



P-ISSN: 2349-8528

E-ISSN: 2321-4902

IJCS 2020; 8(1): 1366-1369

© 2020 IJCS

Received: 16-11-2019

Accepted: 18-12-2019

P Dhal

Department of Vegetable
Science, Odisha University of
Agriculture and Technology,
Bhubaneswar, Odisha, India

GS Sahu

Department of Vegetable
Science, Odisha University of
Agriculture and Technology,
Bhubaneswar, Odisha, India

S Mohanty

Department of Seed Science and
Technology, Odisha University
of Agriculture and Technology,
Bhubaneswar, Odisha, India

SK Dash

Department of Vegetable
Science, Odisha University of
Agriculture and Technology,
Bhubaneswar, Odisha, India

P Tripathy

Department of Vegetable
Science, Odisha University of
Agriculture and Technology,
Bhubaneswar, Odisha, India

A Dhal

Department of Plant Pathology,
Odisha University of Agriculture
and Technology, Bhubaneswar,
Odisha, India

Effect of seed priming on growth behaviour of French bean *Phaseolus vulgaris* L.

P Dhal, GS Sahu, S Mohanty, SK Dash, P Tripathy and A Dhal

DOI: <https://doi.org/10.22271/chemi.2020.v8.i1s.8444>

Abstract

An experiment entitled "Effect of seed priming on growth behaviour of French bean *Phaseolus vulgaris* L." was conducted during Rabi 2018-19 in the Vegetable Research field of Department of Vegetable Science, College of Agriculture, OUAT, Bhubaneswar. The trial was conducted in Randomized Block Design with three replications and ten treatments. The treatments were T₁ (Hydro priming), T₂ (GA₃ 50 ppm), T₃ (KCl 2%), T₄ (Sodium Molybdate 500 ppm), T₅ (Vitavax 2g/kg), T₆ (*Pseudomonas fluorescens* 10%), T₇ (*Trichoderma viride* 10%), T₈ (GA₃ 50 ppm + *T. viride* 10%), T₉ (Sodium Molybdate 500 ppm + *Pseudomonas fluorescens* 10%), T₁₀ (Control) Good quality seeds of French bean variety Harsha was taken and different priming treatments were done in the laboratory followed by growing the crop in the field as per recommended package of practices. However days to 1st flowering & 50% flowering were found non-significant. Days to 50% flowering varied from 45.5 in T₆ to 48.7 in T₄. The average leaf area of the trifoliate compound leaves were taken which was highest in T₃ (148.30) followed by T₈ (140.40 cm²) and was lowest (105.8 cm²) in control. The highest stem girth (2.8 cm) was recorded in KCl (2%) primed seeds closely followed by GA₃ (50PPM) + *T. viride* (10%) (2.7cm) and vitavax treatment @ 2g/kg (2.6 cm). The number of primary branches per plant was maximum in T₃ (4.52) and minimum in T₁₀ (2.87) and this character differed significantly among the treatments. The treatments T₃ (4.52), T₈ (4.32), T₆ (4.14), T₅ (4.01) produced more branches and they are at par.

Keywords: Priming, leaf area, stem girth, flowering

Introduction

Seed priming is a pre sowing treatment which leads to a physiological state that enables seed to germinate more efficiently. Seed priming is a technique which involves water uptake by seeds followed by drying to initiate the early events of germination up to the point of radicle germination, improved seedling vigour and growth under a broad range of environments resulting in better stand establishment and alleviation of phytochrome -induced dormancy in same crop. Seed priming is controlled hydration of seeds to a level that allow pre-germinative metabolic activity to continue, but interrupt the emergence of the radicle. Seed priming improves seed performance, ensures uniformity and better establishment, enhances the yield in diverse environments, greater tolerance to environmental stress and helps to overcome dormancy. Change in seed water content, cell cycle regulation, modification of seed ultrastructure, management of oxidative stress and reserve mobilization are the major physiological and a biochemical change takes places during seed priming.

