

Studies on effect of terminal heat stress on seed quality and its mitigation in chickpea (*Cicer arietinum* L.)

Abstract

A field experiment was conducted to know the effect of terminal heat stress on seed yield and its mitigation in chickpea was carried out during 2021-22 at Seed Unit, University of Agricultural Sciences, Raichur. High temperature during sowing time from early to late and very late directly affect the vegetative and anthesis stages and it was overcome by spraying the plants with heat stress mitigating chemicals. The experiment was laid out with three dates of sowing and ten foliar spray each treatment was replicated twice in a split plot design. Spraying was done at two stages of crop growth *i.e.*, at vegetative (35-40 DAS) and anthesis stage (50-60 DAS) in all dates of sowing. The interaction between dates of sowing and heat stress mitigating chemicals showed highest physiological and seed quality parameters *i.e.* chlorophyll stability index (D₃T₃) (73.06 and 74.63%), minimum cell membrane injury index (55.09 and 63.63 %), maximum proline content (D₃T₃) (6.79 and 9.37 μ mol g⁻¹), relative water content (83.79 and 75.10 %) at 60 DAS and at harvest, first count (90.50 %), final count (100.00 %), speed of germination (43.80), minimum time for radicle emergence (41.00 hrs), root length (17.32 cm), shoot length (10.12 cm), dry weight (27.10 mg), seedling vigour index-I (2643), seedling vigour index-II (2700). Among the treatments, sowing done at October 15th and plants sprayed with salicylic acid @ 400 ppm followed by gibberellic acid (100ppm) (T₁₀) twice at vegetative and anthesis stage was found to be better in obtaining significantly higher physiological and seed quality parameters in chickpea variety JG-11 under heat stress conditions.

Key words: chickpea, dates of sowing, heat stress, foliar spray

2. INTRODUCTION

Chickpea (*Cicer arietinum* L.) is an annual legume and is the third most consumed legume crop, which is widely cultivated as a winter crop for its typically yellow-brown, pea like seeds in arid and semi-arid areas around the world. Chickpea (*Cicer arietinum* L.) belongs to genus *Cicer*, tribe cicerace, family sp. Fabaceae and subfamily Faboideae. Chickpea popularly known as gram, bengal gram, homes, chhola, garbanzo bean is one of the first seed legumes to be domesticated by humans in old world.

Heat stress is the increasing temperature over the optimum range of temperature during the growth and development of plant. Late planting of chickpea in India is very common due to the wide spread intensive cropping system which often delays the sowing of chickpea. As a result, portion of the maturity period of the crop is pushed forward and thus has to face higher temperature of the summer as well as hot spells, often occurring at the time of maturity.

Reproductive growth stage (flowering and podding) in chickpea is known to be very sensitive to changes in external environment and heat stress at this stage leads to reduction in seed yield. Drastic reductions in chickpea seed yields were observed when plants at flowering and pod development stages were exposed to high (35°C) temperatures (Summerfield *et al.*, 1984). Heat stress adversely affects pollen viability, fertilization, seed development, plant photosynthesis, growth, development, reproduction and metabolism. Therefore, the seriousness of high temperature stress depends on its timing, duration and intensity on crops.

Foliar spray is a technique of feeding nutrients to plant by applying liquid chemicals directly to crop canopy. If used widely, can be more efficient, economical, environment friendly, target oriented when used to supplement soil fertilization. Now-a-days, foliar spray is widely adopted strategy in modern crop management where to ensure higher or optimum crop performance by enhancing crop growth. Foliar application overcome soil fertilization limitations, soil unsuitable for fertilizer precipitation, antagonism between certain nutrients, heterogenic soil unsuitable for low dosages and fixation.

Chickpea is grown during *rabi* season under reducing soil moisture conditions without any irrigation. As a result, there was water deficit for crop at critical stages which affects nutrient uptake ultimately causing yield reduction. To increase the yield during heat stress conditions, we have to take into consideration not only the normalization of plant water regime, but also the normalization of plant feeding and elimination of created deficiencies of some elements. Hence, various foliar spray chemicals used for heat stress mitigation can be helpful for achieving better yield from heat affected plants. Various studies have reported that foliar application of plants improves tolerance to heat stress as compared to non-sprayed plants. Through the present investigation, the conditions of cool winter followed by terminal heat stress which is prevalent in the northern dry zone of Karnataka is trying to mitigate by using heat stress mitigating chemicals at different sowing dates.

2. METHODOLOGY

A field experiment was conducted at the seed production block, Seed Unit, Monitoring Agricultural Resources, University of Agricultural Sciences, Raichur, Karnataka. The crop was sown at three different times to achieve normal (October 1st fortnight), late (November 1st fortnight) and very late sowing (December 1st fortnight) conditions (Plate 1) in split-plot design with ten foliar spray treatments viz., control, salicylic acid (800 ppm and 400 ppm), ascorbic acid (10 ppm), KCl (1%), thiourea (400 ppm), cycocel (1000 ppm), KNO₃ (0.3%), chickpea magic (8g/1), gibberellic acid (100ppm) each treatments was replicated twice in a split plot design during *rabi* season 2020-22. Spraying was done at two stages of crop growth i.e., at vegetative (35-40 DAS) and anthesis stage (50-60 DAS) in all dates of sowing.

2.1 Physiological parameters

2.1.1 Chlorophyll stability index (%)

Green plants pigments are thermo-sensitive and degradation occurs when they are subjected to higher temperature. This method is based on pigment changes induced by heating. Chlorophyll stability is the function of temperature and this property of chlorophyll stability was found to have good correlation with drought resistance. Representative leaf sample was placed in two clean tubes with 50 ml of distilled water. One tube was then subjected to heat on water bath at 65 °C ± 1°C for exactly 30 minutes. The chlorophyll in both the samples was extracted by placing the sample in 7 ml of DMSO at 65 °C for 30 minutes. The supernatant was decanted and the tissue will be discarded, then volume was made to 10 ml by DMSO. Finally, the absorbance of the extract was read at 645, 652 and 663 nm using DMSO as blank (Hiscox and Isrealstam, 1979).

$$\text{Total Chlorophyll content (mg/g)} = \frac{[20.2(\text{OD}_{645}) + 8.02(\text{OD}_{663})] \times V}{1000 \times W \times a}$$

Where,

A_{645} = Absorbance of the extract at 645 nm

A_{663} = Absorbance of the extract at 663 nm

a = Path length of light (1 cm)

V = final volume of the chlorophyll extract (10 ml)

W = Fresh weight of the sample (100 mg)

$$\text{CSI (\%)} = \frac{C_s}{C_c} \times 100$$

Where,

CSI = chlorophyll stability index

C_s = chlorophyll content of stressed plant

C_c = chlorophyll content of control plant

a = Path length of light (1 cm)

2.1.2 Cell membrane injury index (%)

The membrane injury index (MII) will be determined according to the method of Premchandra *et al.* (1990). Shoot portion (0.1 g) different treatments and control were thoroughly washed in running tap water and double distilled water and thereafter placed 10 ml of double distilled water at 40 °C for 30 minutes. After the end of this period their electrical conductivity was recorded by EC meter (C_1). Subsequently the same samples were placed on boiling water bath (100 °C) for 10 min and their electrical conductivity is recorded as above (C_2).

$$\text{MII} = (C_1 / C_2) * 100$$

Where,

$C_1 = \text{EC at } 40^\circ\text{C}$

$C_2 = \text{EC at } 100^\circ\text{C}$

2.1.3 Proline content ($\mu\text{ mol g}^{-1}$ fr.wt.)

Leaf sample (0.5 g) was homogenized in 5.0 ml of sulphosalicylic acid (3%) using mortar and pestle. The homogenate is filtered through whatman No. 1 filter paper and filtrate was collected, which was used for the estimation of protein content. 2.0 ml of extract was taken in test tube and to it 2.0 ml of glacial acetic acid was added. The reaction mixture was heated in boiling water bath at 100°C for 30 min brick red color developed after cooling the reaction mixture, 6.0 ml of toluene was added and then transferred to a separating funnel (Plate 2). After through mixing the chromophore containing toluene through mixing the chromophore containing toluene was separated and its absorbance read at 520 nm in spectrophotometer against toluene blank.

$$\text{Proline content} = \frac{\frac{\mu\text{g proline}}{\times\text{ml toluene}}}{\text{ml}} \times \frac{115.13 \mu\text{g}}{\mu\text{mol}} \times \frac{\text{g sample}}{5}$$

2.1.4 Relative water content (%)

Plant leaves generally have lower (more negative) water potential than pure water (0.0), Hence, they osmotically absorb water and become turgid. A measure of this property is

relative water content (RWC) which expresses the leaf water content (%) of the turgid leaf water content (Plate 5).

