

Original Research Article

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Effect of Muriate of Potash (MOP) and Sulphate of Potash (SOP) on Growth Characters of Green gram (*Vigna radiata* (L.) Wilczek) cv. VBN 2 in Pot and Field Condition

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ABSTRACT

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A pot culture and field experiment was conducted to study the effect of potassium on growth characters of green gram cv. VBN 2. The treatments used viz., T₁ - Absolute control, T₂ - control N, P, (-K), T₃ - 10 kg of K₂O ha⁻¹, T₄ - 20 kg of K₂O ha⁻¹, T₅ - 30 kg of K₂O ha⁻¹, T₆ - 40 kg of K₂O ha⁻¹, T₇ - 10 kg of K₂SO₄ ha⁻¹, T₈ - 20 kg of K₂SO₄ ha⁻¹, T₉ - 30 kg of K₂SO₄ ha⁻¹, T₁₀ - 40 kg of K₂SO₄ ha⁻¹. The results revealed that application of T₁₀ - 40 kg of K₂SO₄ ha⁻¹ recorded higher values for growth characters viz., plant height, number of leaves plant⁻¹, number of branches plant⁻¹, number of nodules plant⁻¹, leaf area index and chlorophyll content respectively over control.

Introduction

Pulse consumption is increasing globally due to their high nutritional value and low glycemic index. Pulses are categorized as the most important dietary predictor of survival in older citizens of various ethnicities and are the key factor in increasing the life span of populations (Darmadi-Blackberry *et al.*, 2004). In India pulse crops are grown in an area of 139.09 (MT) with an annual production level of 86.98 (MT) and

productivity of about 639 kg ha⁻¹ during 2016-2017. Production of green gram during *Kharif* season is 1.02 and 1.35 million tonnes in 2015-2016 and 2016-2017 respectively (Anon, 2017). Pulses have double the amount of proteins than cereals. The pulse protein particularly from the lentil is deficient in methionine and cysteine (sulfur containing amino acids) while rich in lysine (Belitz and Grosch, 1996). The pulses also contain a variety of anti-nutrient factors (ANFs) such as the lectins, proteinase inhibitors, non-protein

amino acids, gums, tannins, cynogens, saponins, alkaloids, phytates, etc. which are mostly destroyed in soaking, washing, cooking, and other processes (Ali and Muzquiz, 1998). Green gram is commonly known as *mung* bean in the Indian subcontinent and is widely grown in all Asian countries. 100 g of greengram gives 30 calories and consists approximately 3 g proteins, 6 g carbohydrates and 2 g dietary fibers. It provides about 15% and 45% of the recommended dietary allowance of calcium and iron, respectively. Greengram is almost nil in raffinose or other oligosaccharides and is free of flatulence-causing agent, making it suitable for convalescent, baby or senior citizen foods.

Potassium is one of the essential plant nutrients which play a vital role in various physiological, biochemical activities and are required in high amounts to maintain adequate crop growth and sustainable crop production (Mengel and Kirkby, 1987). Potassium is not only a constituent of the plant structure but it also has a regulatory function in several biochemical processes related to protein synthesis, carbohydrate metabolism and enzyme activation (Hasanuzzaman *et al.*, 2018).

The quantity of potassium absorbed by roots is second to that of nitrogen for most of the cultivated plants. Due to intensive cropping, continuous manuring and limited or no use of potassium fertilizers, the available potassium status of the soils has depleted. Soils have begun to show response to potassium application particularly under intensive use of nitrogen and phosphorus fertilizers. Sufficient amounts of potassium is required for improving the yield and quality of different crops because of its effect on photosynthesis, water use efficiency and plant tolerance to diseases, drought and cold as well for making the balance between proteins and carbohydrates.

Materials and Methods

Pot culture experiment

The pot culture experiment was conducted in pot culture yard of Department of Soil Science and Agricultural Chemistry, Faculty of Agriculture, Annamalai University, Annamalainagar, Cuddalore district, Tamil Nadu, India, located at 11°24' N latitude and 79°41'E longitude with an altitude of + 5.79 m above mean sea level.

Pot preparation

The soil collected from the field shade dried and broken into smaller clods using wooden mallet. Forty kilogram of processed soil was filled in earthen pots, which was maintained 22.5 cm soil depth and 7.5 cm space above the soil surface, so as to provide space for irrigation.

Field experiment

The field experiment was conducted at the Thennavarayanallur, Thiruvarur Taluk, Thiruvarur District- 610103, Tamil Nadu, India, located at between 10°20' 11°07' (N-S) latitude and between 79°15' -79°45' (E-W) longitude with an altitude of 10 m above mean sea level.

Treatment details

T ₁	=	Absolute control
T ₂	=	Control N,P ₂ O ₅ and (-K)
T ₃	=	10 kg of K ₂ O ha ⁻¹
T ₄	=	20 kg of K ₂ O ha ⁻¹
T ₅	=	30kg of K ₂ O ha ⁻¹
T ₆	=	40kg of K ₂ O ha ⁻¹
T ₇	=	10 kg of K ₂ SO ₄ ha ⁻¹
T ₈	=	20 kg of K ₂ SO ₄ ha ⁻¹
T ₉	=	30 kg of K ₂ SO ₄ ha ⁻¹
T ₁₀	=	40 kg of K ₂ SO ₄ ha ⁻¹

Land preparation

The plots selected for experiment was ploughed by power tiller driven rotovator and ploughed several times followed by laddering to obtain a good tilth. Weeds and stubbles were removed and the large clods were broken into smaller pieces to obtain a desirable tilth of soil for sowing of seeds. Finally, the land was leveled and plots were laid according to experimental layout.

Fertilizer application

The experimental plots received a fertilizer schedule according to the treatments. The N, P₂O₅ were applied by basal according to the treatments and required quantity of MOP and SOP were applied in the experimental plots.

After care

The weeding was done by hand hoeing. First hoeing was done on the 15 DAS and the second on 30 DAS. Need based plant protection measures were taken.

Experimental design

The factorial experiment was laid out in a Randomized Block Design (RBD) with three replications. A recommended dose of fertilizers 25 Kg of N: 50 kg of P₂O₅ ha⁻¹ was applied to all plots in the form of urea and SSP respectively. Variable doses of K₂O was applied in the form of MOP and SOP as per treatment schedule except absolute control and control (-K). Five plants from each plot were selected as random and also plants from each pot were tagged for the data collection. The following growth characters *viz.*, plant height, number of leaves plant⁻¹, number of branches plant⁻¹, number of nodules plant⁻¹, leaf area index and chlorophyll content were observed and data collected. Data analysis was done statistically which was suggested by Gomez and Gomez (1984).

