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Effect of foliar application of varying doses of salicylic acid at different growth stages on growth, quality and nutrient uptake efficiency of French bean (*Phaseolus vulgaris* L.)

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Abstract

Background: The ubiquitous presence of salicylic (SA) acid in plant systems has been confirmed by many scientists. Salicylic acid is widely distributed throughout the plant and is proposed to affect a number of phyto-physiological activities. As an endogenous plant growth factor or a hormone SA has number of functions including florigenic and thermogenic role, part in plant-pathogen interaction, signal transduction, etc.

Method: The present investigation was conducted at the Experimental Farm of the Division of Vegetable Science, SKUAST-K, during *Kharif* 2020 to find out the effect of foliar spray of varying doses of salicylic acid (SA) applied at different plant growth stages on growth and quality attributes of French bean var. Shalimar French Bean-1. The experiment was laid out in split plot design with twelve treatment combinations and three replications.

Result: The investigation revealed that the treatment combination T_1D_2 (250ppm of SA applied at plant emergence stage) recorded minimum values for days taken to 50 per cent flowering and days taken to pod formation. Moreover, the same treatment combination T_1D_2 also recorded maximum values for plant height, number of primary branches plant⁻¹, number of leaves plant⁻¹ *et al.*, leaf area and shoot dry weight plant⁻¹. It was further revealed that in the treatment combination T_1D_2 maximum values of vitamin C content, TSS, protein content, leaf chlorophyll content and pod dry matter content were also recorded, while pod anthocyanin content was maximum in the treatment combination T_2D_2 (250ppm of SA applied at vegetative stage). The treatment combination T_1D_2 also significantly enhanced nutrient uptake efficiency of the plants and resulted in lowest values for the available nitrogen, phosphorus and potassium, soil EC and maximum values for organic carbon as compared absolute control. than absolute control D_0 .

Keywords: French bean, purple pod, salicylic acid, growth stage, foliar application, quality, growth, nutrient uptake

Introduction

French bean (Phaseolus vulgaris L.) is one of the most important food legumes cultivated extensively throughout the world. It can be raised as a vegetable crop or as a pulse. Common bean is cultivated prodigiously as a pulse crop but in some pockets, it is especially cultivated for its delicious tender fresh pods as well as dry pods. Beans are one of the delights of vegetarians for their wholesome nutritional properties but in the valley region the cultivation of beans for fresh pods is restricted to a limited area with quite less productivity (1.45 t ha⁻¹) as compared to rest of the country (2.83 t ha⁻¹) (Anonymous, 2018) ^[1]. Being a nutritionally important legume and a short duration crop French bean can be taken up by the famers in the valley for vegetable purpose to obtain good economic returns in less period of time. Besides following the recommended package of practices for the cultivation of beans, foliar application of different doses of salicylic acid at various plant growth stages will augment the growth development along with supplementing to the pod quality. Salicylic acid promotes plant growth and development along with increasing the efficiency of photosynthesis and flowering (Hayat et al., 2007)^[10]. Other physiological roles of salicylic acid include increasing uptake of mineral nutrients from the soil, induction of tolerance against abiotic as well as biotic stresses in the plants, enhancing nitrogen metabolism thereby increasing the protein content and

interfering in ethylene production as a result of which delayed senescence is observed in the plants treated with salicylic acid. All these factors together contribute towards a rampant increase in the yield trends of the plants treated in cause of the application of salicylic acid. Manipulating the levels of salicylic acid in plant through exogenous application may have positive impacts on the productivity as well as the nutritional content of the bean pods.

