

ORIGINAL ARTICLE

Foliar Application of Salicylic Acid and Calcium on Yield, Yield Component and Chemical properties of Strawberry

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ABSTRACT

In order to study effect of salicylic acid and calcium foliar application on growth, yield and yield components of strawberry plants as a factorial in completely randomized experimental design with four replications. This factors included of salicylic acid in 3 levels (0.25, 0.5 and 0.75 mM) and calcium in 2 levels (2.5 and 5 mM) spray on strawberry. Results showed that salicylic acid (0.25 mM) and calcium chloride (2.5 mM) spray either alone or in combination (0.25 mM SA+2.5 mM Ca) affected on vegetative and reproductive growth, significantly. Mean comparisons indicated yield, and quality of strawberry plants was improved in low salicylic acid and calcium chloride concentration. In Finally, salicylic acid and calcium chloride application can be helpful for yield improvement and prevent of decreasing yield.

Keywords: Strawberry, Salicylic acid, Calcium chloride, Yield, Quality

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INTRODUCTION

Foliar feeding of vegetable plants can effectively supplement soil fertilization. It has been found that elements foliar application is in a same level and even more influential compared to soil application. It was suggested that foliar feeding could be applied successfully to compensate shortage of those elements [1]. Salicylic acid, a naturally occurring plant hormone acting as an important signaling molecule adds to tolerance against abiotic stresses. It plays a vital role in plant growth, ion uptake and transport. Salicylic acid can also play a significant role in plant water relations, photosynthesis and growth in plants [2]. In cucumber and tomato, the fruit yield enhanced significantly when the plants were sprayed with lower concentrations of salicylic acid [3]. Khodary [4] and El-Tayeb [5] found that SA treatment increased the chlorophyll, dry weights and carotenoid contents in maize and barley plants. Calcium (Ca^{2+}) is a universal second messenger and it has long been considered as the second messenger in many signaling cascades; including defense signaling [6]. The preharvest nutritional status of fruit, especially with respect to calcium, is an important factor affecting potential storage life [7]. Direct application of calcium to the fruit is the most effective method for increasing fruit calcium content [8]. Preharvest and postharvest treatments with calcium salts have been effective in control of several physiological disorders, reducing the incidence of fungal pathogens and maintaining fruit firmness [9]. Foliar applications of calcium chloride have been reported to delay ripening and retard fungal growth on strawberries [10]. SA and Ca treatments have the potential for commercial control of quality properties and increase shelf life of harvested fruit. Therefore, the aim of this research is to determine the effects of pre-harvest application of SA and Ca in strawberry cultivar 'Selva', specially the change in quality properties.

MATERIALS AND METHODS

Plant material and growth conditions

The experiment was conducted during 2013 and 2014 on strawberry plants at a hydroponic greenhouse located in Iran. Strawberry plants (*Fragaria × ananassa* Duch. cv. Selva) were grown under natural light conditions. The temperature conditions were $25 \pm 2^\circ\text{C}$ and $16 \pm 4^\circ\text{C}$, during days and nights respectively; with mean relative humidity of $70 \pm 5\%$. Runner plants were rooted in plastic pots filled with leca and perlite. They were irrigated three times a day. Rooted plants were transplanted into 3 liter pots. The pots were filled with leca, perlite and peatmoss (1: 3: 1 v/v/v). The plants were fertilized with the following complete nutrient solution containing: N, P_2O_5 , K_2O , Ca, Mg, S, Fe, Mn, Mo and Cu.

Treatments

SA at the rate of 0.25, 0.5 and 0.75 mM, Ca at the rate of 2.5 and 5 mM as CaCl₂ were applied as foliar. The surface of pot was covered with aluminium foil to prevent the adding of SA and CaCl₂ solution into root medium. Untreated plants were left as a control and sprayed with distilled water.

Measurements

Total leaf areas of each plant were recorded with portable leaf area meter. Petiole length was measured using a ruler. The change in petiole diameter was determined for each plant using vernier caliper. To attain a constant weight for biomass estimation (dry weight), plant components were oven dried at 70°C. Total fruit weight of each plant was separately measured and considered as yield. To evaluate fruit quality, total soluble solid (TSS), titratable acidity (TA) and ascorbic acid (vitamin C) were measured. TSS of fruits was determined using a refractometer. Total titratable acid and vitamin C was measured by NaOH (0.1 M) titration and indophenol's method according to Horvitz *et al.*, [11]. The experiment was a factorial in completely randomized design with 6 replications. Data were analyzed by SPSS 16 software and comparing averages was done by Duncan's test and a probability value of %5.