Results and Discussion

Pot culture experiment

Plant height (cm)

Application of increased levels of potassium in the form of MOP and SOP resulted in significantly increased in plant height. The treatment T₁ recorded the lowest plant height of 10.9 cm and the treatment T₁₀ recorded the highest plant height of 33.7 cm. However the treatment T₆ which recorded 33.4 cm was on par with treatment T₁₀. The treatments T₉ and T₅ recorded plant height of 30.1 and 29.6 cm which were found to be statistically similar. Further the treatments T₈ and T₄ recorded plant height of 26.6 and 26.2 cm which were on par with each other. It was followed by treatments T₇ and T₃ recorded a plant height of 23.3 and 22.9 cm which were statistically on par. The treatment T₂ recorded a plant height of 19.5 cm at 30 DAS. The treatments T₁ recorded the lowest plant height of 14.8 cm and the treatment T₁₀ recorded the highest plant height of 43.6 cm. However, the treatment T₆ which recorded 43.2 cm was on par with treatment T₁₀. The treatments T₉ and T₅ recorded plant height of 39.0 and 38.4 cm which were found to be statistically similar. Further the treatments T₈ and T₄ recorded plant height of 34.6 and 34.2 cm which were on par with each other. It was followed by treatments T₇ and T₃ recorded a plant height of 30.5 and 29.9 cm which were statistically on par. The treatment T₂ recorded a plant height of 25.6 cm. The treatment T₁ recorded the lowest plant height of 27.3 cm and the treatment T₁₀ recorded the highest plant height of 60.0 cm. However, the treatment T₆ which recorded 59.6 cm was on par with treatment T₁₀. The treatments T₉ and T₅ recorded plant height of 54.8 and 54.1 cm which were found to be statistically similar. Further the treatments T₈ and T₄ recorded plant height of 49.8 and 49.3 cm which were

on par with each other. It was followed by treatments T₇ and T₃ recorded a plant height of 45.1 and 44.5 cm which were statistically on par. The treatment T₂ recorded a plant height of 39.6 cm at harvest (Fig. 1).

Number of leaves plant⁻¹

The treatment T₁ recorded the lowest number of leaves plant⁻¹ of 9.20 and the treatment T₁₀ recorded the highest number of leaves plant⁻¹ 21.10. However the treatment T₆ which recorded 20.96 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of leaves plant⁻¹ of 19.23 and 18.96 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded number of leaves plant⁻¹ of 17.39 and 17.22 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a number of leaves plant⁻¹ of 15.69 and 15.47 which were statistically on par. The treatment T₂ recorded a number of leaves plant⁻¹ of 13.69 at 30 DAS. The treatment T₁ recorded the lowest number of leaves plant⁻¹ of 12.00 and the treatment T₁₀ recorded the highest number of leaves plant⁻¹ of 24.40. However the treatment T₆ which recorded 24.26 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of leaves plant⁻¹ of 22.45 and 22.17 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded number of leaves plant⁻¹ 20.54 and 20.36 of which were on par with each other. It was followed by treatments T₇ and T₃ recorded a number of leaves plant⁻¹ of 18.77 and 18.54 which were statistically on par. The treatment T₂ recorded a number of leaves plant⁻¹ of 16.68 at 45 DAS. The treatment T₁ recorded the lowest number of leaves plant⁻¹ of 13.40 and the treatment T₁₀ recorded the highest number of leaves of plant⁻¹ 28.00. However the treatment T₆ which recorded 27.83 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of leaves plant⁻¹ of 25.71 and 25.37 which were

found to be statistically similar. Further the treatments T₈ and T₄ recorded number of leaves plant⁻¹ of 23.45 and 23.24 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a number of leaves plant⁻¹ of 21.37 and 21.10 which were statistically on par. The treatment T₂ recorded a number of leaves plant⁻¹ of 18.91 at harvest (Fig. 5).

Number of branches plant⁻¹

Usage of potassium in the form of MOP and SOP resulted in significant increase in number of branches plant⁻¹. The treatment T₁ recorded the lowest number of branches plant⁻¹ of 3.10 and the treatment T₁₀ recorded the highest number of branches plant⁻¹ 6.90. However the treatment T₆ which recorded 6.86 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of branches plant⁻¹ of 6.30 and 6.22 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded number of branches plant⁻¹ of 5.72 and 5.66 which were on par with each other. It was followed by treatments T₇ and T₃ recorded 5.17 and 5.10 number of branches plant⁻¹ which were statistically on par. The treatment T₂ recorded 4.53 number of branches plant⁻¹ at 30 DAS. The treatment T₁ recorded the lowest number of branches plant⁻¹ of 4.90 and the treatment T₁₀ recorded the highest number of branches plant⁻¹ of 8.80. However the treatment T₆ which recorded 8.76 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of branches plant⁻¹ of 8.19 and 8.10 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded 7.59 and 7.53 number of branches plant⁻¹ which was on par with each other. It was followed by treatments T₇ and T₃ which recorded a number of branches plant⁻¹ of 7.03 and 6.96. The treatments T₇ and T₃ were statistically on par with each other. The treatment T₂ recorded a number of branches

plant⁻¹ of 6.37 at 45 DAS. Application of increased levels of potassium in the form of MOP and SOP resulted in significant increase in number of branches plant⁻¹. The treatment T₁ recorded the lowest number of branches plant⁻¹ of 5.10 and the treatment T₁₀ recorded the highest number of branches of plant⁻¹ 10.20. However the treatment T₆ which recorded 10.14 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of branches plant⁻¹ of 9.40 and 9.28 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded number of branches plant⁻¹ of 8.61 and 8.54 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a number of branches plant⁻¹ of 7.88 and 7.79 which were statistically on par. The treatment T₂ recorded a number of branches plant⁻¹ of 7.02 at harvest (Fig. 3).

Leaf area index (LAI)

The treatment T₁ recorded the lowest leaf area index of 1.30 and the treatment T₁₀ recorded the highest leaf area index 1.76. However the treatment T₆ which recorded 1.75 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded leaf area index of 1.69 and 1.68 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded leaf area index of 1.62 and 1.61 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a leaf area index of 1.55 and 1.54 which were statistically on par with each other. The treatment T₂ recorded a leaf area index of 1.47 at 30 DAS. The treatment T₁ recorded the lowest leaf area index 1.65 and the treatment T₁₀ recorded the highest leaf area index of 2.16. However the treatment T₆ which recorded 2.15 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded leaf area index of 2.08 and 2.07 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded leaf area index 2.00 and 1.99 of which were on par with each other. It was followed by treatments

T₇ and T₃ which recorded a leaf area index of 1.93 and 1.92. The treatments T₇ and T₃ were statistically on par. The treatment T₂ recorded a leaf area index of 1.84 at 45 DAS. The treatments T₁ recorded the lowest leaf area index of 1.90 and the treatment T₁₀ recorded the highest leaf area index of 2.42. However the treatment T₆ which recorded 2.41 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded leaf area index of 2.34 and 2.33 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded leaf area index of 2.26 and 2.25 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a leaf area index of 2.18 and 2.17 which were statistically on par. The treatment T₂ recorded a leaf area index of 2.10 (Fig. 7).