The relative water content was estimated based on the method by Barrs and Weatherley (1962). The leaf discs were taken from 3rd fully matured leaf from top of the plant tip and were weighed to indicated as fresh weight (FW). Immediately after weighing, the leaf discs were transferred to petri dishes containing water. After 24 hours, leaf material was surface blotted and were weighed to indicated as turgid weigh (Plate 3). The leaf discs were then oven dried at 80 °C up to 48 hrs and their dry weight was recorded. By using all these parameters, relative water content was calculated.

$$\text{RWC} = \frac{\text{FW}-\text{DW}}{\text{TW}-\text{DW}} \times 100$$

Where,

FW- Fresh weight (mg)

TW- Turgid weight (mg)

DW- Dry weight (mg)

2.2 Seed quality parameters

2.2.1 First count

The seed germination test was conducted in four replicates of 100 seeds each by following between paper method and the rolled towels was incubated in the walk-in seed germination chamber maintained at 25 ± 2 °C temperature and 90 ± 5 per cent relative humidity. The number of normal seedlings from each replication will be counted at the end of the 5 days and the mean germination percentage was calculated.

$$\text{Germination (\%)} = \frac{\text{Number of normal seedlings}}{\text{Total number of seeds}} \times 100$$

2.2.2 Final count

The seed germination test was conducted in four replicates of 100 seeds each by following between paper method and the rolled towels was incubated in the walk-in seed germination chamber maintained at 25 ± 2 °C temperature and 90 ± 5 percent relative

humidity. The number of normal seedlings from each replication was counted at the end of the 8 days and the mean germination percentage was calculated.

$$\text{Germination (\%)} = \frac{\text{Number of normal seedlings}}{\text{Total number of seeds}} \times 100$$

2.2.3 Speed of germination

The speed of germination was calculated by using the formula suggested by Maguire (1962).

$$\text{Speed of germination} = G_1/D_1 + G_2/D_2 + \dots + G_n/D_n$$

Where,

G_1, G_2, \dots, G_n are the number of seeds germinated on D_1, D_2, \dots, D_n day.

2.2.4 Time for radical emergence

From the germination test, 10 seeds were randomly selected from each treatment and replication and looked into it for every 4 hours to check the emergence of radical upto 2 mm in length. The time taken for emergence is recorded and the mean time was calculated and expressed in hours.

2.2.5 Root length (cm)

From the germination test, ten normal seedlings were selected randomly from each treatment on 8th day. The root length was measured from the tip of primary root to base of hypocotyls and mean root length was expressed in centimeters.

2.2.6 Shoot length (cm)

From the germination test, the ten random seedlings were used for measuring shoot length. The shoot length was measured from the base of primary leaf to the base of hypocotyls and the mean shoot length was expressed in centimeter.

2.2.7 Dry weight (mg)

From the germination test the same ten seedlings used for measuring the root and shoot length along with another five seedlings was kept in a butter paper packed and dried in hot air oven maintained at 70 °C for 24 hours. Then the seedlings were cooled in a desiccator for 30 minutes and the weight of dried seedling was recorded using an electronic balance and was expressed in milligram.

2.2.8 Seedling vigour index - I

The seedling vigour index-I was computed using the formula as suggested by Abdul-Baki and Anderson (1973) as follows

$$\text{Seedling vigour index-I} = \text{Germination (\%)} \times \text{Mean seedling length (cm)}$$

2.2.9 Seedling vigour index - II

The seedling vigour index-II was computed by multiplying the germination (%) with seedlings dry weight (g) as follows.

2.3 Statistical analysis and interpretation of data

In order evaluate comparative performance of various treatments, the data was analyzed by the technique of analysis of variance given by Fischer (1950). The collected research data were analyzed statistically by the method of Panse and Sukhatme (1967) wherever the results were significant, the critical difference (CD) was calculated at 1 percent level of significance for laboratory observation ($P < 0.05$).

3 RESULTS AND DISCUSSION

3.1 Physiological parameters

Chlorophyll is one of the major components of the chloroplast and is found to be positively correlated with the photosynthetic rate. High temperature has been found to be associated with chlorophyll content and the ability to stay green. The Chlorophyll stability index showed non-significant results due to the interaction effect of date of sowing and heat stress mitigating chemicals was shown in Table 1. Among the interactions, normal date of sowing and foliar spray of salicylic acid @ 400 ppm (D_1T_3) exhibited lowest chlorophyll stability index (67.76 and 71.06 %) at 60 DAS and at harvest, and it was followed by normal dates of sowing and foliar spray of gibberellic acid @ 100 ppm (D_1T_{10}) (65.74 and 69.36 %) as compared to all other treatments. However, plants sprayed with salicylic acid @ 400 ppm

under very late sowing recorded the maximum chlorophyll stability index (D_3T_3) (73.06 and 74.63%) at 60 DAS and at harvest. Whereas plants without any spray under very late sowing recorded minimum chlorophyll stability index (D_3T_1) (65.96 and 67.63 %) at 60 DAS and at harvest. This increase in chlorophyll stability index at 60 days after sowing and at harvest was observed in salicylic acid sprayed plants under very late sowing condition compared to control might be due to increased activity of chlorophyll which deal with the heat stress leads to increase in chlorophyll in soybean (Ghassemi-Golezani *et al.* 2018).

The membrane injury index is another physiological index that has been widely used to evaluate heat tolerance. Interaction between date of sowing and heat stress mitigating chemicals, the cell membrane injury index showed non-significant results was shown in Table 1. Normal date of sowing and plants sprayed with salicylic acid @ 400 ppm (D_1T_3) exhibited the minimum cell membrane injury index (55.09 and 63.63 %) at 60 DAS and at harvest and it was followed by normal dates of sowing and foliar spray of gibberellic acid @ 100 ppm (D_1T_{10}) (56.17 and 64.05 %) as compared to all other treatments. However, very late sowing and plants sprayed with salicylic acid @ 400 ppm (D_3T_3) (57.63 and 66.12) exhibited the minimum cell membrane injury index when compared to control. Whereas very late sowing without any spray recorded the maximum cell membrane injury index (D_3T_1) (72.03 and 73.15 %) at 60 DAS and at harvest. This decrease in cell membrane injury index at 60 days after sowing and at harvest was observed in salicylic acid sprayed plants under very late sowing condition compared to control might be due to exogenously applied salicylic acid significantly reduced the ion leakage and lipid peroxidation that acts to deal with heat stress leading to higher chlorophyll content in chickpea (Khetrapal *et al.* 2009).

Further, due to the interaction effect between date of sowing and heat stress mitigating chemicals, proline content showed non-significant results was shown in Table 2 (Plate 4). Interaction between normal date of sowing and foliar spray of salicylic acid @ 400 ppm (D_1T_3) exhibited minimum proline content (4.68 and 8.99 $\mu\text{ mol g}^{-1}$) at 60 DAS and at harvest, and it was followed by normal dates of sowing and foliar spray of gibberellic acid @ 100 ppm (D_1T_{10}) (4.54 and 8.75 $\mu\text{ mol g}^{-1}$) as compared to control. However, plants sprayed with salicylic acid @ 400 ppm under very late sowing condition recorded the maximum proline content (D_3T_3) (6.79 and 9.37 $\mu\text{ mol g}^{-1}$) at 60 DAS and at harvest. Whereas plants sprayed with salicylic acid @ 400 ppm (D_1T_1) under normal sowing conditions recorded lowest proline content (5.63 and 8.79 $\mu\text{ mol g}^{-1}$) at 60 DAS and at harvest. In this present

study, the results indicated that, 17.0 and 6.1 per cent increase in proline content at 60 days after sowing and at harvest was observed in salicylic acid sprayed plants under very late sowing condition compared to control might be due to salicylic acid respond to heat stress by accumulating certain specific metabolites such as amino acids, proteins and proline under stress, the increase in proline content had beneficial in enhancing plant resistance to stress thereby developing adaptations in plants to survive under environmental stress in black gram (Baroowa and Gogoi 2012).

Relative water content showed non-significant results due to the interaction of date of sowing and heat stress mitigating chemicals was shown in Table 2. The interactions between normal date of sowing and plants sprayed with salicylic acid @ 400 ppm (D₁T₃) exhibited maximum relative water content (83.79 and 75.10 %) at 60 DAS and at harvest and it was followed by normal dates of sowing and foliar spray of gibberellic acid @ 100 ppm (D₁T₁₀) (82.96 and 74.31 %) as compared to all other treatments. Whereas minimum relative water content was recorded under very late sowing without any spray (D₃T₁) (61.21 and 59.93 %). In the present study, salicylic acid sprayed plants showed 12.0 and 7.2 per cent increase in relative water content at 60 days after sowing and at harvest under very late sowing conditions compared with control, this might be due to the fact that salicylic acid respond to heat stress from osmoregulation by increasing the production of osmolytes, as ions or sugars are often accumulated in plants under heat stress conditions in chickpea (Gunes *et al.* 2008).

