (3.82 - 1.76%). Moreover, SA at 10^{-5} M and 10^{-3} M improved oil quality by increasing the level of sabinene accompanied by a decrease in the proportion of *cis*-sabinene hydrate relative to controls. The data suggest that in both species, SA treatment especially at 10^{-4} M may have higher adaptive capacity to stress, originating from promoting polyamines synthesis and better osmotic adjustment.

Key Words: Ocimum basilicum; Majorana hortensis; Salicylic acid; Vegetative growth; Free proline; Polyamines; Essential oils

INTRODUCTION

Aromatic plants represent a renewable source of flavoring substances, which can be used in the food, perfumery and pharmaceutical industries. The family lamiaceae includes large number of volatile oil plants and the most important members are sweet basil (Ocimum basilicum) and marjoram (Majorana hortensis). Sweet basil is one of the most widespread spices in the world and its dried leaves are used commonly for flavoring many food products. Salicylate content of spices was bio-available, and may contribute to the low cancer incidence in rural India (Paterson et al., 2006). Basil contains a wide range of essential oils rich in phenolic compounds (Simon et al., 1990; Phippen & Simon, 2000) and polyphenols such as flavonoids and anthocyanins (Phippen & Simon, 1998). The volatile basil oil is used in pharmaceutical and perfume industry. On the other hand, marjoram is used world wide as a spice and crude drug. The quality of the product is usually determined by chemical analysis, whereas in food industry applications sensory tests are also practiced. Marjoram essential oil components mainly revealed differences between plant species. In addition, sweet basil essential oil and sweet marjoram water extracts possess high antioxidant activity (Triantaphyllou *et al.*, 2001; Juliani & Simon, 2002). Epidemiological studies have suggested positive associations between the consumption of phenolic-rich foods or beverages and the prevention of disease due to the presence of antioxidant components such as phenolics (Rice-Evans *et al.*, 1996; Scalbert & Wuilliamson, 2000).

The chemical composition of basil and marjoram oils is very variable. The chemovarieties and the environmental conditions caused to this fact. The major components of sweet basil oils from different origins were found to be linalool, eugenol, estragole and methyl cinnamate (Akgül, 1989; Juliani & Simon, 2002). Contrarity, several workers reported that essential oil components of marjoram were terpinen-4-ol, gamma-terpinene, trans-sabinene hydrate, linalool, thujanol, terpinolene and thymol (El-Ghorab *et al.*, 2004).

Salicylic acid (SA) naturally occurs in plants in very low amounts. It has been identified as an important signaling element involved in establishing the local and systemic disease resistance response of plants after pathogen attack (Alvarez, 2000). After a pathogen attack, SA levels often increases and induces the expression of pathogenesisrelated proteins and initiates the development of systemic acquired resistance and hypersensitive response (Grüner et al., 2003; Kachroo et al., 2005). SA, jasmonic acid (JA), and ethylene-dependent signaling pathways regulates plant responses to both abiotic and biotic stress factors. In Arabidopsis ecotype Cvi-0, SA and JA signaling have a role in influencing O₃ -induced cell death and an antagonistic relationship between JA- and SA-signaling pathways in controlling the magnitude of O3 -induced cell death was demonstrated (Mulpuri et al., 2000). During O3 exposure, transgenic plants with a phenotype of reduced O₃ -induced ethylene production accumulated less SA than did wild-type plants. Ethylene promotes SA accumulation by regulating the expression of the *chorismate mutase* and phenylalanine ammonia-lyase genes in O3 -exposed tobacco (Ogawa et al., 2005). Young maize plants exhibited increased cold tolerance upon treatment with SA or aspirin (Janda et al., 1999). Exogenously added SA also increased the heat tolerance of mustard (Dat et al., 2000). Moreover, SA treatments at 0.5 mM strongly or completely suppressed the Cd-induced up-regulation of the antioxidant enzyme activities of barley (Metwally et al., 2003). SA has a direct physiological effect through the alteration of antioxidant enzyme activities. Certain enzymes were activated by SA treatment, while others like catalase, were inhibited. Catalase seems to be a key enzyme in SA-induced stress tolerance, as it was inhibited by binding in several plant species (Chen et al., 1993; Conrath et al., 1995). In grapevine (Vitis amurensis), SA-responsive cis-acting elements and the W-boxes are important for the SA induction of the VCH 3 promoter, which might have a potential use in plant genetic engineering (Hai-Yan et al., 2006). SA induces flowering, increase flower life, retards senescence and increases cell metabolic rate. The sustained level of SA may be a prerequisite for the synthesis of auxin and/or cytokinin (Metwally et al., 2003), yet the influence of these new plant growth hormone on essential oil production and secondary metabolism in general, has received little attention.