During subsequent germination, primed seeds exhibit a faster and more synchronized germination and young seedling are often more vigorous and resistant to abiotic stresses than the seedlings obtained from unprimed seeds. Priming allows some of the metabolic processes necessary for germination to occur without germination. In priming, seeds are soaked in different solutions with high osmotic potential. This prevents the seeds from absorbing in enough water for radicle protrusion thus, suspending the seeds in the lag phase. Seed priming has been commonly used to reduce the time between seed sowing and seedling emergence and to synchronize emergence. Several physiological and biochemical changes by priming in French bean were reported by Sarika *et al.*, (2013) [4]. In seed priming, the osmotic pressure and the period for which the seeds are maintained in contact with the membrane are sufficient to allow pre-germinative metabolic processes to take place within the seeds up to a level limited to that immediately preceding radicle emergence. Since germination and seedling establishment are critical steps in plant life, and the successful establishment of plant, not only depend on rapid

Corresponding Author:**P Dhal**

Department of Vegetable
Science, Odisha University of
Agriculture and Technology,
Bhubaneswar, Odisha, India

and uniform germination of seed but depend on the ability of rapid germination of the seed under environmental conditions. Besides French bean Bassi *et al.*, (2011)^[1] reported in soybean that priming with GA₃ @ 50 ppm for 2 hour enhanced emergence, germination and speed of germination as compared to non-primed seeds. Yari *et al.*, (2010)^[7] reported that seed priming techniques with KH₂ PO₄ and KCl showed good potential to enhance germination, emergence, growth and grain yield of wheat influences the germination and early growth of wheat cultivars. For this purpose, seed pre-priming or priming methods are used to increase the qualitative and quantitative performances of seedlings.

Materials and Method

The field experiment entitled “Effect of different seed priming treatments on vegetative and yield attributing characters of French bean (*Phaseolus vulgaris* L.)” was carried out during Rabi season of the year 2018-19 in the Vegetable Demonstration plot of the Department of Vegetable Science, College of Agriculture, Orissa University of Agriculture and Technology, Bhubaneswar, Odisha.

Preparation of treated seeds

In case of hydro priming the seeds were immersed inside water for about 5 hrs. Then the seeds were taken out as well as air dried followed by sowing in prepared field. In case of GA₃ 50 ppm, KCl (2%), Sodium molybdate (500 ppm), Vitavax (2g /lit), *Pseudomonas fluorescens* (10%), *Trichoderma viride* (10%), (GA₃ 50 ppm+ *Trichoderma viride* 10%), (Sodium molybdate 500 ppm+ *Pseudomonas fluorescens* 10%) solution was prepared and the same procedure was followed for treatment of the seeds.

Sowing and sampling technique

Seeds were hand dibbled to a depth of 3-4 cm, at a row spacing of 50 cm and plant to plant spacing 30 cm, in small furrows opened with the help of trench hoe and well covered with soil. From each treatment ten number plants were selected randomly were tagged for recording various biometric observations. The mean of the ten plants was considered for further analysis. The observations on various growth parameters were recorded from the selected plants. The border plants were excluded while selecting the sample plants.

Phenological Parameters

The number of days required for the plants in each plot to show initiation of flowering were observed and recorded. The number of days taken from sowing to days when 50 percent of the plants come to flowering in the field was calculated and the mean average was represented in days

Growth parameters

Ten trifoliate leaves from each treatment were collected from sample plants and leaf area was measured by leaf area meter and the average was calculated in square centimetre. Ten numbers of compound leaves of French bean were selected from the tagged plants and the total leaf weight was taken and the average leaf weight was expressed in gram. Stem girth of 10 tagged plant from each treatment was measured by thread followed by scale then stem girth per plant was computed from the observed data. During the experiment the numbers of leaves produced by the sample plants at final harvest were counted and average data was used for statistical calculation. The number of branches of the ten selected tagged plants was counted at the final harvest of the crop and the mean was

recorded as number of primary branches per plant. The plant height of ten plants was measured from the base of the plant to the base of the fully opened youngest trifoliate leaf and the average was expressed in cm.