3.2 Seed quality parameters

First count showed non-significant results due to interaction between date of sowing and heat stress mitigating chemicals was shown in (Table 3). Among the interactions, normal date of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) exhibited maximum first count (90.50 %) and it was followed by normal dates of sowing and foliar spray of gibberellic acid @ 100 ppm (D₁T₁₀) (90.00 %) as compared to all other treatments. Whereas very late sowing without any spray recorded the minimum first count (control) (D₃T₁) (86.50 %).

Interaction between date of sowing and heat stress mitigating chemicals, final count yielded non-significant results was shown in Table 3 (Plate 5). Interaction between normal date of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) exhibited maximum final count (100.00 %) and it was followed by normal dates of sowing and foliar spray of

gibberellic acid @ 100 ppm (D₁T₁₀) (99.00 %) as compared to all other treatments. Whereas very late sowing without any spray recorded the minimum final count (D₃T₁) (97.50 %).

The non-significant variation was recorded for speed of germination due to interaction between sowing date and heat stress mitigating chemicals was shown in Table 3 (Plate 5). Among the interaction, normal date of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) exhibited the maximum speed of germination (43.80) and was followed by normal dates of sowing and foliar spray of gibberellic acid @ 100 ppm (D₁T₁₀) (42.50) as compared to all other treatments. Whereas minimum speed of germination was recorded under late sowing without any spray (D₃T₁) (36.70). The increase in germination under normal sowing conditions compared to control might be due to plants sprayed with salicylic acid that induced genes encoding heat resistance on seed germination by enhancing the physiological activity and translocation of food reserves necessary for growth and increase in these traits may also be attributed to the role of salicylic acid in increasing oxygen and nutrient uptake and the activity of enzymes in seeds as reported by (Alamri *et al.*, 2018) in wheat.

Interaction among the dates of sowing and heat stress mitigating chemicals showed non-significant results for time for radicle emergence was shown in Table 3. Among the interactions, the normal date of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) exhibited the minimum time for radicle emergence (41.00 hrs) and was followed by the normal dates of sowing and foliar spray of gibberellic acid @ 100 ppm (D₁T₁₀) (41.50 hrs) as compared to all other treatments. whereas very late sowing without any spray recorded the maximum time for radicle emergence (D₃T₁) (46.00 hrs). This quicker radicle emergence might be due to application of salicylic acid increases oxygen and nutrient uptake and the activity of enzymes in seeds in pulses (Mavi *et al.* 2010).

Root length showed non-significant results due to the interaction between date of sowing and heat stress mitigating chemicals was shown in Table 4 (Plate 6). Interaction between normal date of sowing and foliar spray of gibberellic acid @ 100ppm (D₁T₁₀) exhibited the maximum root length (17.32 cm) and it was followed by normal dates of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) (17.01 cm) as compared to all other treatments. Whereas very late sowing without any spray recorded the minimum seedling root length (D₃T₁) (15.91 cm). The two spray of gibberellic acid was given at vegetative and anthesis stage on chickpea showed the favourable effect on above parameter and it was also observed that, gibberellic acid act as a signal molecule and involved in several

physiological processes controlling the seed germination that promotes stem cell elongation and root length under heat stress conditions in chickpea (Roychowdry *et al.* 2012).

Shoot length showed non-significant results due to interaction effect between date of sowing and heat stress mitigating chemicals as shown in Table 4 (Plate 6). Among the interactions, normal date of sowing and foliar spray of gibberellic acid @ 100ppm (D₁T₁₀) exhibited the maximum shoot length (10.12 cm) and it was followed by normal dates of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) (9.13 cm) as compared to all other treatments. Whereas very late sowing without any spray recorded the minimum shoot length (D₃T₁) (7.82 cm). The experiment findings revealed that shoot length registered significantly superior in the treatment with plants sprayed with gibberellic acid was attributed to decreases the stress by increasing sucrose transport to shoots from cotyledons, which had been reduced under stress conditions and also by increasing invertase activity in shoots in chickpea (Mazid 2014).

Dry weight showed non-significant results due to interaction effect of date of sowing and heat stress mitigating chemicals as shown in Table 4. Interaction among the normal date of sowing and foliar spray of gibberellic acid @ 100ppm (D₁T₁₀) exhibited maximum dry weight (27.10 mg) and it was followed by normal dates of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) (27.00 mg) as compared to all other treatments. Whereas very late sowing without any spray recorded the minimum dry weight (D₃T₁) (22.10 mg). Moreover, plants sprayed with gibberellic acid responded better to above parameter under stress conditions may be due to enhanced the cell division and longer length in chickpea (Roychowdry *et al.* 2012).

Interaction between date of sowing and heat stress mitigating chemicals, the seedling vigour index-I showed non-significant results as shown in Table 4 (Plate 6). Among the interactions, normal date of sowing and foliar spray of gibberellic acid @ 100ppm (D₁T₁₀) exhibited the maximum seedling vigour index-I (2716) and it was followed by normal dates of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) (2614) as compared to all other treatments. Whereas very late sowing without any spray recorded the minimum seedling vigour index-I (D₃T₁) (2313). This experiment revealed even under very late date of sowing the maximum seedling vigour index might be due to gibberellic acid increased

amylase activity in the cotyledons of mung bean seedlings (Keikha *et al.* 2017).

Seedling vigour index-II showed non-significant results due to interaction between date of sowing and heat stress mitigating chemicals was shown in Table 4 (Plate 6). Interaction between normal date of sowing and foliar spray of salicylic acid @ 400 ppm (D₁T₃) exhibited maximum seedling vigour index-II (2700) and it was followed by normal dates of sowing and foliar spray of gibberellic acid @ 100 ppm (D₁T₁₀) (2682) as compared to all other treatments. Whereas very late sowing without any spray recorded the minimum seedling vigour index-II (D₃T₁) (2154). This experiment revealed the minimum seedling vigour index under very late date of sowing might be due to rise in temperature might cause damage to membranes, cellular oxidizing ability and photochemical efficiency in shoots and also reduce the activity of proteins and enzymes under very late sowing conditions in chickpea (Kaur *et al.* 1998).

4 CONCLUSION

It can be concluded that, heat stress adversely affects the physiological and seed quality parameters in chickpea. Among the different dates of sowings, normal date of sowing (October 1st fortnight) recorded highest physiological and seed quality parameters. The chickpea plants sprayed with salicylic acid @ 400 ppm (T₃) followed by gibberellic acid (100ppm) (T₁₀) acid recorded significantly higher physiological and seed quality parameters in chickpea under heat stress conditions (35.1 °C at anthesis stage and 40 °C at pod setting stage) when compared to control.

Disclaimer (Artificial intelligence)

Option 1:

Author(s) hereby declare that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc.) and text-to-image generators have been used during the writing or editing of this manuscript.

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Table 1. Effect of dates of sowing and heat stress mitigating chemicals on chlorophyll stability index and cell membrane injury index in chickpea

Treatments	Chlorophyll stability index (%)								Cell membrane injury index (%)							
	At 60 DAS				At maturity				At 60 DAS				At maturity			
	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean
T ₁	60.04	62.17	65.96	62.72	63.96	65.89	67.63	65.83	70.31	71.69	72.03	71.34	71.38	72.91	73.15	72.48
T ₂	63.63	66.02	68.92	66.19	67.45	69.17	70.52	69.05	60.76	62.09	63.54	62.13	65.61	67.1	67.83	66.85
T ₃	67.76	70.01	73.06	70.28	71.06	73.18	74.63	72.96	55.09	56.47	57.63	56.4	63.63	64.61	66.12	64.79
T ₄	62.67	65.41	67.43	65.17	66.79	68.13	70.01	68.31	62.61	64.5	65.37	64.16	66.09	67.49	68.23	67.27
T ₅	61.02	64.23	66.23	63.83	65.26	67.32	69.2	67.26	66.81	70.27	71.73	69.6	69.71	71.28	72.53	71.17
T ₆	62.93	65.28	67.56	65.26	66.74	68.24	70.54	68.51	64.01	65.49	66.82	65.44	66.81	68.42	69.96	68.4
T ₇	62.09	66.09	68.32	65.5	65.93	67.62	69.93	67.83	65.28	66.59	67.91	66.59	67.81	69.3	70.81	69.31
T ₈	63.06	66.53	68.26	65.95	67.06	68.45	71.02	68.84	62.21	63.64	65.07	63.64	65.09	67.01	67.12	66.41
T ₉	64.19	67.08	69.03	66.77	67.93	69.01	70.34	69.09	58.83	60.13	61.57	60.18	64.67	66.09	67.16	65.97
T ₁₀	65.74	67.86	69.93	67.84	69.36	70.93	72.73	71.01	56.17	57.47	58.86	57.5	64.05	65.45	66.72	65.41
Mean	63.31	66.07	68.47		67.15	68.79	70.66		62.21	63.83	65.05		66.49	67.97	68.96	
	D	T	D x T		D	T	D x T		D	T	D x T		D	T	D x T	
SEm±	0.006	1.032	1.789		0.001	1.08	1.871		0.008	1.038	1.799		0.001	1.088	1.885	
CD @ 5%	0.021	2.997	NS		0.01	3.134	NS		0.028	3.014	NS		0.01	3.159	NS	