The present study was undertaken to examine the effects of foliar application of SA on growth, chemical composition, essential oil production and composition as well as nutrient uptake of *Ocimum basilicum* and *Majorana hortensis*, another common essential oil producing species.

MATERIALS AND METHODS

A pot experiment was conducted at the experimental farm of Helwan University, Cairo, Egypt during 2004/2005. Seeds of *Ocimum baslicum* L. and *Majorana hortensis* (kindly provided by Horticultural Research Institute, Agricultural Research Center, Ministry of Agriculture, Giza, Egypt) were sown in beds on 1st February. Uniform 45 days old seedlings were transplanted per earthenware pot (40 cm in diameter) of 15 kg of sandy loam soil. The pots were arranged in complete randomized blocks design with four

treatments, four replicates per treatment and each replicate represented by 2 plants. Physical and chemical properties of the soil used in the experiment were evaluated according to Jackson (1973): The soil type was sandy loam in texture with pH of 8.7, organic matter 0.35%, electrical conductivity 0.46 m. mhos cm⁻¹. The soil analysis, 2.96% containing CaCO₃, available 4.46, 23.46, 169 and .32.2 mg 100^{-1} g soil of P, K, Mg and Na, respectively and also available 7.2, 9.4, 2.80 and 4.82 ppm soil of Fe, Mn, Cu and Zn, respectively.

Fertilization was carried out for each pot at proportion of 1 g ammonium nitrate (33.5% N), 2 g calcium superphosphate $(15.5\% P_2O_5)$ and 1 g potassium sulphate (48% K₂O). These fertilizers were applied in two doses at 60 and 75 days after planting and repeated after the first and second cuts. Irrigation was regularly carried out at intervals according to weather conditions to keep the moisture content of the soil to field capacity. Plants were foliarly sprayed twice with SA $(10^{-5}, 10^{-4} \& 10^{-3} M)$, the first spray was done after 75 days after sowing, while the second spray was applied one week later after the first one. Treatments were repeated again 2 weeks after the first and second cuts. Control plants were sprayed with distilled water and the volume of the spraying solution was maintained just to cover completely the plant's foliage till drip. Three cuts were taken 2, 4 and 6 month after transplantation.

At full bloom, the plant herbage was cut 10 cm above the soil surface and data on plant height, number of branches, nodes, leaves and spikes per plant as well as fresh and dry weight of herb for plant growth parameter for the 3 cuts was recorded. Plant samples were dried in an oven with drift fan at 70°C until constant dry weight was obtained. Representative fresh samples were taken from each treatment for determination of essential oil content and oil constituents.

Quantitative determination of *O. baslicum* and *M. hortensis* essential oil from the fresh samples was achieved by hydro-distillation at first, second and third cuts. Essential oils from the first cut were separated and analyzed qualitatively by GC/MS at National Research Centre, Dokki, Cairo. The GC analysis was carried out using Varian-3400 GC equipped with a DB-5 fused silica capillary column. Mass spectrometer was a Varian–Finnigan SSQ 7000.

Total carbohydrates were determined in the dried samples of first cut, using phenol sulphuric method (Dubois *et al.*, 1956). Total nitrogen was determined using the modified Micro-Kjeldahl method according to AOAC (1980). Macro and microelements were determined after wet digestion. Phosphorus was determined by using vanadate-molybdate method (Jackson, 1973). Potassium was measured by flame photomtere, Eppendorff. Microelements were estimated by using Inductively Coupled Spectrometry Plasma (ICP) Model Ultima 2-Jobin Yvon. The method of Müting and Kaiser (1963) as modified by EL-Araby (1987) was used for estimation of total amino acids. Free proline was determined according to the method described by Bates *et al.* (1973). Photosynthetic pigments of fresh leaves were determined and calculated according to Wettstein (1957).