Materials and Methods

The field of experiment was situated at the main SKUAST-K campus in Shalimar, Srinagar, which is located on the foothills of the Mahadev hill, 15 kilometres from Srinagar city, at an elevation of 1615 metres above mean sea level and is 34 degrees north latitude and 74 degrees east longitude. The experiment was laid out with 12 treatment combinations and three replications in split plot design as shown in Table A. The recommended dose of fertilizers FYM- 25 t ha-1 and NPK (30:60:60 kg ha⁻¹) was added to the experimental field through DAP, urea and MOP at the time of land preparation. Basal application of entire FYM, P, K and half N is done. Rest of the N is applied after 30 DAS. The flat seed beds were prepared by thoroughly working the soil into fine tilth. The seeds of French bean variety Shalimar French Bean-1 were used for sowing. The seeds were sown on flat beds with a spacing of 30 cm \times 10 cm between and within rows thus accommodating approx. 100 seeds per plot (3 m²). Seeds were soaked in water for 24 hrs prior to sowing so as to soften the hard seed coat and hasten germination. After sowing, the field was immediately irrigated. 10 healthy plants were selected at random from each replication to study observations on different growth and yield traits and tagged for recording the observations. Mean values were worked out for all the characters.

The height of ten tagged plants was measured in centimetres from the ground level to the tip of the plant at the time of last picking and the average was worked out.

At the end of flowering, branches emerging from the main stem, total number of compound leaves present on the whole plant were counted and leaf area of the fully developed 3 leaves was estimated using leaf area meter from ten selected plants and the average was worked out.

At the 50 per cent flowering stage, ten healthy plants were chosen for the calculation of dry shoot weight. The plants were uprooted, the underground root portion was cut and the clean shoot part was placed for 72 hours in an oven at 65°C. The weight was estimated after the required time and then the mean value was expressed in grams.

The days to the 50 per cent flowering appearance set on each tagged plant was recorded from the date of sowing of seeds and the average was worked out.

The days to the full pod set development on each tagged plant was recorded from the date of sowing of seeds and the average was worked out.

The selected fruits were crushed and juice was passed through a double layer of mesh cloth, collected in a tube, shaken well for about 2 minutes and then TSS was determined with the help of hand refractometer and presented as [°]Brix

The total soluble protein was estimated as per the method described by Lowry *et al.* (1951)^[18].

Ascorbic acid content was determined by the method described by Ranganna (1986)^[24].

Chlorophyll estimation was done using acetone method

(Arnon, 1949)^[2].

The pH differential spectrophotometric method described by Lapornik *et al.* (2005) ^[17] based on the principle of the transformation of the anthocyanin to the flavylium cation at the pH of extract decreasing to values between 0.5 and 0.8 was used to measure the anthocyanin content of the samples.

From each plot, 100 g of randomly selected pods (from each tagged plant) was placed in the hot air oven at 60-65 °C for 72 hrs and using the final weight after drying, the pod dry matter percentage was worked out.

The soil samples were converted into soil water suspension of ratio 1: 2.5 and the pH of these suspensions was estimated by a glass electrode digital pH metre.

The Walkley and Black's method (1934) ^[28] was used to determine the soil organic carbon content.

The SOLU-bridge conductivity meter was used to estimate soil electrical conductivity (Jackson, 1973)^[11].

To determine the available phosphorus content of the soil Olsen method (1954) was employed by using NaHCO₃ and estimation was done by using spectrophotometer.

One normal ammonium acetate was used as extractant and the available potassium content was determined by feeding the extract to flame photometer (Jackson, 1973)^[11].

The experimental data was subjected to statistical analysis to test the significance of the findings. Significance of difference in treatment stage, dose and stage-dose interaction was tested at 5% level of significance and CD (critical difference) was worked out.