NS: Non-significant

Sowing window

D₁ = Normal date of sowing

D₂ = Late date of sowing

D₃ = Very late date of sowing

Mitigation treatments

T₁: Control

T₂: Salicylic acid (800 ppm)

T₃: Salicylic acid (400 ppm)

T₄: Ascorbic acid (10 ppm)

T₅: KCl (1%)

T₆: Thiourea (400 ppm)

T₇: Cycocel (1000 ppm)

T₈: KNO₃ (0.3%)

T₉: Chickpea magic (8g/l)

T₁₀: GA₃ (100ppm)

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Table 2. Effect of dates of sowing and heat stress mitigating chemicals on proline content and relative water content in chickpea

Treatments	Proline content ($\mu\text{ mol g}^{-1}$)								Relative water content (%)							
	At 60 DAS				At maturity				At 60 DAS				At maturity			
	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean
T ₁	4.02	4.37	5.63	4.67	8.03	8.56	8.79	8.46	79.1	70.53	61.21	70.28	70.37	62.85	59.93	64.38
T ₂	4.28	5.01	6.23	5.17	8.46	8.72	8.98	8.72	81.36	75.69	66.1	74.38	72.91	65.01	62.2	66.71
T ₃	4.68	5.81	6.79	5.76	8.99	9.09	9.37	9.15	83.79	79.59	69.59	77.66	75.1	67.43	64.61	69.05
T ₄	4.24	4.83	6.13	5.07	8.39	8.67	8.94	8.67	80.96	74.86	65.43	73.75	72.32	64.72	61.89	66.31
T ₅	4.07	4.51	5.71	4.76	8.15	8.53	8.81	8.5	79.85	71.56	62.82	71.41	70.93	63.35	60.53	64.94
T ₆	4.23	4.85	6.13	5.07	8.38	8.65	8.93	8.65	80.51	73.63	64.03	72.72	71.8	64.13	61.13	65.69
T ₇	4.16	4.63	5.96	4.92	8.25	8.58	8.88	8.57	80.2	72.55	63.62	72.12	71.1	63.85	60.9	65.28
T ₈	4.2	4.72	6.01	4.98	8.31	8.61	8.91	8.61	81.15	74.83	65.41	73.8	72.48	64.87	61.97	66.44
T ₉	4.31	5.12	6.3	5.24	8.57	8.77	9.04	8.79	81.73	76.81	66.62	75.05	73.51	65.7	62.79	67.33
T ₁₀	4.54	5.47	6.56	5.52	8.75	8.91	9.36	9.01	82.96	78.28	68.1	76.45	74.31	66.53	63.65	68.16
Mean	4.27	4.93	6.15		8.43	8.71	9		81.16	74.83	65.29		72.48	64.84	61.96	
	D	T	D x T		D	T	D x T		D	T	D x T		D	T	D x T	
SEm \pm	0.0006	0.0798	0.1382		0.0006	0.1364	0.2362		0.012	1.158	2.006		0.0014	1.043	1.807	
CD @ 5%	0.0025	0.2316	NS		0.0024	0.3957	NS		0.041	3.361	NS		0.058	3.028	NS	

NS: Non-significant

Sowing window

D₁ = Normal date of sowing

D₂ = Late date of sowing

D₃ = Very late date of sowing

Mitigation treatments

T₁: Control

T₂: Salicylic acid (800 ppm)

T₃: Salicylic acid (400 ppm)

T₄: Ascorbic acid (10 ppm)

T₅: KCl (1%)

T₆: Thiourea (400 ppm)

T₇: Cycocel (1000 ppm)

T₈: KNO₃ (0.3%)

T₉: Chickpea magic (8g/l)

T₁₀: GA₃ (100ppm)

UNDER PEER REVIEW

Table 3. Effect of dates of sowing and heat stress mitigating chemicals on germination (first & final count), speed of germination and time for radical emergence in chickpea

Treatments	First count (%)				Final count (%)				Speed of germination				Time for radical emergence (hrs)			
	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean
T ₁	88	87.5	86.5	87.3	98.5	98.5	97.5	98.2	37.4	37.2	36.7	37.1	46	46	46	46
T ₂	89.5	88.5	88.5	88.8	98.5	98.5	98	98.3	40.7	40.5	39.8	40.3	46	46	46	46
T ₃	90.5	90.5	89.5	90.2	100	98.5	98	98.8	43.8	43.2	42.8	43.3	41	42	46	43
T ₄	89.5	88.5	88	88.7	99	98.5	98	98.5	40.7	40.6	40.3	40.5	46	46	46	46
T ₅	89.5	88.5	87	88.3	98.5	99	98	98.5	37.7	36.7	36.5	37	46	46	46	46
T ₆	89.5	88.5	88	88.7	98.5	98	98	98.2	40.5	39.9	39.5	40	46	46	46	46
T ₇	89.5	88.5	87.5	88.5	99	98	98.5	98.5	40.6	40.3	39.7	40.2	46	46	46	46
T ₈	90	89.5	88.5	89.3	98.5	98	98.5	98.3	41.3	40.9	40.5	40.9	46	46	46	46
T ₉	90	89	88.5	89.2	98.5	98	98.5	98.3	41.8	41.1	40.9	41.3	46	46	46	46
T ₁₀	90	89	88.5	89.2	99	99	98.5	98.8	42.5	42.3	41.3	42	41.5	42	46	43.1
Mean	89.6	88.8	88.05		98.8	98.4	98.2		40.7	40.2	39.8		45	45.2	46	
	D	T	D x T		D	T	D x T		D	T	D x T		D	T	D x T	
SEm±	0.177	0.334	0.578		0.318	0.395	0.684		0.016	0.887	1.537		0.04	1.008	1.747	
CD @ 1%	0.565	0.985	NS		1.371	0.81	NS		0.05	2.821	NS		0.175	2.07	NS	

NS: Non-significant

Sowing window

D₁ = Normal date of sowing

D₂ = Late date of sowing

D₃ = Very late date of sowing

Mitigation treatments

T₁: Control

T₂: Salicylic acid (800 ppm)

T₃: Salicylic acid (400 ppm)

T₄: Ascorbic acid (10 ppm)

T₅: KCl (1%)

T₆: Thiourea (400 ppm)

T₇: Cycocel (1000 ppm)

T₈: KNO₃ (0.3%)

T₉: Chickpea magic (8g/l)

T₁₀: GA₃ (100ppm)

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Table 4. Effect of dates of sowing and heat stress mitigating chemicals on chickpea root length, shoot length, dry weight, seedling vigour-I & seedling vigour-II

Treatments	Root length (cm)				Shoot length (cm)				Dry weight (mg)				Seedling vigour index-I				Seedling vigour index-II			
	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean	D ₁	D ₂	D ₃	Mean
T ₁	16.6 1	16.5 2	15.9 1	16.3 4	7.9	7.91	7.8 2	7.87	23.4	22.6	22.1	22.7	241 4	240 4	231 3	2377	230 4	222 6	215 4	2228
T ₂	16.9 2	16.9	16.5 1	16.7 7	8.92	8.92	8.7	8.84	27	26.2	25	26.1	254 5	254 3	247 0	2519	265 9	258 0	245 0	2563
T ₃	17.0 1	16.9 1	16.6	16.8 4	9.13	9.04	8.9 1	9.02	27	26.3	25.2	26.2	261 4	255 6	249 9	2556	270 0	259 0	246 9	2586
T ₄	16.7 2	16.9	16.5 2	16.7 1	8.7	8.7	8.5 2	8.64	26	26	24.5	25.5	251 6	252 1	245 3	2496	257 4	256 1	240 1	2512
T ₅	16.7	16.6 2	16	16.4 4	8.01	8.01	7.9	7.97	23.4	22.8	22.5	22.9	243 3	253 7	234 2	2437	230 4	225 7	220 5	2255
T ₆	16.7	16.3	16.3 1	16.4 3	8.52	8.42	8.4 1	8.45	25	24	23	24	248 4	242 2	242 3	2443	246 2	235 2	225 4	2356
T ₇	16.7	16.0 2	16.5 3	16.4 1	8.6	8.53	8.5	8.54	25.6	24.5	23.5	24.5	250 4	240 5	246 5	2458	253 4	240 1	231 4	2416
T ₈	16.7 1	16.7	16.2	16.5 3	8.52	8.44	8.5 2	8.49	24.5	23.6	23	23.7	248 5	246 3	243 4	2460	241 3	231 2	226 5	2330
T ₉	16.7	16.8 1	16.5 1	16.6 7	8.7	8.61	8.6	8.63	26	26	23.7	25.2	250 1	249 1	247 3	2488	256 1	254 8	233 4	2481
T ₁₀	17.3 2	17.1	16.6	17	10.1 2	9.6	9.5 2	9.74	27.1	26.6	25.6	26.4	271 6	264 3	257 2	2643	268 2	263 3	252 1	2612
Mean	16.8 7	16.6 6	16.3 6		8.71	8.61	8.5 4		25.5	24.9	23.8		252 1	249 8	244 4		251 9	244 6	233 6	
	D	T	D x T		D	T	D x T		D	T	D x T		D	T	D x T		D	T	D x T	