Fresh herb of sweet basil and marjoram plants developed from the third cut were used for the extraction of polyamines according to (Flores & Galston, 1982) as adopted by (Deabes, 2000) and HPLC was used for measurement of individual polyamines (Smith *et al.*, 1985). Data obtained were subjected to analysis of variance and the values of least significant differences (LSD at 5% level) were calculated to compare the means of different treatments according to (Snedecor & Cochran, 1980).

RESULTS AND DISCUSSION

Growth criteria. Salicylic acid (SA) increased growth in terms of plant height, number of branches, spikes and leaves per plant, leaf area and fresh and dry weights of herb in both species during three cutting (Fig. 1 & 2). This promoting effect was at maximal at 10⁻⁴ M SA for all growth parameters except plant height and leaf area, showing maximum values at 10⁻⁵ M. Similarly, treatment of pepper seeds with SA and sulfosalicylic acid (both at 10^{-4} M) was effective in inducing highest leaf number, a tallest seedling and more plant fresh and dry weight, compared to lower (10^{-3} M) and higher (10^{-5} M) , which showed a negative effect on seedling final fresh and dry weight (Mendoza et al., 2002). Kord and Hathout (1992) found that foliar application of salicylaldehyde at 10⁻⁵ M stimulated different morphological and growth criteria of tomato plants but inhibitory effects were observed at 10⁻³ M. Spraving mungbean with SA, significantly enhanced the total seed vield (Jaiwal & Bhambie, 1989). Moreover, the increase in plant height was due to increase in number of internodes. while the increase in the fresh and dry weights of basil and marjoram at low concentration might be attributed to an increase in number of branches and leaves as well as leaf area, leading to increased photosynthetic activity. Hamada (1998) found that soaking of wheat grains in 100 ppm aspirin showed a marked increase in the fresh and dry matter gain in shoots and roots through the regulation of some photosynthetic reactions. A significant increase in leaf area per basil and marjoram was also noticed at 10^{-5} M of SA but markedly decreased with 10^{-4} and 10^{-3} M, especially at 1st and 2nd cuttings. Large leaf size of some types of basil with industrial values coupled with the high content of essential oils has potential use in herbal tea trade (Simon, 1992).

Oil content. Foliar spray of both basil and marjoram with SA increased essential oil yield per plant for the three cuts compared with their corresponding controls and also increased oil % gradually specially with the second cut in basil but showing the opposite response in control marjoram (Fig. 3). Similarly, Bottcher *et al.* (1999) found that *M. hortensis* herbs (1st cut) contained 22% more essential oils

Fig. 1. Effect of foliar spray with salicylic acid on the vegetative growth of *Ocimum basilicum* L plants. Vertical bars represent LSD at 5%



than physiologically younger plants (2^{nd} cut). The highest mean values of essential oil percentage of the three cuts when pooled together increased significantly with about (26.52% in basil & 36.06% in marjoram), while on a per plant basis oil yield increased by 90.33% and 100.09% relative to control by SA application at 10^{-4} M in basil and 10^{-3} M in marjoram, respectively. The increment in oil yield might be due to the increase in vegetative growth, nutrients uptake or changes in leaf oil gland population and monoterpins biosynthesis. In accordance, total N and P uptake as well as essential oil yield of *Coriandrum sativum* increased with increasing rates of N and P application up to 60 and 45 kg ha⁻¹, respectively (Tiwari & Banafar, 1995).

The mean values of oil percentage and yield per plant of the three cuts when pooled together increased 5.49 and 26.66%, respectively over control plants by SA application at 10^{-3} M. Ram *et al.* (1997) reported that SA application (100 ppm) had no effect on the herbage and essential oil yields in *Pelargonium graveolens, Mentha arvensis* and *Cymbopogon martini.* They concluded that either the synthesis of essential oil constituents occurs constitutively, without the intervention of SA or the amount of SA required for the induction of synthesis of essential oil constituents are already available in these plants.