Results and Discussion

Influence of priming on flowering

Days to first flowering as influenced by various treatments is presented in Table- 1. It was found that maximum number of days to first flowering was recorded in control that is T₁₀ (44.50) followed by 43.70 T₄ and T₈ 43.20. In treatment T₃ (KCl 2%) only 40.60 days was required for first flowering. There is no significant difference among the treatments though the control plot recorded more days to first flowering. Days to first flowering and 50% flowering was found to be much influenced by various methods of priming. The hydro priming approach resulted the least 40.3 days to first flowering closely followed by 40.6 days in KCl 2% and *P. fluorescens* (10%). However, the treatments did not produce any significant differences in days required for first flowering. The present findings are in conformity with earlier reports of Maiti *et al* (2013)^[3] and Singh *et al* (2014)^[5].

The number of days taken to 50% flowering is presented in Table-1. It was observed that in T₄ 48.70 days were taken for days to 50% flowering, followed by T₁₀ (48.50) and T₈ (47.80). It was noticed that T₆ took lowest number of days (45.50) for 50% flowering followed by T₂ in which 45.60 days was required for flowering. It was observed that none of the treatment produced any significant effect for days to 50% flowering. There is also no significant changes in days to 50% flowering. However, the treatments T₁₀ (Control), T₂ (GA₃ 50 ppm) and T₃ (KCl 2%) recorded comparatively less 45.5, 45.6 and 46.2 days to fifty percent flowering respectively. Singh *et al.* (2014)^[5] also recorded earlier plant growth due to osmo, hydro and halo priming in cowpea seeds.

The average leaf area of compound leaves is presented in Table 1. Highest leaf area of 148.3 cm² was recorded in T₃ followed by 140.4 cm² in T₈, 138.40cm² in T₂ and 132.6 cm² in T₅. The lowest average leaf area was found with T₁₀ (105.8cm²). It was observed that all of the treatments produced significant difference so far as average leaf area was concerned as compared to control. The priming with KCl (2%) recorded the highest average leaf area (148.3 cm²) closely followed by T₈ (GA₃ 50 ppm+ *T. viridae* 10%) 140.4, T₂ (GA₃ 50 ppm) 138.4, T₅ (Vitavax 2g/kg) 132.6, T₇ (*T. viridae* 10%) 132.5 and T₆ (*P. fluorescens* 10%) 131.7 which are statistically at par and significantly different from T₁ (Hydro priming) 128.9, T₄ (Sodium Molybdate 500 ppm) 128.4, T₉ (Sodium Molybdate 500 ppm + *P. fluorescens* (10%) 125.6 and control (105.8) However the leaf size in all the treatments were found to have more leaf area as compared to control.

The observation on fresh weight of leaf is presented in Table 1. The average fresh weight of leaf was highest in T₈ (3.24g) followed by T₂ (3.14g), T₃ (3.10g) and T₉ (2.91g) and T₆ (2.85) which are statistically at par. The lowest average leaf weight 2.44g was recorded in T₁. Among the treatments hydro priming recorded the least fresh weight of leaf (2.44g) in T₁. Results revealed the highest fresh leaf weight was in T₈ (3.24g) followed by T₂ (3.14 g), T₃ (3.10g), T₉ (2.91g) and T₆ (2.85) and they are at par. There is not any significant variation among the treatments showing that different priming agents as well as their combination did not have any impact on fresh weight of leaf.

The observation on stem girth is presented in Table 1. It was observed that due to different priming treatment the highest

stem girth of 2.8 cm was recorded in T₃. However T₈ recorded stem girth of 2.7cm closely followed by 2.6 cm in T₅. The lowest stem girth of 2.1 cm was recorded with control. Highest stem girth of 2.8 cm was recorded in T₃ (2.8 cm) closely

followed by T₈ (2.7 cm). However halo priming, hormo priming, combination of hormo and bio priming as well as chemical seed treatment recorded higher stem girth than control and other treatments.