SEm±	0.00 2	0.26 3	0.45 5		0.00 2	0.13 3	0.2 3		0.00 2	0.38 4	0.665 5		0.21 1	38.4 8	66.6 4		0.33 4	38.3 5	66.4 3	
CD @ 1%	0.00 7	NS	NS		0.00 6	0.38 6	NS		0.00 7	1.11 4	NS		1.08 8	111. 6	NS		1.03 5	111. 3	NS	

NS: Non-significant

Sowing window

D₁ = Normal date of sowing

D₂ = Late date of sowing

D₃ = Very late date of sowing

Mitigation treatments

T₁: Control

T₂: Salicylic acid (800 ppm)

T₃: Salicylic acid (400 ppm)

T₄: Ascorbic acid (10 ppm)

T₅: KCl (1%)

T₆: Thiourea (400 ppm)

T₇: Cycocel (1000 ppm)

T₈: KNO₃ (0.3%)

T₉: Chickpea magic (8g/l)

T₁₀: GA₃ (100ppm)

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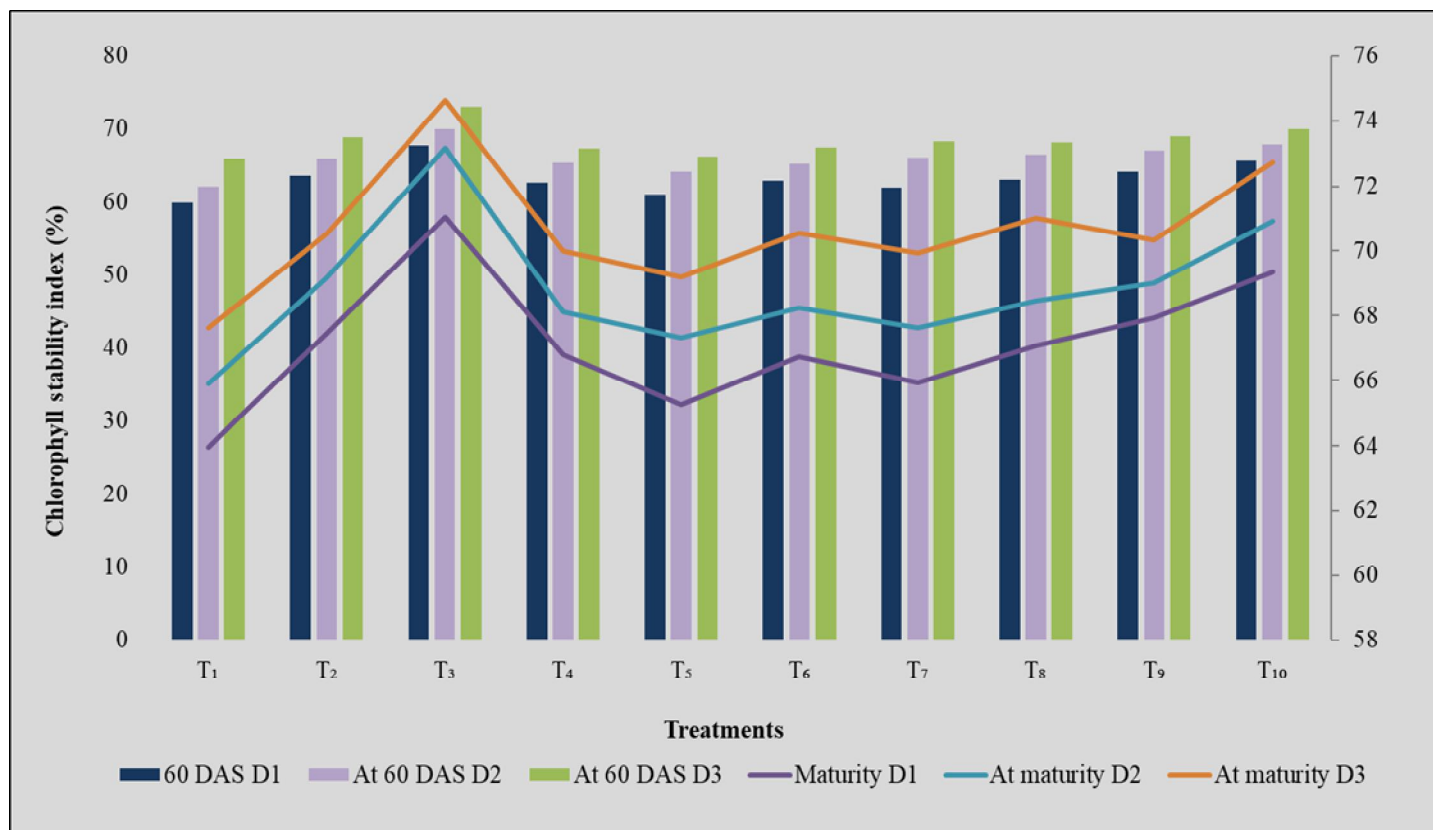


Fig. 1. Effect of dates of sowing and heat stress mitigating chemicals on chlorophyll stability index in chickpea

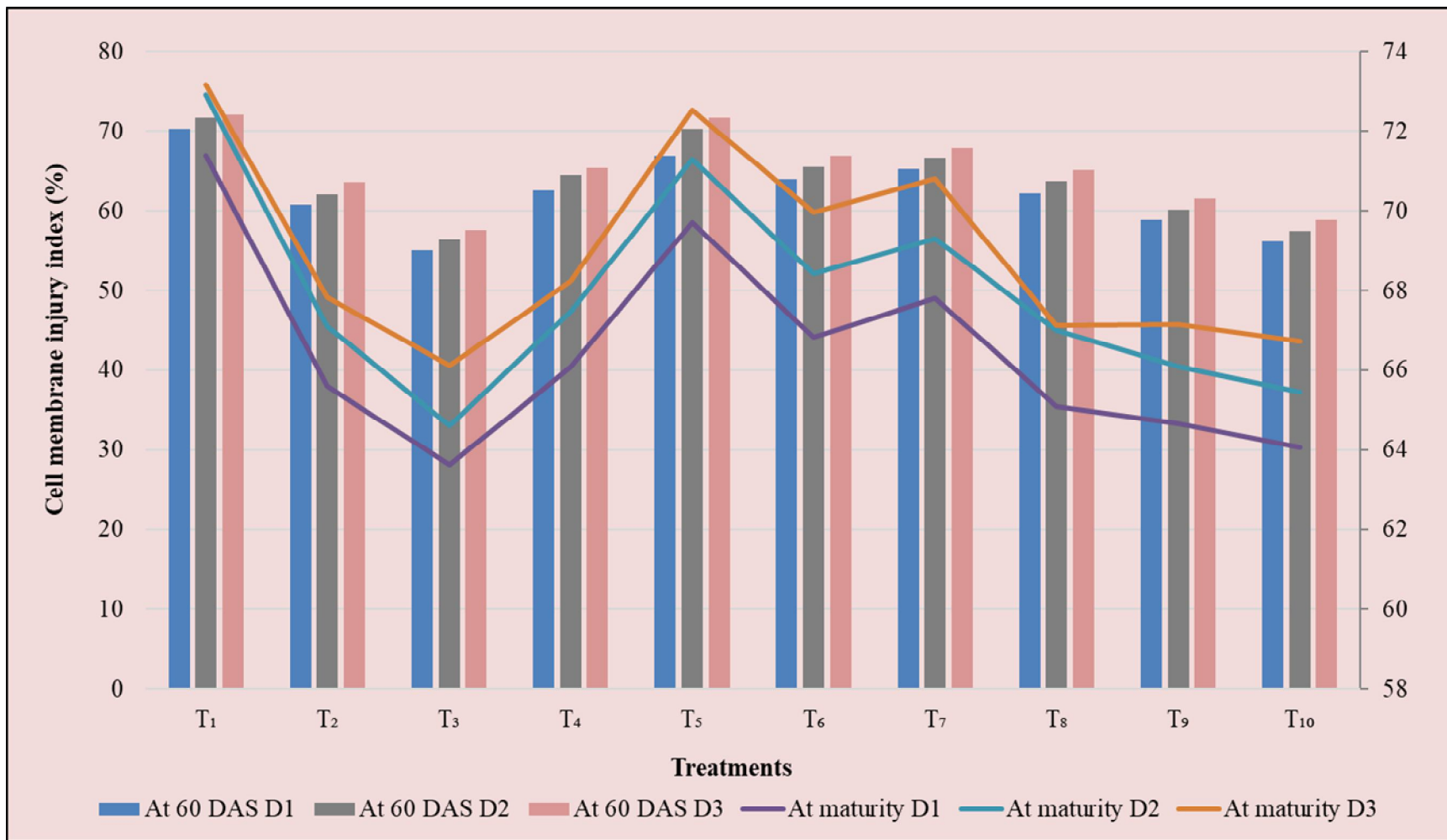


Fig. 2. Effect of dates of sowing and heat stress mitigating chemicals on cell membrane injury index