Fig. 2. Effect of foliar spray with salicylic acid on the vegetative growth of *Origanium hortensis* plants. Vertical bars represent LSD at 5%

Fig. 3. Effect of foliar spray with salicylic acid on the oil content of *Ocimum basilicum* L and *Origanium hortensis* plants. Vertical bars represent LSD at 5%



Main components of the essential oil. The *Ocimum* basilicum essential oil appeared to be complex and riche in flavour notes and have higher proportions of linalool representing 46.63% in the oil, but intermediate proportions of methyl eugenol (13.83%), 1, 8-cineol (13.20%), eugenol (7.16%) and α -cadinol (4.46%). Juliani and Simon (2002) found that the major oil components of sweet basil was linalool, eugenol and germcrene D accompanied by lesser amount of α -pinene, camphene, sabinene, β -pinene,

myrcene, 1, 8 -cineol, cis-ocimene, terpinolene, camphore, borneol, 4 -terpinol, α -terpineol, bornvl acetate, α -cubebene, α -copaene, β -elemene, methyleugenol, β -caryophyllene, α humulene, a-amorphene, biclogermacrene, guaiene and cadinene. Our results indicated that most percentage of the oil was oxygenated compounds with much higher quantities of linalool and 1, 8 -cineol (Table I & II), but also other compounds are nearly similar to our findings. The oil of Egypt basil can be classified as Linalool Rich Type (Nacar & Tansi, 2000). However, there was a pronounced difference in the essential oil of Ocimum basilicum content due to SA application. Common components of essential oil in all treatments were linalool (46.63 - 43.32%), methyl eugenol (13.83 - 5.68%), 1, 8 -cineol (13.20 - 4.43%), eugenol (12.64 - 7.16%) and α -cadinol (9.59 - 4.46%). In addition, SA application increased the eugenol level from 7.13% in control to 12.64% at maximal yield with SA at 10⁻ ³ M and was accompanied by an increase in 1, 8 -cineol content (from 13.20 to 15.31%), all other components exhibited a slight change. In this regard, the relative percentage of eugenol was correlated with the antioxidant activity of Sweet basil essential oil in two assays (Juliani & Simon, 2002). Therefore, a high percentage of eugenol in the SA treated herb is of particular interest in basil, which could constitute new sources of antioxidant phenolics in the diet.

On the other hand, the proportions of terpinen-4-ol, *cis*-sabinene hydrate, p-cymene and γ -terpinene of control marjoram essential oil are closer to those of marjoram grown on Reunion Island (Vera & Ming, 1999). Under SA application, marjoram exhibited the same main components. Possible changes in the proportions of the major components were monitored under the influence of SA; the essential oil was rich in cis-sabinene hydrate (37.50 -14.27%), terpinen-4-ol (24.33 - 13.99%), p-cymene (18.21 -2.29%), sabinene (17.69 - 9.79%), γ-terpinene (10.64 -4.77), α -terpineol (5.52 - 3.96%), trans-sabinene hydrate (8.19 - 5.45%), α -terpinene (2.41 - 0.00%) and β carvophyllene (3.82 - 1.76%). SA at 10^{-5} M and 10^{-3} M produced an increase in the level of sabinene, which was accompanied by a decrease in the proportion of cis-sabinene hydrate relative to controls (Table II). Similar changes were evident in Majorana hortensis by salt stress (El-Keltawi & Croteau, 1987). On the other hand, the levels of numerous other monoterpenes including a-pinene, a-myrcene, aterpinene, p-cymene, y-terpinene as well as of the sesquiterpenes carvophyllene and α -humulene were increased. The level of cis-sabinene hydrate was concomitantly decreased relative to controls at 10⁻⁵ M and 10^{-3} M SA. At, 10^{-4} M the proportion of *cis*-sabinene hydrate, y-terpinene and terpinen-4-ol markedly increased with only a minor influence on α -terpinenes. The Egyptian marjoram oil is rich in the main component, terpinen-4-ol (Refaat et al., 1990).