Results and Discussion

- 1. Growth Parameters: The experimental results depicted that foliar application of different concentrations of SA at different growth stages had a significant influence on the growth of French bean plants. Among all the treatment combinations, the treatment combination T_1D_2 i.e., application of 250ppm of SA at plant emergence stage resulted in significantly superior values for most of the growth attributes as compared to control treatment. From the data in tables 1.1 to 1.7 it is comprehensible that foliar spray of 250ppm of SA significantly increased the plant height, no. of branches per plant, no. of leaves per plant, leaf area and dry shoot weight per plant when applied at plant emergence stage. The intensity of growth processes of plants treated with SA was noticeably higher than in control. This may be attributed to the growth stimulating effect of SA. Application of SA promotes the accumulation of auxin and cytokinin which consequently increases the mitotic index of the apical meristems. The increase in the mitotic activity subsequently leads to enhanced growth (Hayat et al., 2007) [10]. These results were in agreement with the findings of Kumar et al. (1997) in Pisum sativum; Khatun et al. (2016) ^[15] in Glycine max; Yadav et al. (2019)^[30] in Vigna unguiculata and Saha et al. (2020) ^[27] in *Phaseolus vulgaris*. Days taken to 50 per cent flowering and days taken to pod formation *-were also found to be significantly reduced with the application of SA. Application of SA hastened the floral bud emergence by 2-5 days due to its florigenic role. SA promotes the movement of solutes to the growing buds which causes better development of the buds and precocious flowering (Watanabe et al., 1981)^[29].
- **2. Quality Parameters:** Phenolic compounds like SA are acknowledged to have a significant influence on the

quality attributes like total soluble solids (TSS), protein content, ascorbic acid content, pod dry matter and pigments like chlorophyll and anthocyanins. From studying the experimental findings in tables 2.1 to 2.5, it was found that application of 250 ppm of SA at both plant emergence or vegetative growth stage significantly enhanced the bio-chemical attributes as compared to rest of the treatment combinations. The promotive effect of the foliar spray on chlorophyll content may be attributed to increased N uptake from the soil which is a principal component of the chlorophyll structure. Similar results were observed by, Khan et al. (2003) [14] in soybean and Khan et al. (2012) in wheat. Due to increased chlorophyll content; leaf area and no. of leaves per plant, subsequently the photosynthetic efficiency of the plant will be increased and higher amounts photosynthates will be produced which may contribute towards increased TSS and pod dry matter content. The increase in level of proteins is presumably because of better resistance mechanism possessed by the application of SA, since most of the pathogenesis related proteins residing in the leaf intercellular fluids are responsible to provide resistance to the plants against diseases (Chandra et al., 2007)^[3]. These results coincided with the findings of Maity and Bera et al. (2009) ^[19] in Vigna mungo; Javaheri et al. (2012) ^[12] in tomato, Chaudhary (2019)^[5] in pea and Chattoo et al. (2020)^[4] in onion. Application of SA enhanced the ascorbic acid content in the pods. Increase in ascorbic acid content with the application of SA might been reported to be due to indirect activation of ascorbic acid biosynthesis from carbohydrates such as sucrose and glucose, as reported by Pérez-Balibrea et al. (2011) [23] in broccoli sprouts and inhibited ascorbic acid oxidase (AAO) enzyme activity, which is responsible for ascorbic acid oxidation as suggested by Rao et al. (2011) in sweet pepper. Anthocyanin content was markedly increased with the application of 250 ppm SA at vegetative stage. Anthocyanins and ascorbic acid contents are main components of fruit antioxidant activity and SA is an established natural plant signalling molecule that may stimulate the antioxidant activity of fruit by enhancing total phenolics, flavonoids, anthocyanins and ascorbic acid contents. Dokhanieh et al. (2013)^[7] demonstrated that SA might be a potential molecule for activating phenylpropanoid-flavonoids pathways in the cherry fruit which stimulates the accumulation of phenolics, flavonoids and anthocyanins by activating their biosynthetic pathways. Anthocyanins have shown to be potent antioxidant and ROS scavengers (Hassanpour et al., 2011)^[9] and SA in also known to have a significant role in promoting anti-oxidant activity. Razmi et al. (2013) [26] also reported that treatment of soybean with SA enhanced

the total pigment content.