Fig. 3. Effect of dates of sowing and heat stress mitigating chemicals on proline content in chickpea

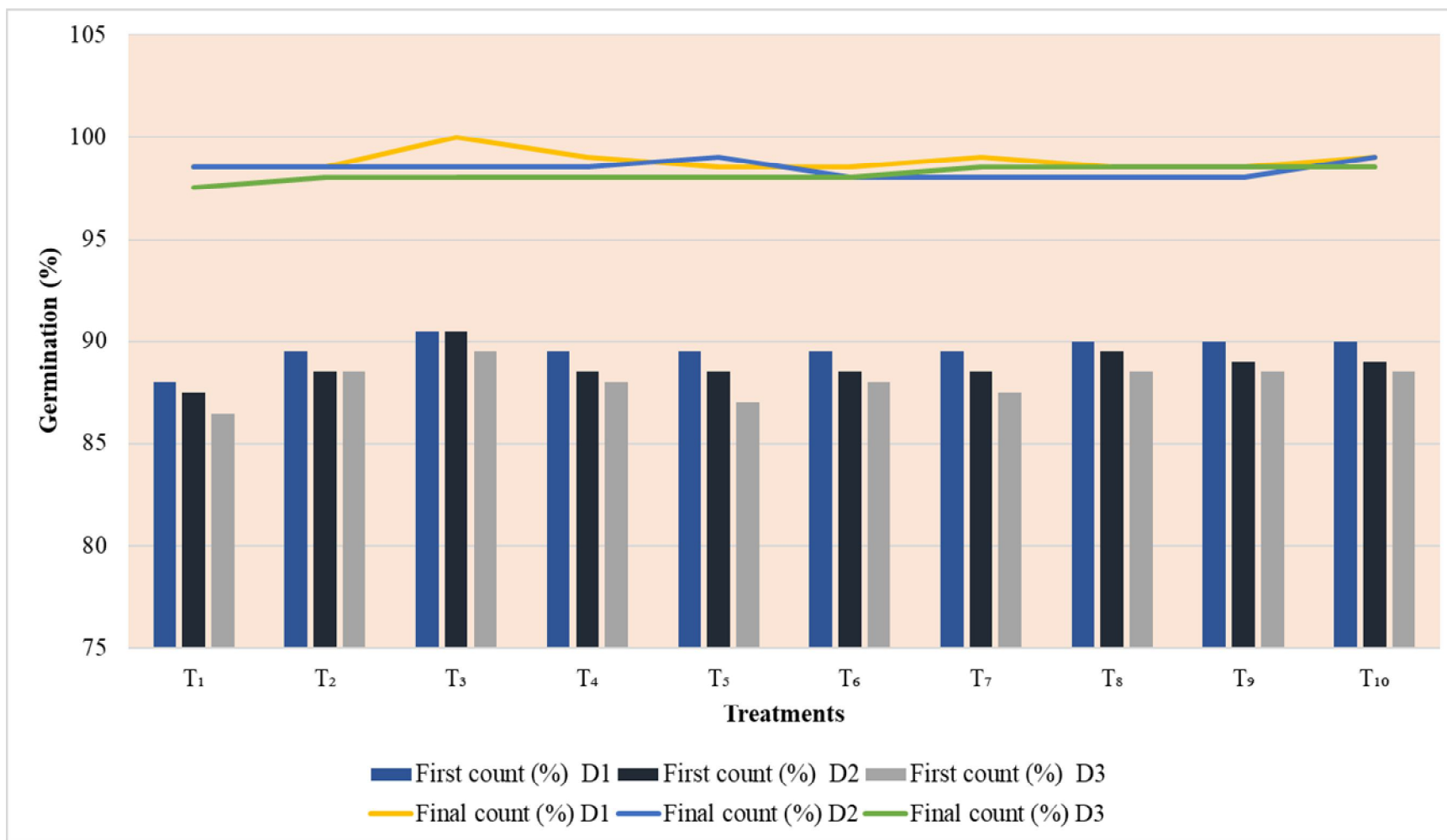


Fig. 4. Effect of dates of sowing and heat stress mitigating chemicalson germination (%) in chickpea

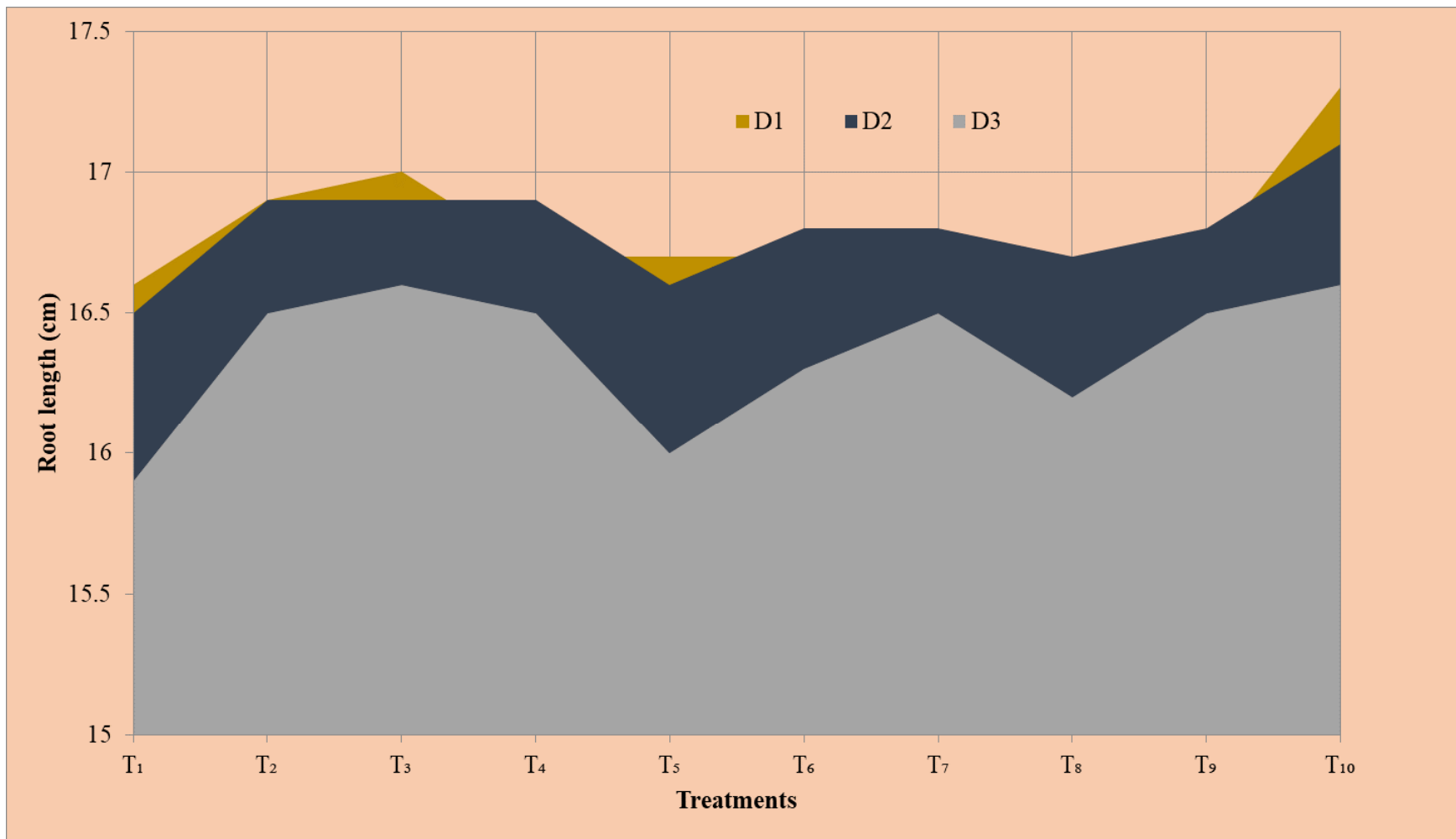


Fig. 5. Effect of dates of sowing and heat stress mitigating chemicals on root length in chickpea