Carbohydrate, protein, total amino acids and free proline. The effect of foliar application of SA on the total

Oil components (%)	Control	Salicylic acid			
_		10 ⁻⁵ M	10 ⁻⁴ M	10 ⁻³ M	
α-Pinene	0.45	0.46	0.05	0.58	
Camphene	0.04	0.08	-	-	
Sabinene	1.46	1.47	0.21	2.01	
α-myrcene	1.30	1.50	0.26	1.70	
Limonen	0.05	-	-	-	
1,8-cineol	13.20	14.44	4.43	15.31	
Ocimene	1.16	1.73	0.54	1.38	
γ-terpinene	0.37	0.33	0.29	0.29	
Linalool	46.63	45.62	46.63	43.32	
Camphor	0.76	0.60	0.26	0.47	
4-terpineol	0.48	0.69	0.46	-	
Borneol	-	-	-	0.44	
α-terpineol	0.79	1.18	0.98	1.44	
Endobornyl actate	0.97	1.45	1.63	1.46	
Methyl cinnamate	1.60	1.50	1.78	0.67	
β-caryophyllene	0.01	-	-	-	
α-copaene	0.01	0.01	0.07	0.01	
Eugenol	7.16	8.83	9.69	12.64	
Methyl eugenol	13.83	8.91	10.70	5.68	
α-bergamotene	-	-	1.78	-	
α-cubebene	0.01	0.07	0.11	0.02	
α-humulene	0.83	0.84	0.95	1.05	
δ -muurolene	-	0.09	-	0.01	
Germacrene D	0.96	0.40	2.64	0.78	
Bicycloge- rmacrene	0.59	0.68	1.59	0.74	
α-guaiene	0.23	0.43	-	0.50	
δ-cadinene	1.66	2.27	2.22	2.55	
α-muurolene	0.04	0.08	0.05	0.09	
Nerollidol	0.13	0.12	1.19	0.13	
Spathulenol	0.06	0.05	0.16	0.03	
Iso-spathulenol	0.04	0.04	0.11	0.04	
Carotol	0.50	0.61	1.18	0.71	
α-cadinol	4.46	4.94	9.59	5.21	
α-eudesmol	-	0.35	0.65	0.25	
α-bisabolol	-	0.05	0.09	0.08	
Farnesol	-	0.03	-	0.04	
Unidentified	0.22	0.15	0.71	0.37	

Table I. Effect of foliar spray with salicylic acid on the composition of essential oil of *Ocimum basilicum* L plants

carbohydrate content was accompanied by similar effects on crude proteins, total free amino acids and proline content of O. basilicum and M. hortensis at first cut (Table III & Fig. 4). Data revealed that maximum values of total carbohydrates and crude proteins of both species were obtained by foliar application of 10⁻⁴ M SA. In other studies, the highest increase in the level of the total soluble carbohydrate and proteins of wheat and tomato plants was obtained with 10^{-4} M and 10^{-5} M salicylaldehyde, respectively (Mohamed et al., 1989; Kord & Hathout, 1992). A similar trend of total free amino acid and proline content was noted in basil and marioram as a result of using 10^{-4} M SA. At high concentration, free proline protects membranes and proteins against the adverse effects of inorganic ions. Temperature extremes can act as an enzymestabilizing agent under salinity (Samaras et al., 1995; Demir & Koccacaliskan, 2001) and reduce peroxidative damage to the lipid membranes due to salt dependent oxidative stress (Jain et al., 2001). Present data suggest that in both species, SA has higher adaptative capacity to stress, originating from

Table II. Effect of foliar spray with salicylic acid on the composition of essential oil of *Origanum marjorana* plants

Oil components (%)	Control		Salicylic acid			
• • •		10 ⁻⁵ M	10 ⁻⁴ M	10 ⁻³ M		
α-pinene	1.59	1.92	0.96	2.34		
Sabinene	9.79	14.78	4.11	17.69		
α- myrcene	1.87	2.75	0.88	2.51		
Phellandrene	0.02	-	0.05	-		
a-terpinene	2.41	-	2.61	0.38		
p-cymene	6.28	17.68	2.29	18.21		
Γ-terpinene	4.77	9.68	5.21	10.64		
trans-sabinene hydrate	6.72	8.19	6.21	5.45		
cis-sabinene hydrate	37.50	14.83	40.63	14.27		
cis-2-p-menthen-1-ol	0.67	0.39	1.47	0.50		
Terpinen-4-ol	15.71	16.60	24.33	13.99		
α-terpineol	4.80	3.96	5.52	5.42		
Linalyl acetate	1.55	1.40	0.29	1.07		
Endobornyl actate	0.07	0.03	0.21	0.11		
α-terpinyl propionate	0.22	0.21	0.14	0.18		
Nerolidol	0.01	0.26	0.07	0.42		
Bicycloelemene	0.31	0.17	0.26	0.1 7		
Farnesyl acetate	0.07	-	0.04	-		
β-caryophyllene	2.63	3.82	1.76	2.96		
Aromadendrene	1.15	0.55	0.98	-		
α-humulene	0.19	0.24	0.15	0.60		
Leden	0.91	0.91	0.71	0.81		
α-cadinene	0.06	0.03	0.15	0.05		
Spathulenol	0.25	0.35	0.12	0.48		
α-elemene	0.11	0.18	-	0.35		
Unidentified compounds	0.34	1.07	0.85	1.40		