Nutrient Uptake: The efficiency of plants to absorb 3. available nutrients from the soil depends on the physiological health of the plant. The healthier the plant, better will be its root system and higher will be it efficiency to absorb nutrients from the soil. As a result of this higher plant biomass will be produced and inflated yield will be achieved. This would ultimately exhaust the soil reservoir. Application of SA increase the nutrient uptake efficiency of the plant by improving the source and sink relationship, due to which higher amounts of nutrients will be absorbed from the roots and moved to the sink i.e., leaves and pods (Hayat et al., 2007) [10]. Ultimately, a result of this is that there is decline in soil nutrient status. The data in tables 3.1 to 3.6 shows that application of 250 ppm of SA at plant emergence and vegetative growth stage resulted in a significant decrease in the values of soil pH, soil EC, available soil N, P and K. This can be correlated to increased uptake of nutrients from the soil with the foliar spray of SA. Due to better establishment of roots at younger stages with the application of SA the nutrient uptake the plant is increased, consequently the amounts of mineral salts in the soil will be decreased. This might be the reason behind the lower values recorded for available nitrogen, phosphorous and potassium in the soil reservoir. The same treatment combination resulted in a significant increase in the organic carbon content in the soil. The increase in organic carbon content may be attributed to better development of root biomass which increases the organic matter content in the soil. This increase in organic matter might also be responsible for the corresponding decline in soil pH as well as EC. These findings can be supported through the investigations of Gunes *et al.* (2007)^[8] in maize, Khan *et al.* (2010)^[13] in mung bean (Vigna radiata L.), Nazar et al. (2011)^[21] in mung bean, Dawa et al. (2015)^[6] common bean (Bronco cultivar) and Metwaly and El-Shatoury (2017) [20] in potato cv. Spunta.

Tables

Table A: Treatment details

Fac	ctors	Levels
Α.	Treatment Stages	T_1 – Plant Emergence (2-4 leaf
	(Major Factor)	stage) (15 DAS)
		T_2 – Vegetative Stage (30 DAS)
		T_3 – Flowering Stage (45DAS)
В.	Doses of Salicylic	D_0 – Control (No Spray)
	Acid (Minor	D ₁ – 200 ppm
	Factor)	$D_2 - 250 \text{ ppm}$
		D ₃ – 300 ppm

Table 1.1: Effect of salicylic acid application on final plant height (cm)

Stage Dose	Stage Dose			200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence	(T ₁)	45.88		48.92	51.21	47.59	48.40
Vegetative (T	2)		45.87	47.47	48.93	47.72	47.50
Flowering (Ta)		45.59	46.77	47.79	46.41	46.64
Mean		45.78		47.72	49.31	47.24	
C.D. $(p \le 0.05)$							
	Stage	:	1.34				
	Dose	:	0.53				
	$Stage \times Dose$:	0.77				

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Dose	Stage	Control (D ₀)		200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(7	[1]	2.600		3.133	3.733	2.800	3.067
Vegetative (7	[₂)	4	2.533	2.800	2.867	2.733	2.733
Flowering (7	3)	4	2.667	2.600	2.667	2.600	2.633
Mean		4	2.600	2.844	3.089	2.711	
C.D. $(p \le 0.05)$							
	Stag	e	0.389				
	Dose :		0.142				
	Stag × Do		0.207				

 Table 1.2: Effect of salicylic acid application on number of branches

 plant⁻¹

 Table 1.3: Effect of salicylic acid application on number of leaves

 plant⁻¹

Dose	Stage		ntrol D ₀)	200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emer (T ₁)	gence	38.33		42.53	45.6	41.87	42.08
Vegetative	(T_2)	38.2		41.53	43.93	40.07	40.93
Flowering	(T ₃)	3	9.07	40.27	41.67	39.27	40.07
Mean		38.53		41.44	43.73	40.40	
C.D. $(p \le 0.05)$							
	Stage	:	0.66				
	Dose	:	0.78				
	Stage × Dose	:	1.13				

Table 1.4: Effect of salicylic acid application on leaf area (cm²)

S Dose	tage	Control (D ₀)		-	200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emerge (T1)	ence	34.64		/	38.08	39.31	36.63	37.17
Vegetative (T2)	(1)	34.1	3	36.79	37.39	35.00	35.83
Flowering (Γ3)	34.4		7	35.12	35.86	34.14	34.92
Mean		(T)	34.45		36.66	37.52	35.26	
C.D. (p ≤ 0.05)								
	Sta	Stage		1.05				
	Do	Dose		0.64				
	· ·	age × Dose		0.94				