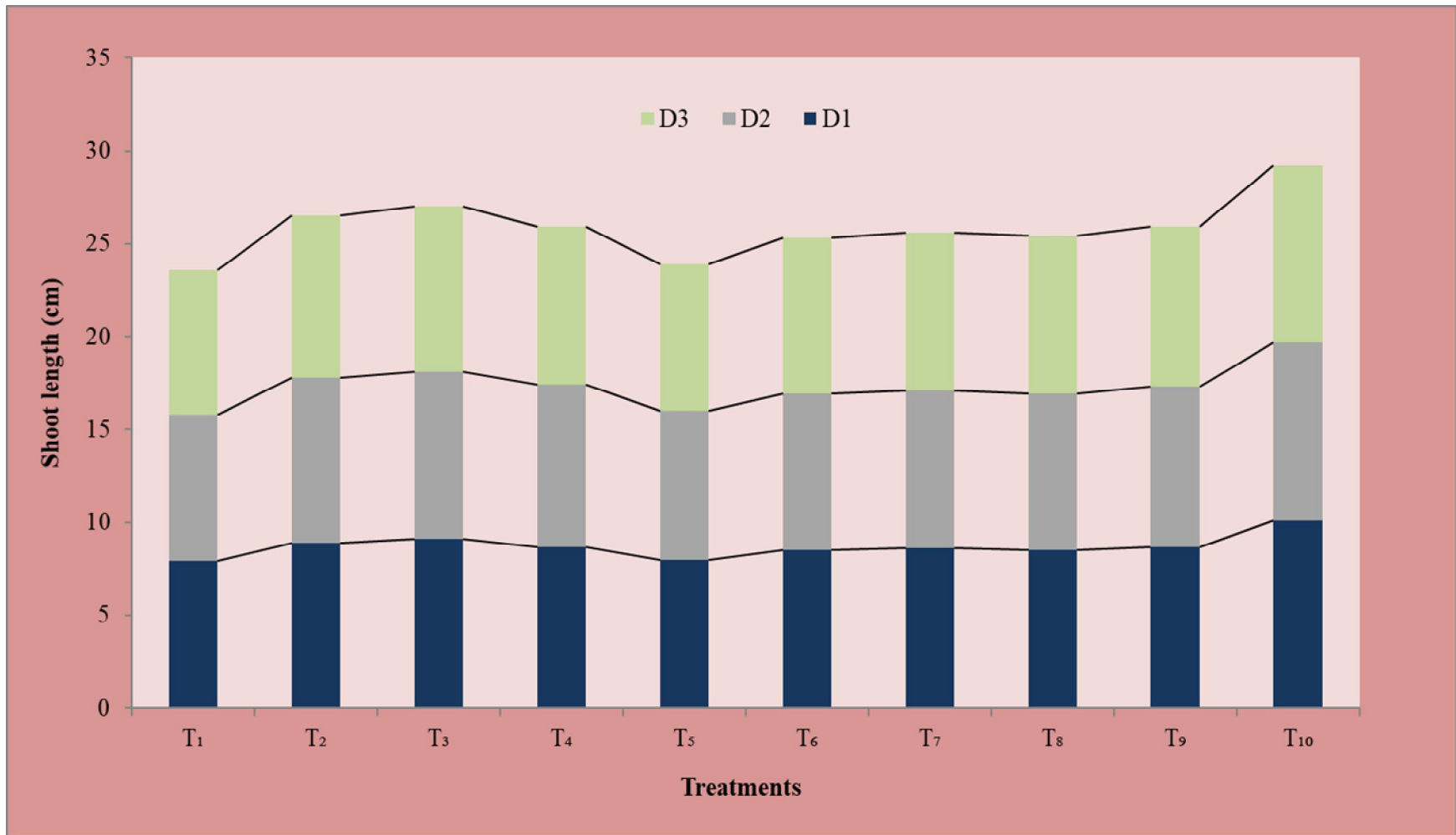


Fig. 6. Effect of dates of sowing and heat stress mitigating chemicalson shoot length in chickpea

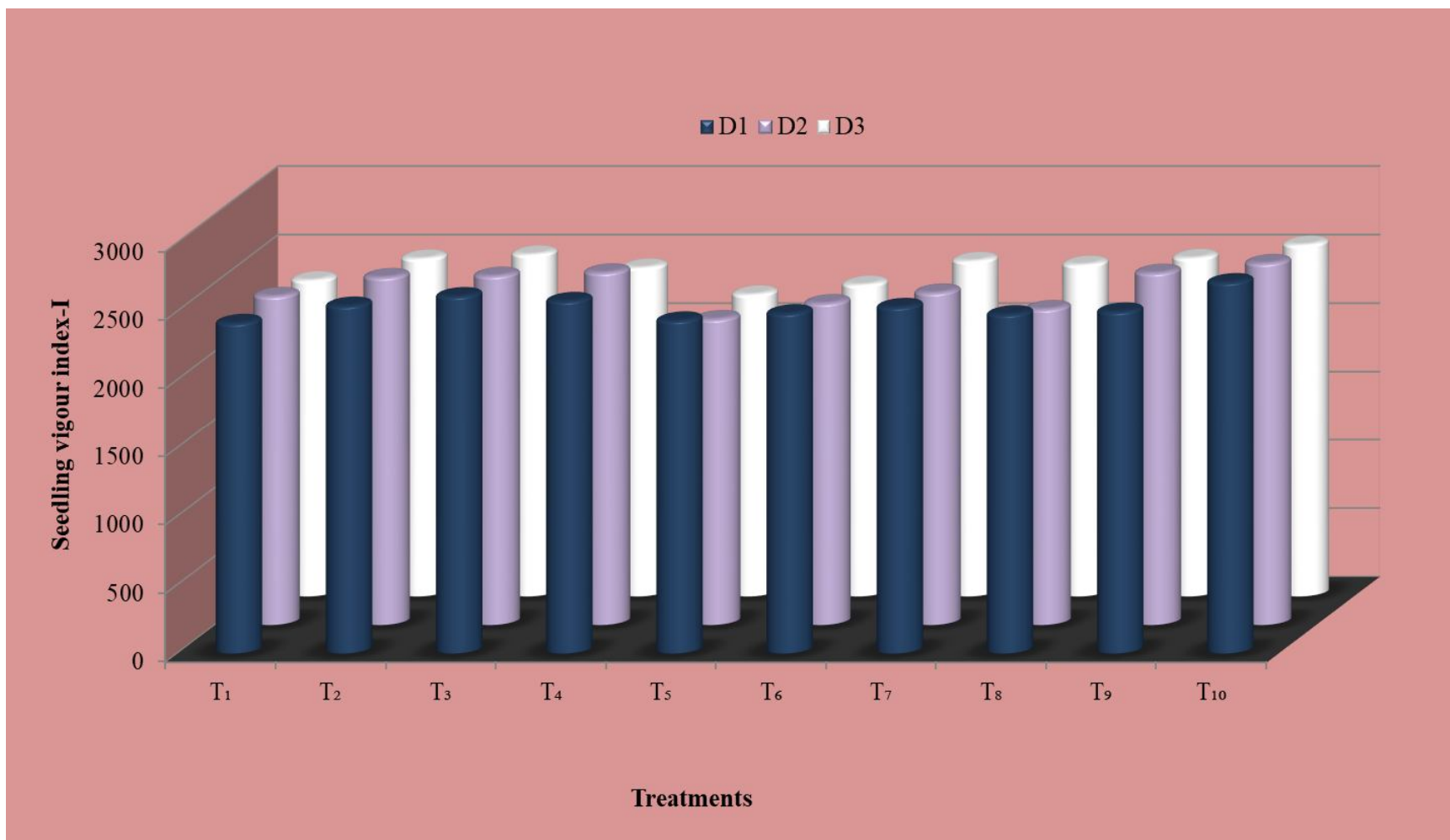


Fig. 7. Effect of dates of sowing and heat stress mitigating chemicals on seedling vigour index-I in chickpea