Chlorophyll content (mg g⁻¹)

Increased levels of potassium in the form of MOP and SOP resulted in significant increase in chlorophyll a and chlorophyll b. The treatment T₁ recorded the lowest chlorophyll a and chlorophyll b of 0.27 and 0.25 mg g⁻¹ and the treatment T₁₀ recorded the highest chlorophyll a and chlorophyll b 0.51 and 0.47 mg g⁻¹. However the treatment T₆ which recorded 0.51 and 0.47 mg g⁻¹ was on par with treatment T₁₀. The treatments T₉ and T₅ recorded chlorophyll a and chlorophyll b of 0.47, 0.44 mg g⁻¹ and 0.47, 0.43 mg g⁻¹ which were found to be statistically similar. Further the treatments T₈ and T₄ recorded chlorophyll a and chlorophyll b of 0.44, 0.40 mg g⁻¹ and 0.43, 0.40 mg g⁻¹ which were on par with each other. It was followed by treatments T₇ and T₃ recorded a chlorophyll a and chlorophyll b of 0.40, 0.37 mg g⁻¹ and 0.39, 0.37 mg g⁻¹ and which were statistically on par. The treatment T₂ recorded a chlorophyll a and chlorophyll b of 0.36 and 0.33 mg g⁻¹ at 30 DAS. The treatment T₁ recorded the lowest chlorophyll a and chlorophyll b 0.46 and 0.45 mg g⁻¹ and the treatment T₁₀ recorded the highest chlorophyll a and chlorophyll b of 0.89 and

0.88 mg g⁻¹. However the treatment T₆ which recorded 0.89 and 0.88 mg g⁻¹ was on par with treatment T₁₀. The treatments T₉ and T₅ recorded chlorophyll a and chlorophyll b of 0.82, 0.81 mg g⁻¹ and 0.81, 0.80 mg g⁻¹ which were found to be statistically similar. Further the treatments T₈ and T₄ recorded chlorophyll a and chlorophyll b readings 0.76, 0.74 mg g⁻¹ and 0.75, 0.74 mg g⁻¹ which were on par with each other. It was followed by treatments T₇ and T₃ which recorded a chlorophyll a and chlorophyll b readings of 0.69, 0.68 mg g⁻¹ and 0.69, 0.67 mg g⁻¹ respectively. The treatments T₇ and T₃ were statistically similar. The treatment T₂ recorded a chlorophyll a and chlorophyll b of 0.62 and 0.61 mg g⁻¹. Significant differences in total chlorophyll content were observed due to the application of potassium (MOP and SOP) in different treatments. The treatment T₁ recorded the lowest total chlorophyll 0.52 and 0.91 mg g⁻¹ and the treatment T₁₀ recorded the highest total chlorophyll of 0.98 and 1.77 mg g⁻¹. However the treatment T₆ which recorded 0.97 and 1.76 mg g⁻¹ was on par with treatment T₁₀. The treatments T₉ and T₅ recorded total chlorophyll of 0.91, 1.63 mg g⁻¹ and 0.90, 1.62 mg g⁻¹ which were found to be statistically similar. Further the treatments T₈ and T₄ recorded total chlorophyll readings 0.84, 1.50 mg g⁻¹ and 0.83, 1.49 mg g⁻¹ which were on par with each other. It was followed by treatments T₇ and T₃ which recorded total chlorophyll readings of 0.77, 1.38 mg g⁻¹ and 0.76, 1.36 mg g⁻¹ respectively. The treatments T₇ and T₃ were statistically on par. The treatment T₂ recorded total chlorophyll of 0.69, 1.23 mg g⁻¹ at 30, 45 DAS respectively.

Field experiment

Plant height (cm)

The treatment T₁ recorded the lowest plant height of 11.2 cm and the treatment T₁₀ recorded the highest plant height of 35.2 cm.

However the treatment T₆ which recorded 34.9 cm was on par with treatment T₁₀. The treatments T₉ and T₅ recorded plant height of 31.4 and 30.8 cm which were found to be statistically similar. Further the treatments T₈ and T₄ recorded plant height of 27.7 and 27.3 cm which were on par with each other. It was followed by treatments T₇ and T₃ recorded a plant height of 24.3 and 23.8 cm which were statistically on par. The treatment T₂ recorded a plant height of 20.2 cm at 30 DAS.

Application of increased levels of potassium in the form of MOP and SOP resulted in significant increase in plant height. The treatment T₁ recorded the lowest plant height of 16.6 cm and the treatment T₁₀ recorded the highest plant height of 46.0 cm. However the treatment T₆ which recorded 45.6 cm was on par with treatment T₁₀. The treatments T₉ and T₅ recorded plant height of 41.3 and 40.7 cm which were found to be statistically similar. Further the treatments T₈ and T₄ recorded plant height of 36.8 and 36.4 cm which were on par with each other. It was followed by treatments T₇ and T₃ recorded a plant height of 32.6 and 32.1 cm which were statistically on par. The treatment T₂ recorded a plant height of 27.6 cm at 45 DAS. Potassium has been used in the form of MOP and SOP resulted in significant increase in plant height. The treatment T₁ recorded the lowest plant height of 28.8 cm and the treatment T₁₀ recorded the highest plant height of 60.8 cm. However the treatment T₆ which recorded 60.4 cm was on par with treatment T₁₀. The treatments T₉ and T₅ recorded plant height of 55.7 and 55.0 cm which were found to be statistically similar. Further the treatments T₈ and T₄ recorded plant height of 50.8 and 50.3 cm which were on par with each other. It was followed by treatments T₇ and T₃ recorded a plant height of 46.2 and 45.6 cm which were statistically on par. The treatment T₂ recorded a plant height of 40.8 cm at harvest (Fig. 2).

Number of branches plant⁻¹

The treatment T₁ recorded the lowest number of branches plant⁻¹ of 3.8 and the treatment T₁₀ recorded the highest number of branches plant⁻¹ 7.40. However the treatment T₆ which recorded 7.36 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of branches plant⁻¹ of 6.83 and 6.75 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded number of branches plant⁻¹ of 6.28 and 6.23 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a number of branches plant⁻¹ of 5.76 and 5.70 which were statistically on par. The treatment T₂ recorded a number of branches plant⁻¹ of 5.16 at 30 DAS. Increased levels of potassium in the form of MOP and SOP usage resulted in significant increase in number of branches plant⁻¹. The treatment T₁ recorded the lowest number of branches plant⁻¹ of 5.60 and the treatment T₁₀ recorded the highest number of branches plant⁻¹ of 9.80. However the treatment T₆ which recorded 9.75 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of branches plant⁻¹ of 9.14 and 9.04 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded number of branches plant⁻¹ of 8.49 and 8.43 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a number of branches plant⁻¹ of 7.89 and 7.81 which were statistically on par. The treatment T₂ recorded a number of branches plant⁻¹ of 7.18 cm at 45 DAS. The treatment T₁ recorded the lowest number of branches plant⁻¹ of 6.10 and the treatment T₁₀ recorded the highest number of branches of plant⁻¹ 12.00. However the treatment T₆ which recorded 11.93 cm was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of branches plant⁻¹ of 11.07 and 10.94 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded number of branches plant⁻¹ of 10.16 and 10.08

which were on par with each other. It was followed by treatments T₇ and T₃ recorded a number of branches plant⁻¹ of 9.32 and 9.21 which were statistically on par. The treatment T₂ recorded a number of branches plant⁻¹ of 8.33 at harvest (Fig. 4).