 Table 1.5: Effect of salicylic acid application on shoot dry weight
 (g)

Dose	Stage		ontrol (D ₀)	200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean		
Plant Emergen	$ce(T_1)$	7	.439	9.983	11.843	8.294	9.39		
Vegetative ((T ₂)	²) 7		7.869		9.704	11.132	8.259	9.241
Flowering (T3)	8.037		9.163	9.645	8.095	8.735		
Mean		7.782		9.617	10.874	8.216			
C.D. (p≤0.05)									
	Stage	:	0.056						
	Dose	:	0.064						
	Stage × Dose	:	0.094						

 Table 1.6: Effect of salicylic acid application on days taken to 50 per cent flowering

Dose	stage	Control (D ₀)			200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T	[1]	45.4		5.4	40.93	40.33	42.20	42.22
Vegetative (7	Γ2)		46	5.13	43.47	42.87	43.93	44.10
Flowering (T	3)	46.33		5.33	46.13	46.00	46.27	46.18
Mean			45.96		43.51	43.07	44.13	
C.D. (p≤0.05)								
	Stag	ge	•••	0.70				
	Dose		•••	0.24				
	Stag Dos		:	0.35				

 Table 1.7: Effect of salicylic acid application on days taken to pod formation

Dose	Stage	-	ontrol (D ₀)	200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emerge	ence(T ₁)	4.	53.20	48.53	47.13	50.00	49.72
Vegetative	Vegetative (T ₂)		54.13	49.87	49.07	50.53	50.9
Flowering	Flowering (T ₃)		54.27	52.73	53	53.87	53.47
Mear	1	53.87		50.38	49.73	51.47	
C.D. (p≤0.05)							
	Stage	:	0.67				
	Dose	:	0.21				
	Stage × Dose	:	0.31				

Table 2.1: Effect of salicylic acid application on pod TSS (° Brix)

Dose	Stage	Contr	rol (D ₀)	200 ppm (D1)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emerge	$ence(T_1)$	8.16		8.45	8.79	8.51	8.48
Vegetative	e (T ₂)	8.02		8.40	8.74	8.30	8.37
Flowering	(T ₃)	8.	13	8.36	8.63	8.22	8.34 ±
Mean	l	8.10		8.4	8.72	8.34	
C.D. (p≤0.05)							
	Stage	:	0.22				
	Dose	:	0.12				
	$Stage \times Dose$:	0.17				

Table 2.2: Effect of salicylic acid application on pod protein content (%)

Stage	Control (D ₀)	200 ppm (D1)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T ₁)	1.693	1.873	2.333	1.867	1.942
Vegetative (T ₂)	1.677	1.86	2.197	1.790	1.881

Flowering (T ₃)	Flowering (T ₃) 1.733			2.017	1.757	1.838
Mean	1.	701	1.859	2.182	1.804	
C.D. (<i>p</i> ≤0.05)						
Stag	e :	0.098				
Dos		0.054				
Stag	$e \times Dose$:	0.079				

Table 2.3: Effect of salicylic acid application on pod vitamin C content (mg 100g-1)

Dose	Stage	Cont	rol (D ₀)	200 ppm (D1)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence	$ce(T_1)$	13.51		14.37	15.16	13.71	14.19
Vegetative (T2)	13.36		14.79	14.86	13.69	14.18
Flowering (Γ3)	1.	3.17	13.98	14.53	13.75	13.86
Mean		1.	3.35	14.38	14.85	13.72	
C.D. (p ≤ 0.05)							
	Stage	••	0.62				
	Dose	••	0.34				
	Stage \times Dose	••	0.49				

Table 2.4: Effect of salicylic acid application on leaf chlorophyll content (mg g⁻¹)

Dose	Stage	Cor	ntrol (D ₀)	200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T ₁)		0.993		1.210	1.590	1.140	1.233
Vegetative (T ₂)			0.980	1.220	1.483	1.063	1.187
Flowering (T ₃)			0.913	1.153	1.323	1.007	1.099
Mean		0.962		1.194	1.466	1.070	
C.D. (p ≤ 0.05)							
	Stage	:	0.088				
	Dose	:	0.066				
	Stage \times Dose	:	0.096				