Table 1: Days to first flowering, 50% flowering, leaf characters and stem girth of French bean crop as influenced by priming

	Treatment	Days to first flowering	Days to 50% flowering	Average leaf area (cm) ²	Fresh weight of leaf (g)	Stem girth in (cm)
T1	Hydro priming	40.30	47.30	128.9	2.44	2.4
T2	GA ₃ (50 ppm)	42.00	45.60	138.4	3.14	2.5
T3	KCl (2%)	40.60	46.20	148.3	3.10	2.8
T4	Sodium Molybdate 500 ppm	43.70	48.70	128.4	2.68	2.5
T5	Vitavax (2g/kg)	42.50	46.50	132.6	2.62	2.6
T6	<i>P. fluorescens</i> (10%)	40.80	45.50	131.7	2.85	2.5
T7	<i>T. viride</i> (10%)	41.20	46.50	132.5	2.56	2.4
T8	GA ₃ (50 ppm) + <i>T. viride</i> (10%)	43.20	47.80	140.4	3.24	2.7
T9	Sodium Molybdate 500 ppm + <i>P. fluorescens</i> (10%)	41.40	46.40	125.6	2.91	2.4
T10	Control	44.50	48.50	105.8	2.70	2.1
	SE (m) ±	2.00	2.17		6.09	0.14
	CD 5%	6.08	6.59		18.46	0.41
	CV%	8.26	8.02		8.04	8.32

Effect on Number of leaves, primary branch and plant height

The number of leaves per plant was recorded at harvest and is presented in Table 2. The highest number of leaves (20.5) were obtained in T₇ followed by T₃ (19.2) and T₈ (18.6) and T₂ (18.4) which are statistically at par. However, the number of leaves per plant was lowest in T₁₀ (15.2) followed by T₉ (15.6). T₇, T₃, T₈ & T₂ produced significantly higher leaves than T₁₀ (control). The no. of leaves were found to be highest (20.5) in T₇ (*T. viridae* 10%) closely followed by T₃ (KCl 2%) that is 19.2. Though there is not any definite pattern for increase or decrease of no. of leaves by priming approach either sole or in combination but the unprimed control recorded the lowest no. of leaves (15.2) during growth period. This finding is in close conformity with the results of Singh *et al.* (2016) who experimented and found priming with bio control agents increased the number of leaves and other plant growth characters.

The number of primary branches per plant as recorded at harvest is presented in Table 2. It was found that there was significant differences in number of branches per plant and it was highest with T₃ (4.52) followed by T₂ (4.14), T₈ (4.32), T₅ (4.01) and T₇ (3.98) and they are statistically at par. The minimum number of branches was recorded in T₁₀ (2.87) followed by T₄ (2.95), T₁ (3.27) and T₆ (3.29). The number of

branches were found to be highest (4.52) in (KCl 2%) treatment followed by T₈ (4.32), T₂ (4.14), T₅ (4.01) and T₇ (3.98). The least no. of branches were recorded in control (2.87). It may be concluded that when number of branches increased, there is increase in number of leaves. However there is significant difference among the treatments in producing number of branches per plant and number of leaves per plant indicating that these two characters respond well to priming approach and different environmental, edaphic, climatic and nutritional factors which had profound influence on those two characters during the period of experimentation. Soliman *et al.* (2016) also reported this type of findings in faba bean.

The data on plant height is presented in Table 2 revealed that all the priming treatments were significantly different from control (39.83). Highest plant height of 45.91 cm was observed in T₃ (KCl 2%) which was followed at par with other treatments but significantly higher than the unprimed control. Priming treatments with KCl 2% recorded the highest plant height (45.91 cm) which was at par with other treatments but significantly higher than the unprimed control (39.83 cm). This finding is in close agreement with the result of Maiti *et al.* (2013)^[3] & Ghobadi *et al.* (2014)^[2] who recorded higher plant height in halo priming when investigated in different vegetable crops.