RESULTS AND DISCUSSIONS

The results indicated that 0.25 mM SA and 2.5 mM Ca both caused a significant increase in vegetative and reproductive growth compared to other levels ($p \leq 0.05$). The combination between SA and Ca (0.25 mM SA + 2.5 mM Ca) on vegetative and reproductive growth was significant, as well (Table 1). The dry weight increased to its maximum (15.13 g) with 0.25 mM SA + 2.5 mM Ca application (Table 1). The maximum number of runners (6.12), leaf area (30.12 cm), length of roots number (27.12), number of flowers (14.51), length of flowering period (35.12), weight of primary fruit (18.9 g), weight of secondary fruit (17 g), number of achenes of primary fruit (215.12) and number of achenes of secondary fruit (195) was recorded with 0.25 mM SA + 2.5 mM Ca (Table 1-2). Salicylic Acid (SA) is a growth regulator which participates in the regulation of physiological processes in plants. It stimulates flowering in a range of plants, increases flower life, controls ion uptake by roots and stomatal conductivity [12]. Previous studies have demonstrated that a wide range of responses might appear after exogenous SA application as follows: height plant increases, fruit weight and fruits per plant [13-14]. Application of SA also significantly increased dry weights of root and top part of barley and soybeans [15]. The mechanism of salicylic acid was reported by Oata [16] and Pieterse and Muller [17] who concluded that salicylic acid induced flowering by acting as a chelating agent. This view was supported by Raskin *et al.*, [18] who confirmed that salicylic acid functioned as endogenous growth regulators of flowering and florigenic effects. Deficiency of Ca decreases plant height by decreasing mitotic activity in the terminal meristem [19]. Thus, the application of calcium increases plant height [20]. Application of Calcium carbide also stimulates root growth and early onset of flowering in agronomic and vegetable crops [21]. The highest percentage of TSS and TA and ascorbic acid content was attained in fruits treated with 0.25 mM SA + 2.5 mM Ca and the lowest was achieved in control (Table 4). Also highest of pH (2.21 %) content was attained in fruits treated with 0.25 mM of foliar SA concentration. Table 3, indicates the effect of SA and Ca on total phenolics (TP), flavonoids (TF), and non-flavonoids (TNF) and lipid peroxidation (MDA). 0.25 mM SA + 2.5 mM Ca of increased total phenolics (TP), flavonoids (TF), and non-flavonoids (TNF) of strawberry. In general, application of SA and Ca produced significantly decreased lipid peroxidation (MDA) of fruits compared to control. It was reported that the foliar application of salicylic acid on soybean also enhanced the flowering and pod formation [22]. Sayyari *et al.*, [23] has shown that the amount of acidity and TSS was influenced by SA treatment in pomegranate. Chandra *et al.*, [24] reported that application of salicylic acid increased total soluble sugar and soluble protein of cowpea plants. Vitamin C, pH, TSS and titratable acidity (TA) of fruits treated with higher 5-SSA concentrations was higher than those of control fruits. It has been suggested that TA decreases in fruits in result of breakup of acids to sugars during respiration [25]. Han and Li [26] have also reported that apple fruits treated with SA had increased TA content at the end of storage. Our results showed that SA had significant effect on maintaining higher content of vitamin C in peach fruits. Kazemi *et al.*, [27] have also reported that fruits treated with SA were observed with maximum vitamin C content. Ghasemnezhad *et al.*, [28] reported that the decrease of total phenolic levels might be due to breakdown of cell structure in order to senescence phenomena during the storage period. The effect of salicylic acid treatments on maintain of total phenolics content plausibly may be attributed to delay in senescence process. Our results are in agreement with those of kazemi *et al.*, [27], who reported treatment with salicylic acid significantly reduced the membrane permeability and MDA content. In conclusion, foliar application of SA and Ca treatments were generally effective on vegetative growth, photosynthetic pigments, minerals, yield and quality of strawberry fruit.

Table 1) Effect of pre-harvest application of SA and Ca on dry weight, number of runners, leaf area, number of flowers and length of the roots of strawberry

| Treatment | | dry weight(g) | Number of runners | Leaf area (cm) | Length of roots (cm) | Number of flowers |
|-------------|----------|---------------|-------------------|----------------|----------------------|-------------------|
| Control | 0 | 6c | 3c | 20.67c | 16.03c | 9.78c |
| SA | 0.25 | 13ab | 5.98a | 28ab | 26.67a | 13.6ab |
| | 0.5 | 10.11b | 4.43b | 28b | 23.34ab | 12.07b |
| | 0.75 | 10b | 4.4b | 27.45b | 20.12b | 12.01b |
| Ca | 2.5 | 13.12ab | 6a | 28.67ab | 26.78a | 13.67qb |
| | 5 | 9.97b | 4.47b | 27.78b | 19.89b | 11.87b |
| Combination | 0.25*2.5 | 15.13a | 6.12a | 30.12a | 27.12a | 14.51a |
| | 0.25*5 | 10b | 4.49b | 27b | 20.12b | 12.11b |
| | 0.5*2.5 | 9.78b | 4.35b | 26.45b | 18.12bc | 12b |
| | 0.5*5 | 9bc | 4.3b | 23.56bc | 18.43bc | 10.12bc |
| | 0.75*2.5 | 9bc | 3.45bc | 23.34bc | 18.08bc | 10.08bc |
| | 0.75*5 | 8.89bc | 3.39bc | 23.12bc | 18bc | 10bc |

Means followed by same letter are not significantly different at 5% probability using Duncan's test.