Leaf area index (LAI)

The treatment T₁ recorded the lowest leaf area index of 1.00 and the treatment T₁₀ recorded the highest leaf area index 1.90. However the treatment T₆ which recorded 1.89 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded leaf area index of 1.76 and 1.74 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded leaf area index of 1.62 and 1.61 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a leaf area index of 1.49 and 1.47 which were statistically on par. The treatment T₂ recorded a leaf area index of 1.34 at 30 DAS. The treatment T₁ recorded the lowest leaf area index 1.60 and the treatment T₁₀ recorded the highest leaf area index of 2.24. However the treatment T₆ which recorded 2.23 was on par with treatment T₁₀ (Fig. 8).

The treatments T₉ and T₅ recorded leaf area index of 2.14 and 2.12 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded leaf area index 2.04 and 2.03 of which were on par with each other. It was followed by treatments T₇ and T₃ recorded a leaf area index of 1.95 and 1.94 and which were statistically on par. The treatment T₂ recorded a leaf area index of 1.84 at 45 DAS. The treatments T₁ recorded the lowest leaf area index of 1.90 and the treatment T₁₀ recorded the highest leaf area index of 2.60. However the treatment T₆ which recorded 2.59 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded leaf area index of 2.49 and 2.47 which were found to be statistically similar.

Further the treatments T₈ and T₄ recorded leaf area index of 2.38 and 2.37 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a leaf area index of 2.28 and 2.27 which were statistically on par. The treatment T₂ recorded a leaf area index of 2.16 at harvest.

Total chlorophyll (mg g⁻¹)

The treatment T₁ recorded the lowest chlorophyll a and chlorophyll b of 0.28 and 0.25 mg g⁻¹ and the treatment T₁₀ recorded the highest chlorophyll a and chlorophyll b 0.49 and 0.46 mg g⁻¹. However the treatment T₆ which recorded 0.49 and 0.46 mg g⁻¹ was on par with treatment T₁₀. The treatments T₉ and T₅ recorded chlorophyll a and chlorophyll b of 0.46, 0.45 mg g⁻¹ and 0.45, 0.42 mg g⁻¹ which were found to be statistically similar. Further the treatments T₈ and T₄ recorded chlorophyll a and chlorophyll b of 0.42, 0.39 mg g⁻¹ and 0.42, 0.39 mg g⁻¹ which were on par with each other. It was followed by treatments T₇ and T₃ which recorded a chlorophyll a and chlorophyll b of 0.39, 0.36 mg g⁻¹ and 0.39, 0.36 mg g⁻¹. The treatments T₇ and T₃ were statistically on par. The treatment T₂ recorded a chlorophyll a and chlorophyll b of 0.36 and 0.33 mg g⁻¹ at 30 DAS. Significant increase in chlorophyll a and chlorophyll b was observed due to potassium application. The treatment T₁ recorded the lowest chlorophyll a and chlorophyll b 0.47 and 0.45 mg g⁻¹ and the treatment T₁₀ recorded the highest chlorophyll a and chlorophyll b of 0.89 and 0.84 mg g⁻¹. However the treatment T₆ which recorded 0.90 and 0.89 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded chlorophyll a and chlorophyll b of 0.82, 0.78 mg g⁻¹ and 0.81, 0.77 mg g⁻¹ which were found to be statistically similar. Further the treatments T₈ and T₄ recorded chlorophyll a and chlorophyll b 0.76, 0.72 mg g⁻¹ and 0.75, 0.71 mg g⁻¹ of which were on par with each other. It was followed by treatments T₇ and T₃

recorded a chlorophyll a and chlorophyll b of 0.70, 0.66 mg g⁻¹ and 0.69, 0.66 mg g⁻¹ and which were statistically on par. The treatment T₂ recorded a chlorophyll a and chlorophyll b of 0.63 and 0.60 mg g⁻¹ at 45 DAS. Significant differences in total chlorophyll content were observed due to the application of potassium (MOP and SOP) in different treatments. The treatment T₁ recorded the lowest total chlorophyll 0.54 and 0.93 mg g⁻¹ and the treatment T₁₀ recorded the highest total chlorophyll of 0.96 and 1.80 mg g⁻¹. However the treatment T₆ which recorded 0.96 and 1.79 mg g⁻¹ was on par with treatment T₁₀. The treatments T₉ and T₅ recorded chlorophyll of 0.89, 1.66 mg g⁻¹ and 0.88, 1.64 mg g⁻¹ which were found to be statistically similar. Further the treatments T₈ and T₄ recorded total chlorophyll readings 0.83, 1.53 mg g⁻¹ and 0.82, 1.52 mg g⁻¹ were on par with each other. It was followed by treatments T₇ and T₃ which recorded total chlorophyll readings of 0.77, 1.40 mg g⁻¹ and 0.76, 1.39 mg g⁻¹ respectively. The treatments T₇ and T₃ were statistically on par. The treatment T₂ recorded a total chlorophyll of 0.70, 1.26 mg g⁻¹ at 30, 45 DAS respectively.

Number of nodules plant⁻¹

Application of increased levels of potassium in the form of MOP and SOP resulted in significant increase in number of nodules plant⁻¹. The treatment T₁ recorded the lowest number of nodules plant⁻¹ of 13.80 and the treatment T₁₀ recorded the highest number of nodules plant⁻¹ of 27.00. However the treatment T₆ which recorded 26.85 was on par with treatment T₁₀. The treatments T₉ and T₅ recorded number of nodules plant⁻¹ of 24.93 and 24.62 which were found to be statistically similar. Further the treatments T₈ and T₄ recorded number of nodules plant⁻¹ of 22.89 and 22.70 which were on par with each other. It was followed by treatments T₇ and T₃ recorded a number of nodules plant⁻¹ of 21.00 and 20.76 which were statistically on par. The

treatment T₂ recorded a number of nodules plant⁻¹ of 18.78 at harvest (Fig. 6).

The application of potassium 40 kg ha⁻¹ MOP and 40 kg ha⁻¹ SOP recorded statistically growth parameters such as significant higher plant height, significant higher number of branches per plant, significant higher number of leaves per plant at different growth stages, compared to control and lower doses of muriate of potash and sulphate of potash. Plant height showed an increase with increasing levels of potassium. Plants received more potassium along with nitrogen might have encouraged the vegetative growth. In field experiments the plant height increased due to soil application of SOP produced higher plant height 30, 45 DAS and at harvest. The plant height were recorded (35.2, 46.6 and 60.80cm) which were 68, 64 and 53% higher compared to absolute control (11.2, 16.6 and 28.8cm). Similarly in pot experiments the plant height increased due to soil application of SOP produced higher plant height 30, 45 DAS and at harvest. The plant height were recorded (33.7, 43.6 and 60.0cm) which were 67, 66 and 55% higher compared to absolute control (10.9, 14.6 and 27.3cm). Potash levels along with uniform dose of nitrogen increased the plant height significantly. Redistribution of resources leading to cell enlargement and cell division (Karivaratharaju and Ramakrishnan, 1985). This increase in plant height under higher K level was mainly associated with adequacy of nutrients in soil after application and application of potassium fertilizer improves length of stem, branches, pods, seed weight and seed yield. Similar results obtained by Buriro *et al.*, (2015) and Fathima *et al.*, (2001). Kumar *et al.*, (2014) resulted significant increase in plant height with potash application can be attributed to the fact that potash enhances plant vigour and strengthens the stalk. K⁺ is essential for attaining full activity of enzyme which has an impact on numerous physiological processes

(Das, 1999). Some of them are of major relevance for the plant growth and production. These results are also in conformity with Tak *et al.*, (2013).