Table 2: Number of leaves and number of branches at harvest of French bean crop as influenced by different treatments

	Treatments	Number of leaves at harvest	Primary branches at harvest	Plant height at harvest (cm)
T1	Hydropriming	16.2	40.30	3.27
T2	GA ₃ (50 ppm)	18.4	42.43	4.14
T3	KCl (2%)	19.2	45.91	4.52
T4	Sodium Molybdate 500 ppm	16.2	40.97	2.95
T5	Vitavax (2g/kg)	17.5	41.20	4.01
T6	<i>P. fluorescens</i> (10%)	15.8	40.20	3.29
T7	<i>T. viridae</i> (10%)	20.5	41.72	3.98
T8	GA (50 ppm) + <i>T. viridae</i> (10%)	18.6	43.23	4.32
T9	Sodium Molybdate 500 ppm + <i>P. fluorescens</i> (10%)	15.6	41.40	3.31
T10	Control	15.2	39.83	2.87
	SE (m) ±	0.81	0.28	1.93
	CD5%	2.47	0.86	5.84
	CV%	8.14	13.33	8.00

Conclusion

Days to first flowering mean varied from 40.30 in T₁ to 44.5 in T₁₀. There was no significant difference in days to first flowering among the treatments. Days to 50% flowering varied from 45.5 in T₆ to 48.7 in T₄. There was also no significant difference for days to 50% flowering among the treatments. The average leaf area of the trifoliate compound leaves were taken which was highest in T₃ (148.30) followed by 140.40 cm² in T₈ and was lowest 105.8 cm² in T₁₀. The unprimed control treatment recorded significantly the least average leaf size. Fresh weight of leaves as found during the experiment varied from 2.44 to 3.24 among all the treatments and there was not much variation for this character. However the unprimed plot recorded significantly less weight of fresh leaf compared to T₈ & T₃. The highest stem girth (2.8 cm) was recorded in KCl (2%) primed seeds closely followed by GA₃ (50PPM) + *T. viride* (10%) (2.7cm) and vitavax treatment @ 2g/kg (2.6 cm). The number of leaves per plant ranged from 15.2 to 20.5 and T₉ (15.60), T₆ (15.8), T₁ & T₄ (16.2), T₅ (17.5) were found to be significantly differed from other treatments for this character. The number of primary branches per plant was maximum in T₃ (4.52) and minimum in T₁₀ (2.87) and this character differed significantly among the treatments. The treatments T₃ (4.52), T₈ (4.32), T₆ (4.14), T₅ (4.01) produced more branches and they are at par.

References

1. Bassi G, Sharma S, Gill BS. Pre-sowing seed treatment and quality in-vigouration in soybean (*Glycine max* (L) Merrill). Seed Res. 2011; 31:81-84.
2. Ghobadi M, Abnavi MS, Mohammadi R. Effect of Hormonal Priming (GA₃) and Osmo priming on Behavior of Seed Germination in Wheat (*Triticum aestivum* L.). Journal of Agricultural Science. 2014; 4(9):212-215.
3. Maiti R, Rajkumar D, Jagan M, Parmanik K, Vidyasagar P. Effect of Seed Priming on Seedling Vigour and Yield of Tomato and Chilli. International Journal of Bio-resource and Stress Management. 2013; 4(2):119-125.
4. Sarika G, Basavaraju GV, Bhanuprakash K, Chaanakeshava V, Paramesh R, Radha BN. Investigation on seed viability and vigour of aged seed by priming in French bean. Veg Sci. 2013; 40:169-173.
5. Singh A, Dahiru R, Musa M, Haliru BS. Effects of osmo-priming duration on germination, emergence and early growth of cowpea (*Vigna unguiculata* (L.) Walp.) In the Sudan Savannah. Nigeria. Int J Agron. 2014; 4(3):64-68.
6. Soliman MH, Al-Juhani RS, Hushas MA, Al-Juhani FM. Effect of seed priming with Salicylic acid on seed germination and seedling growth of broad bean (*Vicia faba* L.). International Journal of Agricultural Technology. 2016; 12(6):1125-1138.
7. Yari L, Aghaalikani M, Khazaei F. Effect of seed priming duration and temperature on seed germination behaviour of bread wheat (*Triticum aestivum* L.). ARPN J Agric Biol Sci. 2010; 5:1-7.