Table 2.5: Effect of salicylic acid application on pod anthocyanin content ($\mu g g^{-1}$)

Dose	Stage	Co	ontrol (D ₀)	200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T ₁)		13.890	15.563	16.303	14.773	15.133
Vegetative (T ₂)			13.480	15.867	17.307	14.683	15.334
Flowering (T ₃)			13.337	15.08	15.960	14.393	14.693
Mean		13.569		15.503	16.523	14.617	
C.D. (p ≤ 0.05)							
	Stage	:	0.029				
	Dose	:	0.023				
	Stage \times Dose	:	0.034				

 Table 2.6: Effect of salicylic acid application on pod dry matter content (%)

Dose	Stage	Con	trol (D ₀)	200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T)		9.860	11.580	13.157	10.167	11.191
Vegetative (T ₂)			9.797	10.983	13.053	10.087	10.980
Flowering (T ₃)		9.637		10.687	12.963	9.900	10.797
Mean		9.764		11.083	13.058	10.051	
C.D. (p ≤ 0.05)							
	Stage	:	0.100				
	Dose	:	0.026				
	Stage \times Dose	e :	0.038				

Table 3.1: Effect of salicylic acid application on soil pH after crop harvest

Dose	Contr	ol (D	(o)	200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T1)	7.243	(2.87	1)	7.127 (2.851)	7.103 (2.847)	7.140 (2.853)	7.153 (2.855)
Vegetative (T ₂)	7.217	(2.86	6)	7.113 (2.848)	7.113 (2.848)	7.197 (2.863)	7.160 (2.857)
Flowering (T ₃)	7.257	(2.87	3)	7.227 (2.868)	7.220 (2.867)	7.237 (2.87)	7.235 (2.87)
Mean	7.239	(2.87	7)	7.156 (2.856)	7.146 (2.854)	7.191 (2.862)	
C.D. (<i>p</i> ≤0.05)							
	Stage	:	0.037				
	Dose	:	0.041				
	Stage \times Dose	:	0.060				

Data within parenthesis are square root transformed values. Initial soil pH = 7.27

Table 3.2: Effect of salicylic acid application on soil organic carbon (%) after crop harvest

Dose	Control (D ₀)	200 ppm (D1)	250 ppm (D ₂)	300 ppm (D 3)	Mean
Plant Emergence(T ₁)	0.879 (0.938)	0.899 (0.948)	0.938 (0.969)	0.884 (0.940)	0.900 (0.949)
Vegetative (T ₂)	0.845 (0.919)	0.879 (0.938)	0.922 (0.960)	0.881 (0.939)	0.882 (0.939)
Flowering (T ₃)	0.819 (0.905)	0.87 (0.933)	0.919 (0.959)	0.859 (0.927)	0.867 (0.931)
Mean	0.848 (0.921)	0.883 (0.940)	0.927 (0.963)	0.875 (0.935)	
C.D. (p ≤ 0.05)					
	Stage	:	0.021		
	Dose	:	0.008		
	Stage \times Dose	:	0.011		

Data within parenthesis are square root transformed values.

Initial soil OC = 0.910%

Table 3.3: Effect of salicylic acid application on EC (dS m⁻¹) after crop harvest

Dose	Control (D ₀)			200 p	opm (D1)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T ₁)	0.226	(0.475)		0.193	3 (0.439)	0.190 (0.436)	0.197 (0.444)	0.202 (0.449)
Vegetative (T ₂)	0.224	(0.473)		0.197	7 (0.444)	0.190 (0.436)	0.212 (0.46)	0.206 (0.454)
Flowering (T ₃)	0.221	0.221 (0.47)		0.206	5 (0.454)	0.192 (0.438)	0.217 (0.466)	0.209 (0.457)
Mean	0.224	(0.473)		0.199	9 (0.446)	0.191 (0.437)	0.209 (0.457)	
C.D. (p ≤ 0.05)								
	Stage	:	0.	019				
	Dose	:	: 0.					
	Stage \times Dose	:	: 0.					