better osmotic adjustment.

Photosynthetic pigments. Foliar spray of SA increased photosynthetic pigments in both species under study. Here again SA applied at 10^{-4} M was the most effective (Table III & Fig. 4). These results are in agreement with those obtained by Khurana and Makeshwari (1978), who found that in *Spirodela Polyrrhiza* SA at 10^{-5} M stimulated total chlorophyll synthesis whereas 10^{-3} M has a reverse effect. Stomatal index and stomatal density of pepper seeds were negatively affected by treatment with SA and SSA (10^{-3} M). On the other hand, SA and SSA at 10^{-4} and 10^{-5} M increased the stomatal index and its density on abaxial side, showing the opposite response in the adaxial side (Mendoza *et al.*, 2002).

Macro and microelements. As evident from Table IV. SA treatment increased the content of N, P, K, Mn, Fe, Zn, Na and Cu, in *O. basilicum* and *M. hortensis*. The maximum mean values of all macro and micronutrients content of both species were obtained as a result of SA at 10^{-4} M and 10^{-5} M. Sarangthem and Singh (2003) found that the level of N, proteins and nitrate reductase activity were increased in *Phaseolus vulgaris* by foliar application of SA at 0.1%. In this study, increased nutrient content seems to be involved in stress-tolerance mechanism and play an important role to enhance the activity of enzymes responsible for drought resistance (Wu *et al.*, 1999; Cherki *et al.*, 2002). SA applied at 10^{-3} M showed a variable pattern of various nutrients (Table V). It is slightly increased the uptake of all macro