Potassium fertilizer gave the highest rates of plant height with significance difference from the rest of interactions and that can be explained by availability of the humidity and element of potassium in soils from the beginning of plant growth that led to an increase the speed of photosynthesis and that reflected positively on the plant height. These results are also in conformity with Shahzad *et al.*, (2014) and Mustafa *et al.*, (2016). The number of branches plant⁻¹ increased significantly with increased levels of potassium. In field experiment the number of branches plant⁻¹ increased due to the soil application of 40 kg ha⁻¹ SOP produced higher number of branches plant⁻¹ at 30, 45 DAS and at harvest (7.40, 9.80 and 12 respectively). The values were 49, 43 and 49% higher compared to absolute control (3.80, 5.60 and 6.10 respectively). The pot experiments also followed similar pattern of results. The higher number of branches plant⁻¹ recorded at 30 DAS, 45 DAS and at harvest were 6.90, 8.80 and 10.20 respectively were 55, 44 and 50% higher compared to absolute control which recorded 3.10, 4.90 and 5.10 respectively. The results of this experiment were in accordance with these Buriro *et al.*, (2015), who noticed with the application of K, the plants grew vigorously to produce more branches plant⁻¹. Number of leaves plant⁻¹ showed significant increase with increasing level of potassium. In pot culture experiment the number of leaves plant⁻¹ increased due to the soil application of 40 kg ha⁻¹ SOP. The treatment produced higher number of leaves plant⁻¹ at 30, 45 DAS and at harvest (21.10, 24.40 and 28.00) which were 56, 51 and 52% higher compared to absolute control (9.20, 12.00 and 13.40).

Fig.1 Effect of potassium on plant height (cm) of green gram VBN 2 (Pot culture experiment)

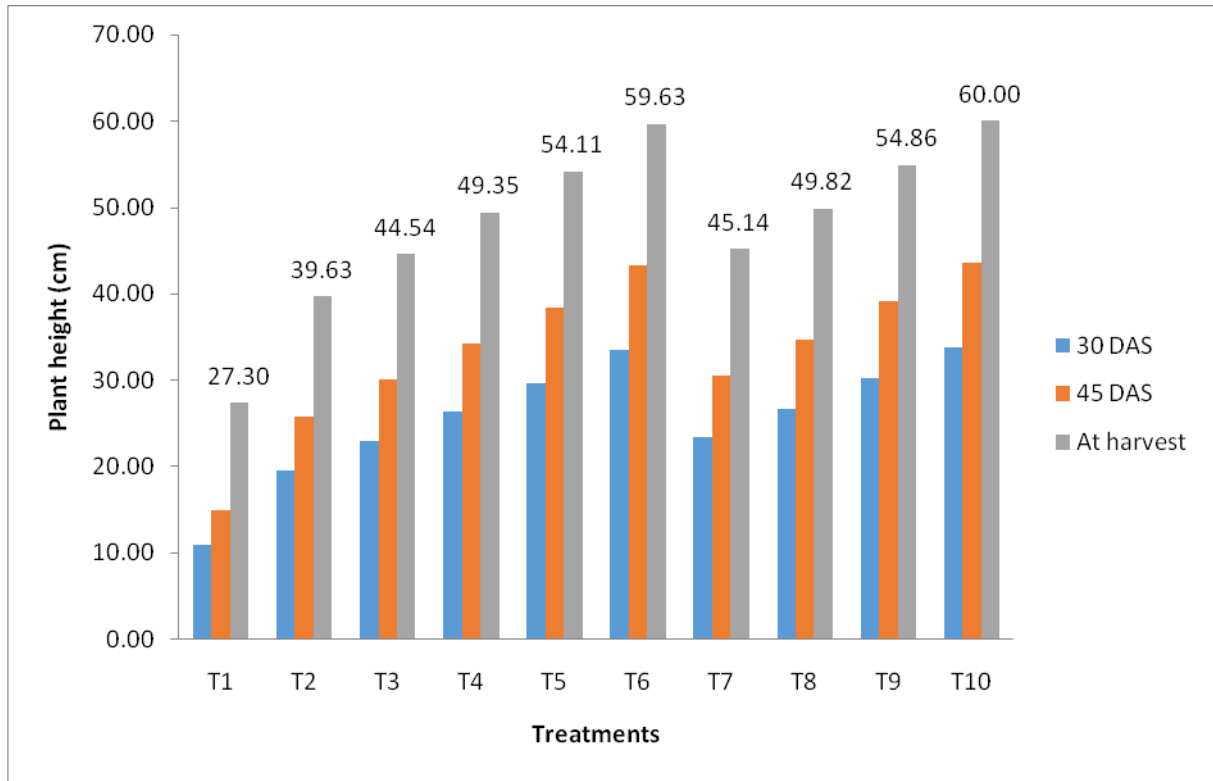


Fig.2 Effect of potassium on plant height (cm) of green gram VBN 2 (Field experiment)