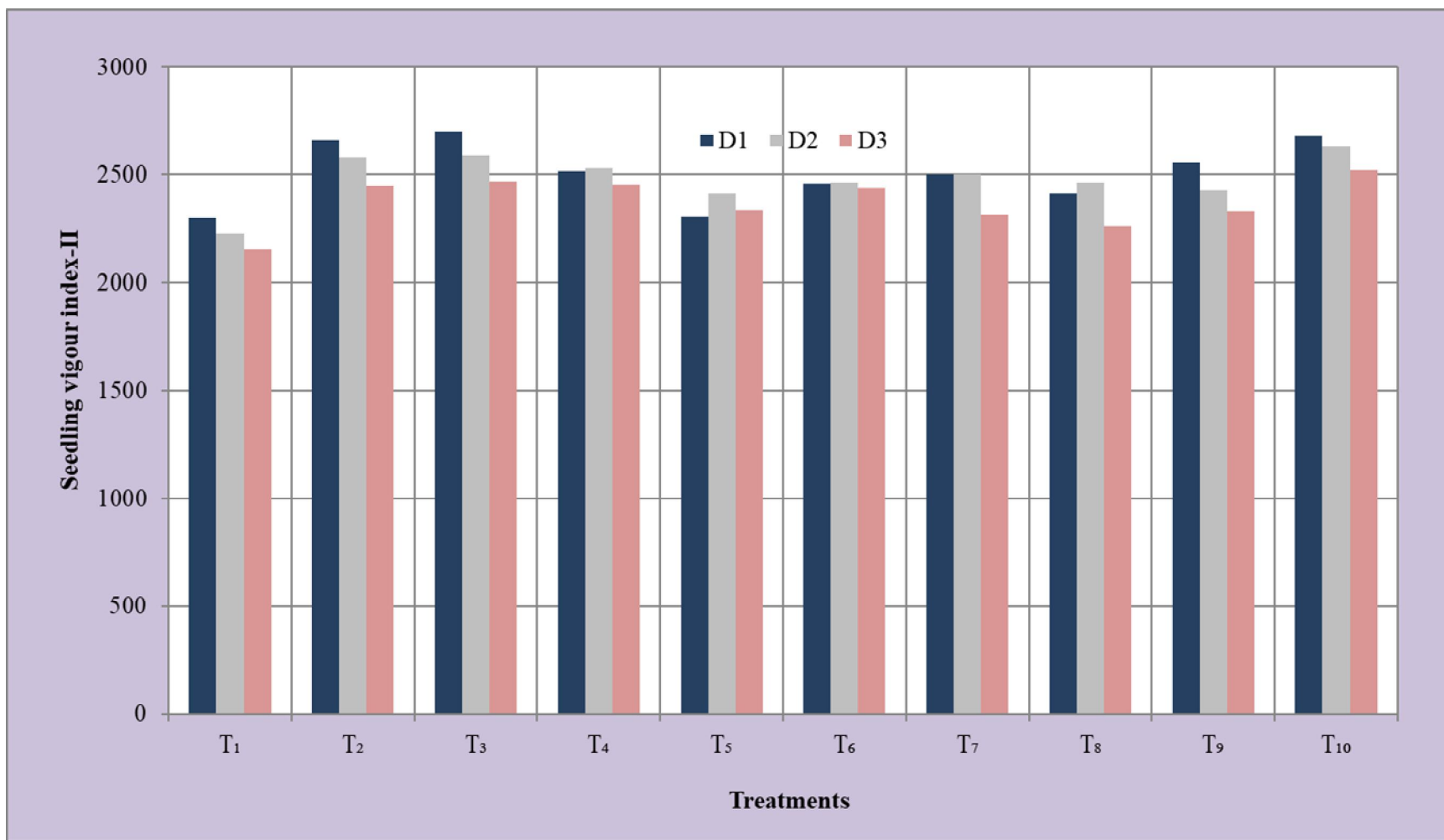


Fig. 8. Effect of dates of sowing and heat stress mitigating chemical on seedling vigour index-II in chickpea

Normal sowing (D_1)

Late sowing (D_2)

Very late sowing (D_3)



Plate 1: General view of experimental plot



At 60 DAS



At maturity

Normal sowing



At 60 DAS



At maturity

Very late sowing

Plate 2: Effect of dates of sowing and heat stress mitigating chemicals on proline content





D₁T₃: Normal sowing + Salicylic acid @ 400 ppm



D₁T₁: Normal sowing + Control



D₃T₃: Very late sowing + Salicylic acid @ 400 ppm



D₃T₁: Very late sowing + Control

Plate 3: Effect of dates of sowing and heat stress mitigating chemicals on seed germination

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D₁T₁₀: Normal sowing + Gibberellic acid @ 100ppm



D₁T₁: Normal sowing + Control



D₃T₃: Very late sowing + Gibberellic acid @ 100ppm



D₃T₁: Very late sowing + Control

Plate 4: Effect of dates of sowing and heat stress mitigating chemicals on seedling shoot and root length



Crushed leaf sample (0.5g)



Homogenate
filtration



2 ml of glacial acetic acid
2 ml of acid ninhydrin



100 °C for 30 min



Colour changed slightly



6 ml of toluene



Brick red colour



Absorbance at 520nm



Plate 5. Procedure for estimation of proline content

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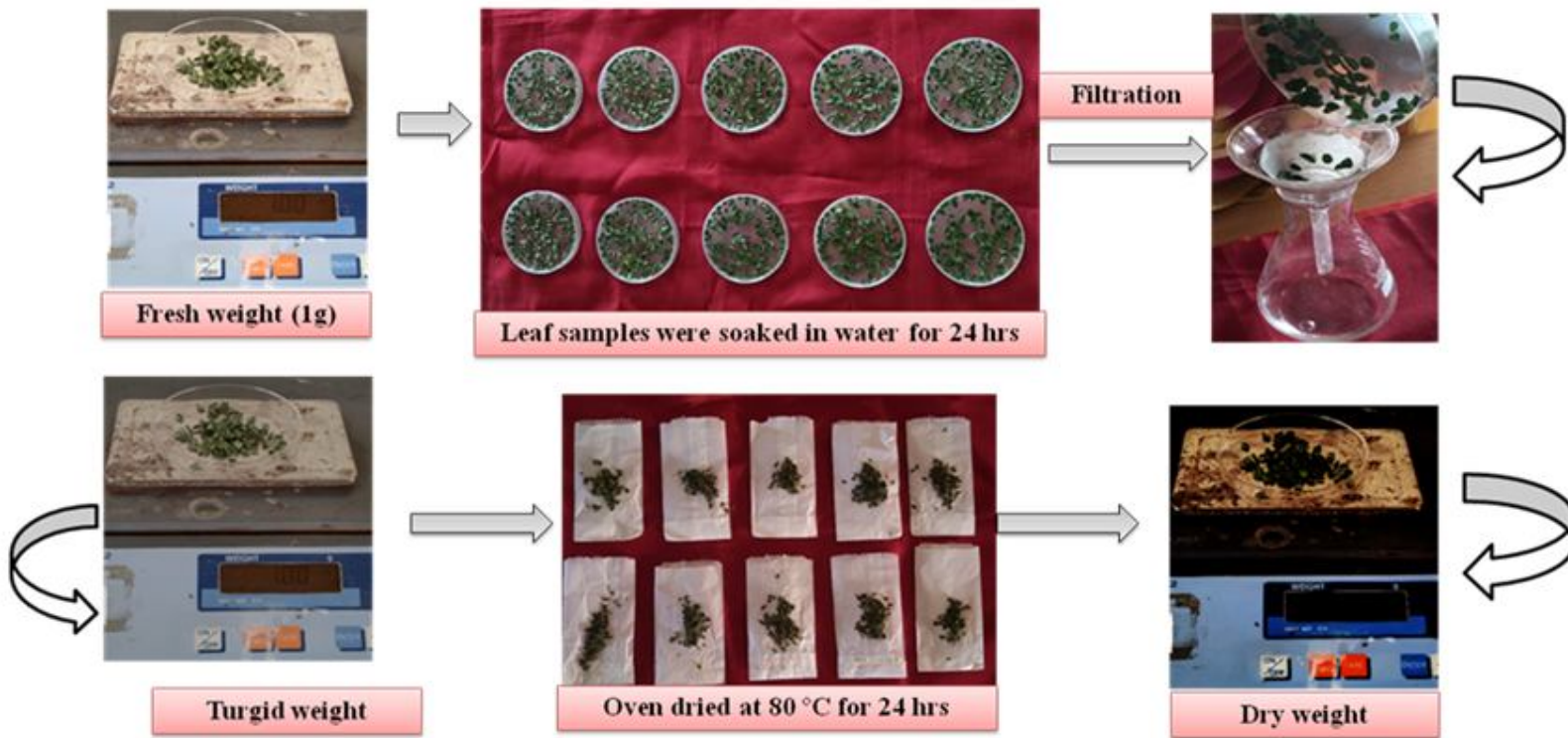


Plate 6. Procedure for estimation of relative water content

Plate 6. Procedure for estimation of relative water content

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