Ocimum basilicum	Carbohydrates	Crude	Free amino Free proline			Photosynthetic pigments (mg/g dry wt)			
		proteins	acids						
Treatments	%	%	Mg/g dry wt	mg/g dry wt	Ch.a	Ch.b	Ch.a+b	Carotenoids	Total pigments
Control	19.25 <u>+</u> 0.52	19.00 <u>+</u> 0.25	4.8 <u>+</u> 0.03	6.84 <u>+</u> 0.10	2.90 <u>+</u> 0.07	2.01 <u>+</u> 0.02	4.91 <u>+</u> 0.07	1.21 <u>+</u> 0.07	6.12 <u>+</u> 0.14
SA (10 ⁻⁵ M)	21.69 <u>+</u> 0.26	25.15 <u>+</u> 0.16	5.48 <u>+</u> 0.35	14.03 <u>+</u> 0.36	3.32 <u>+</u> 0.05	2.12 <u>+</u> 0.01	5.43 <u>+</u> 0.04	1.46 <u>+ 0.05</u>	6.89 <u>+</u> 0.06
SA (10 ⁻⁴ M)	22.98 <u>+</u> 0.28	27.49 <u>+</u> 0.26	9.30 <u>+</u> 0.11	19.37 <u>+</u> 0.45	3.74 <u>+</u> 0.03	2.32 <u>+</u> 0.05	6.05 <u>+</u> 0.04	1.74 <u>+</u> 0.06	7.79 <u>+</u> 0.06
SA (10 ⁻³ M)	18.33 <u>+</u> 0.44	22.02 <u>+</u> 0.16	6.06 <u>+</u> 0.47	18.99 <u>+</u> 0.43	3.05 <u>+</u> 0.11	1.35 <u>+</u> 0.10	4.39 <u>+</u> 0.20	1.45 <u>+</u> 0.07	5.85 <u>+</u> 0.14
L.S.D. at 5%	090	0.45	0.60	0.57	0.10	0.13	0.23	0.15	0.23
Origanium hortensis									
Control	20.19 <u>+</u> 0.52	10.00 <u>+</u> 0.13	2.09 <u>+</u> 0.10	3.75 <u>+</u> 0.08	1.96 <u>+</u> 0.08	1.04 <u>+</u> 0.10	2.99 <u>+</u> 0.07	2.45 <u>+</u> 0.05	5.44 <u>+</u> 0.09
SA (10 ⁻⁵ M)	22.39 <u>+</u> 0.21	11.25 <u>+</u> 0.19	2.35 <u>+</u> 0.07	5.19 <u>+</u> 0.02	2.66 <u>+</u> 0.04	1.29 <u>+</u> 0.07	3.95 <u>+</u> 0.02	2.62 <u>+</u> 0.02	6.57 <u>+</u> 0.03
SA (10 ⁻⁴ M)	24.94 <u>+</u> 0.13	12.50 0.50	2.65 <u>+</u> 0.03	5.41 <u>+</u> 0.03	3.18 <u>+</u> 0.06	1.55 <u>+</u> 0.06	4.73 <u>+</u> 0.08	2.84 <u>+</u> 0.03	7.57 <u>+</u> 0.07
SA (10 ⁻³ M)	23.73 <u>+</u> 0.39	10.00 <u>+</u> 013	2.45 <u>+</u> 0.01	3.76 <u>+</u> 0.03	1.62 <u>+</u> 0.03	0.72 <u>+</u> 0.03	2.34 <u>+</u> 0.04	2.22 <u>+</u> 0.02	4.56 <u>+</u> 0.04
L.S.D. at 5%	0.65	0.62	0.15	0.11	0.13	0.16	0.13	0.07	0.13

Table III. Effect of foliar spray with salicylic acid on chemical constituents of *Ocimum basilicum* L. and *Origanium hortensis* plants <u>+</u> standard deviation

Table IV. Effect of foliar spray with salicylic acid on macro and micronutrients content of *Ocimum basilicum* L. and *Origanium hortensis* plants

Ocimum basilicum	Macronutrients content (%)				Mic			
Treatments	Ν	Р	K	Mn	Fe	Zn	Na	Cu
Control	3.04	0.29	3.50	122.50	888.00	78.00	16.79	31.50
Salicylic acid 10-5 M	4.02	0.41	3.61	148.50	1063.00	96.00	18.86	32.00
Salicylic acid 10 ⁻⁴ M	4.40	0.45	3.68	154.50	1098.00	235.00	20.70	33.00
Salicylic acid 10-3 M	3.52	0.18	3.13	130.00	904.50	93.00	14.03	30.00
Origanium hortensis								
Control	1.60	0.34	2.08	84.00	1399.00	61.00	14.95	20.50
Salicylic acid 10 ⁻⁵ M	1.80	0.49	2.32	85.00	1735.00	67.00	18.86	21.50
Salicylic acid 10-4 M	2.00	0.59	2.39	103.00	1770.00	73.00	35.88	47.50
Salicylic acid 10 ⁻³ M	1.60	0.31	2.22	91.00	1576.00	49.50	13.34	17.50

Table V. Effect of foliar spray with salicylic acid on macro and micronutrients uptake of *Ocimum basilicum* L. and *Origanium hortensis* plants

Ocimum basilicum	Macron	utrients uptake	t (mg plant ⁻¹)		Micronutrients uptaket (mg plant ⁻¹)			
Treatments	Ν	Р	K	Mn	Fe	Zn	Na	Cu
Control	154.43	14.53	177.85	62.23	451.10	39.62	8.53	16.26
Salicylic acid 10 ⁻⁵ M	286.63	29.45	257.18	105.88	757.92	68.45	13.45	22.82
Salicylic acid 10 ⁻⁴ M	479.60	48.61	400.79	168.41	1196.82	256.15	22.56	35.97
Salicylic acid 10 ⁻³ M	234.08	12.10	208.28	86.45	601.49	61.85	9.33	19.95
Origanium hortensis								
Control	118.56	24.82	153.83	62.24	1036.66	45.20	11.08	15.19
Salicylic acid 10 ⁻⁵ M	148.86	40.69	192.11	70.30	1434.85	55.41	15.60	17.78
Salicylic acid 10 ⁻⁴ M	198.80	58.15	237.86	102.38	1759.38	72.56	35.66	47.22
Salicylic acid 10-3 M	126.88	24.82	175.81	91.00	1576.00	49.50	13.34	17.50