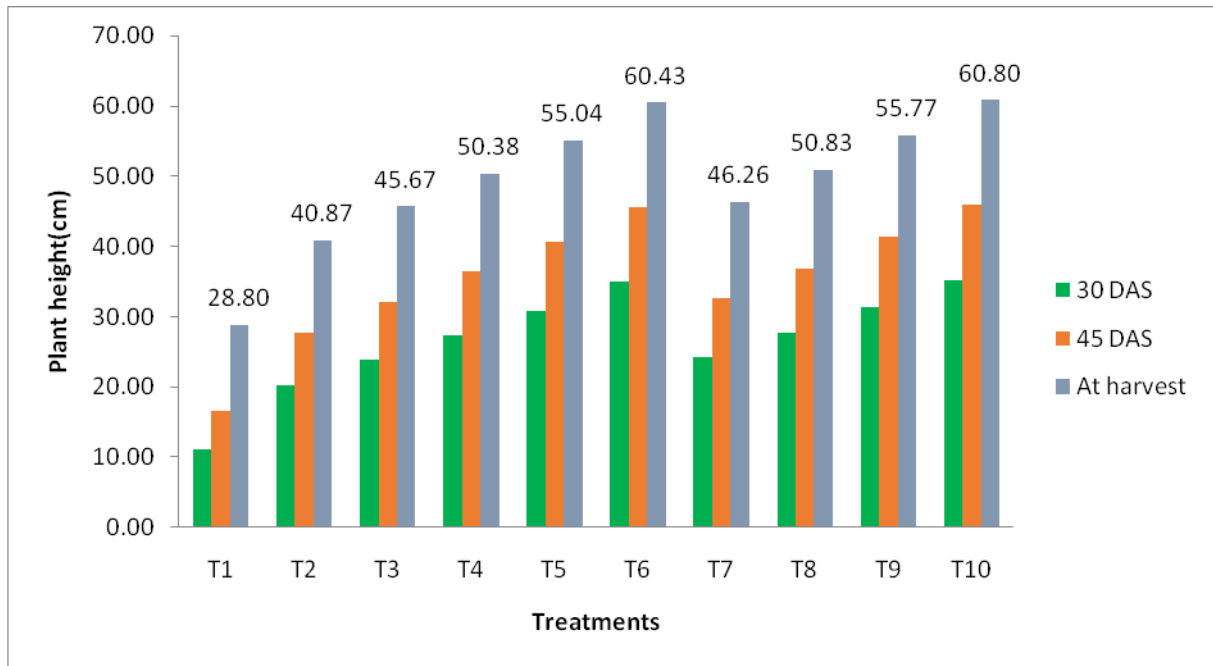


Fig.3 Effect of potassium on number of branches plant-1 of green gram VBN 2 (Pot culture experiment)

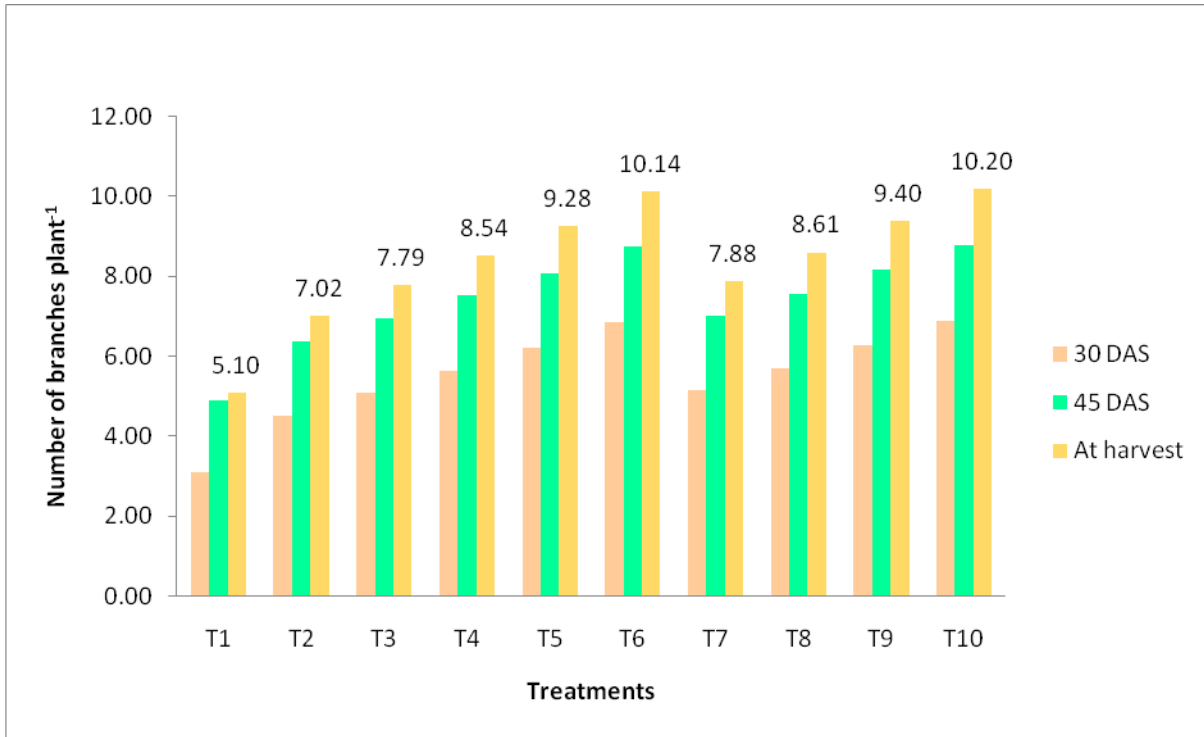


Fig.4 Effect of potassium on number of branches plant-1 of green gram VBN 2 (Field experiment)

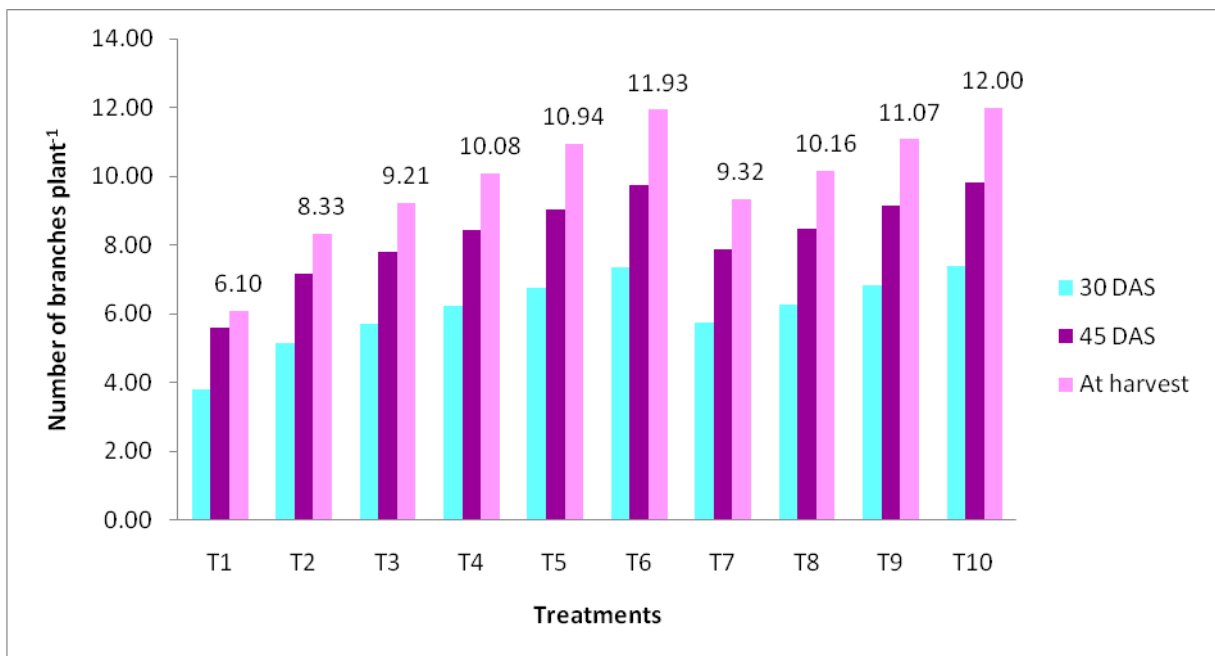


Fig.5 Effect potassium on number of leaves plant⁻¹ of green gram VBN 2 (Pot culture experiment)