Table 2) Effect of pre-harvest application of SA and Ca on Length of flowering period, weight of primary fruit, weight of secondary fruit, number of achenes of primary fruit and Number of achenes of secondary fruit

| Treatment | | Length of flowering period (day) | Weight of primary fruit (g) | Weight of secondary fruit (g) | Number of achenes of primary fruit | Number of achenes of secondary fruit |
|-------------|----------|----------------------------------|-----------------------------|-------------------------------|------------------------------------|--------------------------------------|
| Control | 0 | 12.12c | 10c | 7.88c | 110c | 98c |
| SA | 0.25 | 19.12ab | 17.98ab | 16.01ab | 200ab | 191.12ab |
| | 0.5 | 15.14b | 15.11b | 12bc | 189.12b | 160b |
| | 0.75 | 15b | 15b | 12.7b | 167.12bc | 145.21bc |
| Ca | 2.5 | 18.98b | 18ab | 16.21ab | 199.23ab | 187.12ab |
| | 5 | 16b | 15.65b | 13.13b | 187.12b | 167.12b |
| Combination | 0.25*2.5 | 21.12a | 18.97a | 17a | 215.12a | 195a |
| | 0.25*5 | 16.09b | 16.01b | 12.12b | 170.12bc | 167.12b |
| | 0.5*2.5 | 16b | 15.34b | 13b | 168.12bc | 160.09b |
| | 0.5*5 | 15.34bc | 15b | 12.34b | 165.32bc | 150.09bc |
| | 0.75*2.5 | 15bc | 13.14bc | 11.12bc | 163.09bc | 148.12bc |
| | 0.75*5 | 15.01bc | 13bc | 11bc | 161.23bc | 146bc |

Means followed by same letter are not significantly different at 5% probability using Duncan's test.

Table 3) Effect of pre-harvest application of SA and Ca on TP, TF, TNF and MDA

| Treatment | | TP w(gallic acid)/(mg/kg) | TF w(gallic acid)/(mg/kg) | TNF w(gallic acid)/(mg/kg) | MDA (nmol gfw-1) |
|-------------|----------|---------------------------|---------------------------|----------------------------|------------------|
| Control | | 67.41c | 20.11c | 21.12c | 28.12a |
| SA | 0.25 | 119.23a | 40.12ab | 45.32a | 11c |
| | 0.5 | 151.23b | 33.11b | 30.12b | 20bc |
| | 0.75 | 149.21b | 32.32b | 29.34b | 20.12bc |
| Ca | 2.5 | 119a | 41.14ab | 44.54a | 12.12c |
| | 5 | 142.12b | 33b | 31.12b | 19.12bc |
| Combination | 0.25*2.5 | 121.12a | 44.12a | 46.12a | 10.11c |
| | 0.25*5 | 150.12b | 40.12ab | 40.12ab | 12.12c |
| | 0.5*2.5 | 147.21b | 31.12b | 32.12b | 19.67bc |
| | 0.5*5 | 149.21b | 28.23bc | 30.12b | 19.56bc |
| | 0.75*2.5 | 145.12b | 27.56bc | 30b | 19.34bc |
| | 0.75*5 | 144.54b | 26.34bc | 28.12bc | 19bc |

Means followed by same letter are not significantly different at 5% probability using Duncan's test.

Table 4) Effect of pre-harvest application of SA and Ca on pH, TSS, TA and Vit.C

| Treatment | | pH | TSS | TA g(citric acid)/(g/L) | Vitamin C (mg/100 g) |
|-------------|----------|-------|--------|-------------------------|----------------------|
| Control | | 1c | 5c | 1.43c | 20.12c |
| SA | 0.25 | 2.21a | 7.8ab | 3a | 30.12ab |
| | 0.5 | 1.67b | 6.45b | 2.45b | 25.12b |
| | 0.75 | 1.6b | 6.35b | 2.4b | 25b |
| Ca | 2.5 | 1.98a | 7.67ab | 2.98a | 31.12ab |
| | 5 | 1.6b | 6.3b | 2.32b | 26.12b |
| | 0.25*2.5 | 2.11a | 8a | 3.12a | 33.11a |
| Combination | 0.25*5 | 1.6b | 6.45b | 2.32b | 25b |
| | 0.5*2.5 | 1.56b | 6.4b | 2.3b | 24.98b |
| | 0.5*5 | 1.52b | 5.31bc | 1.67bc | 22.11bc |
| | 0.75*2.5 | 1.5b | 5.3bc | 1.6bc | 22bc |
| | 0.75*5 | 1.5b | 3.13bc | 1.54bc | 21.12bc |

Means followed by same letter are not significantly different at 5% probability using Duncan's test.

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