Data within parenthesis are square root transformed values. Initial soil $\vec{EC} = 0.239 \text{ dS m}^{-1}$

Table 3.4: Effect of salicylic acid application on available N (kg ha-1) in soil after crop harvest

Stage	Control (D ₀)			200 ppm (D ₁)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T ₁)	274.77 (16.	58)	270.80 (16.46)	268.17 (16.38)	271.13 (16.47)	271.22 (16.47)
Vegetative (T ₂)	279.87 (16.	73)	271.54 (16.48)	271.72 (16.48)	272.77 (16.52)	273.98 (16.55)
Flowering (T ₃)	277.69 (277.69 (16.66)		274.26 (16.56)	273.07 (16.52)	275.99 (16.61)	275.25 (16.59)
Mean	277.45 (16.	56)	272.2 (16.5)	270.99 (16.46)	273.3 (16.53)	
C.D. (p ≤ 0.05)							
	Stage	:	2.74				
	Dose	:	1.91				
	Stage \times Dose	:	2.78				

Data within parenthesis are square root transformed values.

Initial soil available N = 281.96 kg ha⁻¹

Table 3.5: Effect of salicylic acid application on available P (kg ha⁻¹) in soil after crop harvest

Dose	Control (D ₀)			200 ppm (D1)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T ₁)	21.79 (4.	67)		19.91 (4.46)	19.50 (4.42)	20.68 (4.55)	20.47 (4.52)
Vegetative (T ₂)	23.08 (4.	80)		21.23 (4.61)	20.03 (4.48)	22.04 (4.69)	21.59 (4.65)
Flowering (T ₃)	22.66 (4.	22.66 (4.76)		22.11 (4.70)	21.52 (4.64)	22.58 (4.75)	22.22 (4.71)
Mean	22.51 (4.	74)		21.08 (4.59)	20.35 (4.51)	21.77 (4.67)	
C.D. (p ≤ 0.05)							
	Stage	•••	1.60				
	Dose	:	0.61				
	Stage \times Dose	:	0.89				

Data within parenthesis are square root transformed values.

Initial soil available $P = 23.89 \text{ kg ha}^{-1}$

Table 3.6: Effect of salicylic acid at different growth stages on available K (kg ha⁻¹) in soil after crop harvest

Dose	Control (D ₀)	200 ppm (D1)	250 ppm (D ₂)	300 ppm (D ₃)	Mean
Plant Emergence(T ₁)	190.69 (13.81)	186.58 (13.66)	185.49 (13.62)	189.85 (13.78)	188.15 (13.72)
Vegetative (T ₂)	192.95 (13.89)	188.55 (13.73)	187.57 (13.70)	190.94 (13.82)	190.00 (13.78)
Flowering (T ₃)	191.22 (13.83)	190.09 (13.79)	189.18 (13.75)	192.33 (13.87)	190.71 (13.81)
Mean	191.62 (13.84)	188.41 (13.73)	187.41 (13.69)	191.04 (13.82)	

C.D. (p≤0.05)				
	Stage	:	1.18	
	Dose	:	0.93	
	Stage \times Dose	:	1.36	
Data within paranthasis	are square root tra	ofo	rmad valu	as a

Data within parenthesis are square root transformed values. Initial soil available K =194.79 kg ha⁻¹

Conclusion

Application of varying doses of SA at different growth stages had a significant influence on the plant growth and quality attributes of French bean. However, the treatment combination T_1D_2 (foliar spray of 250 ppm of SA at plant emergence stage) resulted in the superior values for majority of the parameters. The treatment combination T_1D_2 (foliar spray of 250 ppm of SA at plant emergence stage) increased the nutrient uptake efficiency of the plant which resulted in excellent growth and consequently lead to increased yield of French bean. Therefore, a spray of 250 ppm at plant emergence stage can be done to achieve virtuous plant growth and pod quality.

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