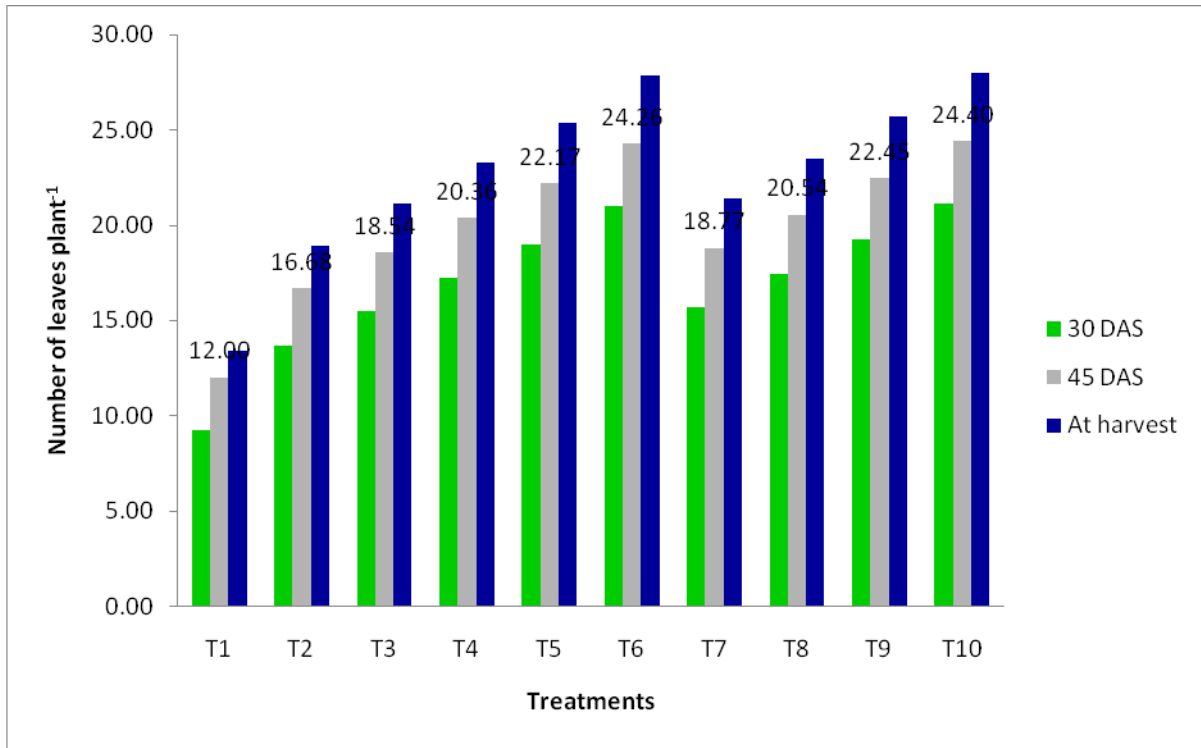


Fig.6 Effect of potassium on number of nodules plant⁻¹ of green gram VBN 2 (Field experiment)

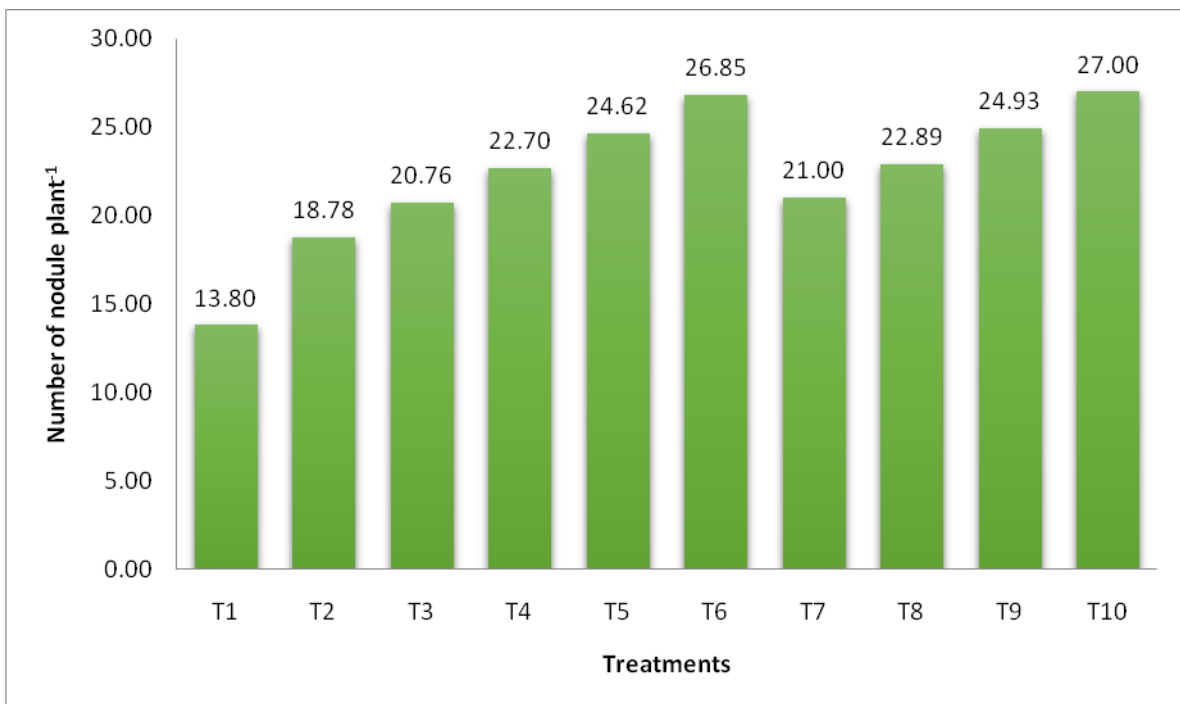


Fig.7 Effect of potassium on leaf area index of greengram at VBN 2 (Pot culture experiment)