and micro-element of herbs, except P (in case of basil) and Zn, Na and Cu content (in case of marjoam). Similarly, in tomato, significant increase in the uptake of elements was obtained with 10^{-4} M and 10^{-5} M salicylaldehyde (Kord & Hathout, 1992). Application of SA might improve physiological performance in terms of production of photosynthates, total oil and dry matter yield, which can be related to increased nutrient uptake by SA-treated plants (Abad-Farooq & Misra, 1983; Cheol *et al.*, 2001).

Polyamines. In control basil plants spermidine (Spd) was dominant followed by putrescine (Put), while spermine (Spm) was undetectable (Table VI & Fig. 5). Contrarily, control marjoram not only showed greater total polyamines (TPAs) but also differential pattern of individual polyamines, spermine being the main, followed by spermidine and putrescine. In both species, endogenous putrescine (Put), spermidine (Spd) as well as total polyamines (TPAs) contents were gradually increased with the increase of SA concentration up to 10^{-4} M relative to their controls. In an earlier study, the application of polyamines and SA at 400 mg L⁻¹ to *Citrus reticulate* increased endogenous PA and SA concentrations, leading to an improvement in fruit quality and prolonged storage life. Moreover, SA, Spm and Spd treatments were better than a Put (Zheng & Zhang, 2004). Polyamines are considered anti-senescence in nature and regulate the action of the prosenescence ethylene (Cassol & Mattoo, 2002). Spermine was the main polyamine in marjoram under control condition, which was un-detectable in basil. Furthermore, at 10^{-5} M of SA, a marked decrease in spermine (reduced to

Fig. 4. Effect of foliar spray with salicylic acid on chemical constituents of *Ocimum basilicum* L. and *Origanium hortensis* plants. Vertical bars represent \pm standard deviation.



Fig. 5. HPLC analysis for endogenous concentrations of polyamines of *Ocimum basilicum* L and *Origanium hortensis* plants sprayed with salicylic acid.



approximately 63.9% of the control) was observed in marjoram plants, whereas it was un-detectable in basil. Again, SA at 10^{-4} M displayed the maximum values of spermine in both species, relative to controls. Polyamines accumulation under conditions of water-deficit stress in drought-tolerant alfalfa may be a general strategy for improvement. Consequently, SA treatment may activate polyamine oxidase and improve abiotic stress tolerances in sweet basil and marjoram plants.

Table VI. HPLC analysis for endogenous concentrations of polyamines of *Ocimum basilicum* L. and *Origanium hortensis* plants sprayed with salicylic acid

Ocimum basilicum	Polyamines levels (mg 100 ⁻¹ g F.Wt)								
Treatments	Put	Spm	Spm	Total					
Control	0.332	0.625	N.S	0.957					
SA (10 ⁻⁵ M)	1.077	2.923	N.S	4.000					
SA (10 ⁻⁴ M)	2.071	4.951	0.219	7.241					
SA (10 ⁻³ M)	0.744	2.556	0.204	3.504					
Origanium hortensis									
Control	0.142	0.493	0.535	1.170					
SA (10 ⁻⁵ M)	0.746	5.375	0.193	6.314					
$SA(10^{-4}M)$	0.278	5.944	0.585	6.807					
SA (10 ⁻³ M)	0.772	3.854	0.489	5.115					

CONCLUSION

SA applied on basil and marjoram stimulated the growth and oil yield by enhancing photosynthesis and nutrient uptake. Basil may be a new source of antioxidant phenolics in the diet due to the greater production of eugenol by SA. All SA treatments enhanced total free amino acid, free proline, Put, Spd as well as total polyamines TPAs, whilst differently affecting the level of spermine. However, further investigations are required to elucidate the possible role of SA on plant growth regulating activity.

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