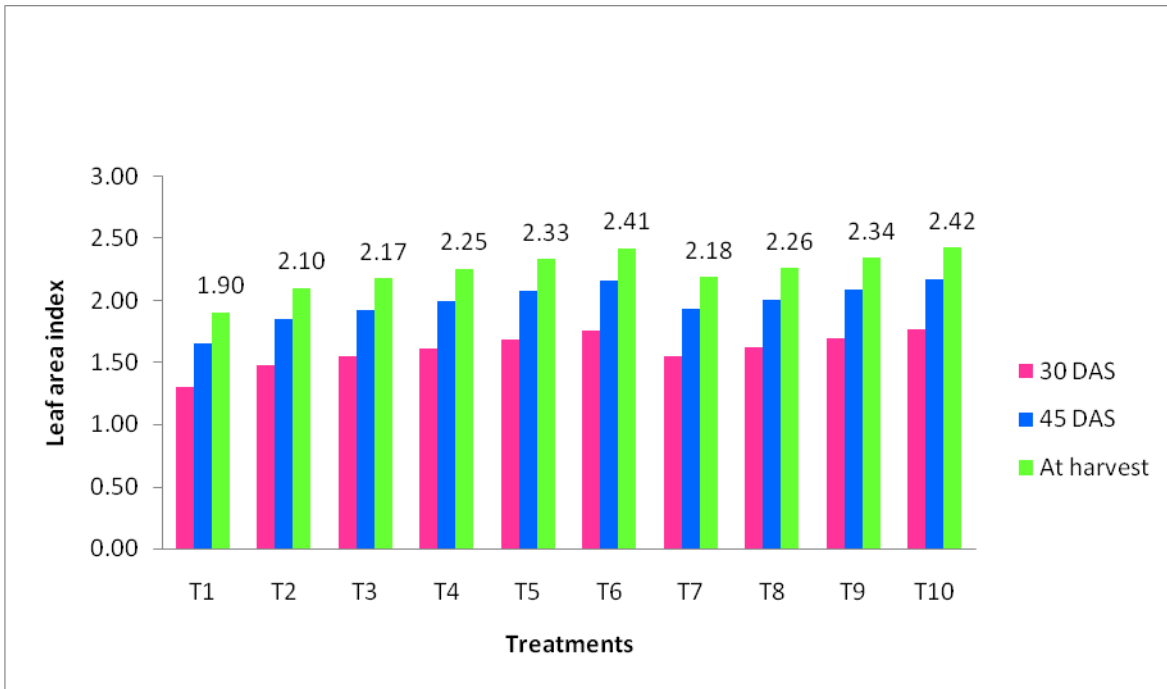
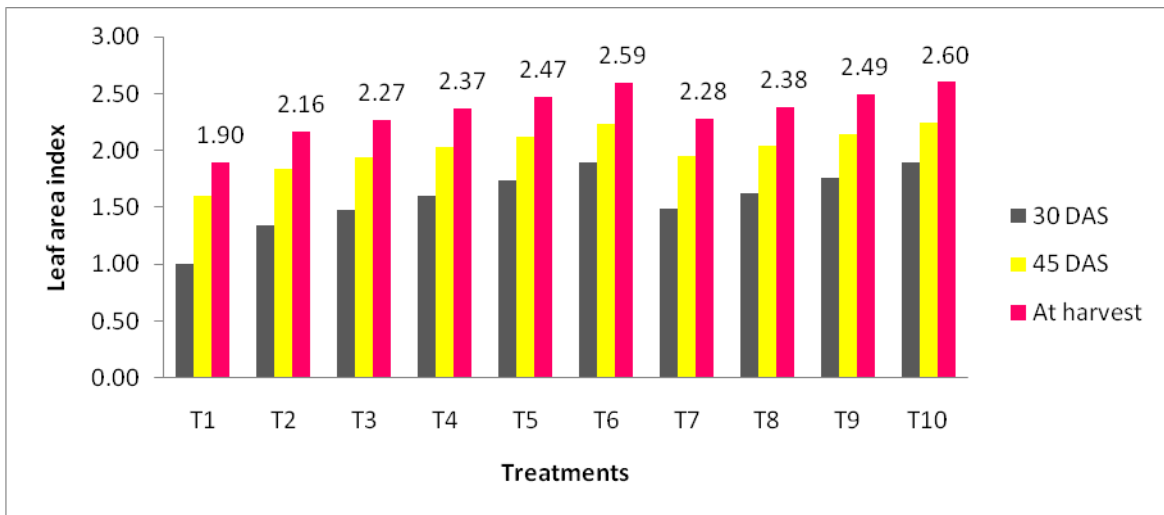


Fig.8 Effect of potassium on leaf area index of green gram VBN 2 (Field experiment)



Potassium application increased the availability of nitrogen and phosphorus, which resulted in better more number of branches plant⁻¹. Similar findings were reported by Kumar *et al.*, (2014) and Sahai (2004). Highest number of branches may be due to K application increased the availability of nitrogen and phosphorous. Similar findings

were reported by Ali *et al.*, (1996) and Khairul Mazed *et al.*, (2016). Total Chlorophyll content showed an increase with increasing level of potassium. Chlorophyll content increased due to the soil application of SOP @ 40 Kg ha⁻¹ produced higher chlorophyll content at 30, 45 DAS were recorded 0.96, 1.80 which were 44 and 48%

higher compared to absolute control in field experiment. The total chlorophyll content at 30, 45 DAS were recorded 0.98, 1.77 which were 47 and 49% higher compared to absolute control in pot experiment. K application not only enhanced the availability of other nutrient but also increased the photosynthesis rate of mung bean Kumar *et al.*, (2018). Adequate supply of potassium and phosphorus nutrient increase chlorophyll content in plants. These results are found to be similar with the results from Fletcher *et al.*, (1982), Mfillage *et al.*, (2014), Zhao *et al.*, (2001). Significantly highest chlorophyll, carotenoids content as a result of foliar K nutrition could be attributed to the mode of action of macro elements in enhancing the photosynthetic activity Doss *et al.*, (2013). Number of nodules plant⁻¹ showed an increase with increasing level of potassium. Number of nodules plant⁻¹ increased due to the soil application of 40 Kg ha⁻¹ of SOP.

The treatment produced higher number of nodules plant⁻¹ at harvest was 49% higher compared to absolute control in field experiment. The above results obtained in the study are in conformity with the results of Khan and Prakash (2014), Kurdali *et al.*, (2002), Mir *et al.*, (2012) and Tahir *et al.*, (2013). Increase in the number of nodules plant⁻¹ might be due to addition of K applied in the initial stage which might have helped in the formation and growth of roots and formation of nodules has been reported by Sathiyamoorthi *et al.*, (2008). Application of potassium (macronutrients) might have caused increased internal root growth, and enhanced the rhizobium activity in legumes. Similar findings were reported by Jack *et al.*, (2000) and Suryalakshmi (2013). Leaf area index showed an increase with increasing level of potassium. Leaf area index significantly increased due to the soil application of 40 Kgha⁻¹ of SOP. The treatment produced higher leaf area index at

30, 45 DAS and harvest stage (which were 47, 29 and 27% higher compared to absolute control in field experiments. Similar trend of results were observed in pot experiments. The leaf area index at 30, 45 DAS were 26, 24 and 21% higher compared to absolute control. The observed higher leaf area due to K⁺ may be ascribed to its role in augmenting the cell size. The important role of potassium in the process of division and elongation of the cells and that reflects positively on leaf area Mengal and Arneke (1982), Tak *et al.*, (2013). Application of potassium fertilizer gave the highest leaf area and with a significance difference. Leaf area index is owing to more number of branches and leaves. This might be due to optimum supply of nutrients which increased the plant growth, leaf number, leaf length and breadth. Similar results were also observed by Geetha and Velayutham (2009).

The present study concludes that application of potassium in the form of SOP (K₂SO₄) at 40 kg ha⁻¹ increases the growth characters. This increase was due to the fact that role of potassium induces the process of division and elongation of cells and that reflects positive growth in plants.

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