

Induction of Salt Tolerance in Tomato Through Seed Priming

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Abstract: A pot experiment was conducted in the Net House of “Field Laboratory of Plant Stress Management” in the Horticulture farm of Sher-e-Bangla Agricultural University, Dhaka, during the period from October 2016 to March 2017. The two factors experiment was laid out in Complete Randomized Design with five replications. Factor A is three tomato varieties viz. V_1 = Exotic line 1 (Korean), V_2 = Exotic line 2 (Taiwan) and V_3 = BARI tomato 14 and factor B is seed priming treatment viz. P_0 = No priming (Control), P_1 = Hydropriming (distilled water), P_2 = NaCl priming (50 mM) and P_3 = KNO_3 priming (200 mM). The total treatment combinations were (4×3) 12 and 8 dS/m fixed salinity maintained for all the pots. The experimental results exhibited that seed priming treatment significantly affected growth, yield and quality parameters of tomato. The highest plant height (137.10 cm), number of fruits per plant (40.92) and fruit yield per plant (585.00 g) were found from V_1 under 8 dS/m salinity level. In case of seed priming, the highest plant height (150.10 cm), number of fruits per plant (48.11) and fruit yield per plant (755.80 g) were recorded from P_2 mostly at 8 dS/m salinity level. Regarding the combined effect, the highest plant height (187.00 cm), number of fruits per plant (55.00) and fruit yield per plant (829.30 g) were found from V_1P_2 under 8 dS/m salinity level. So, Exotic line 1 with NaCl priming (50mM) showed better performance for growth, yield and quality of tomato under saline condition.

Keywords: Tomato, Stress, Seed Priming, Salt Tolerance

1. Introduction

Tomato (*Lycopersicon esculentum* Mill.) is important amongst the most substantial vegetables and broadly cultivated in Bangladesh. It is the second biggest vegetable cultivated after potato in Bangladesh. It is considered as a significant "defensive nourishment" as a result of its exceptional nutritive value. The whole area under tomato production in Bangladesh is around 68, 366 acres of land with a yield of 3, 88, 725 tons and a productivity of 5.69 tons/acres of land [1]. In Bangladesh, it is commonly termed as "poor man's apple" and widely grown throughout the country. It is mainly cultivated as Rabi crop in Bangladesh.

The coastal area covers about 20% of the country and over thirty percent of the net cultivable area. It extends inside up to 150 km from the coast. Out of 2.85 million hectares of the

coastal and offshore areas about 0.83 million hectares are arable lands, which cover over 30% of the total cultivable lands of Bangladesh. [2].

During the lifecycle, tomato crop come across a number of biotic and abiotic stresses which severely limit the production. Among the abiotic stresses, salinity, drought, temperature, mineral toxicity, U. V radiations are vital for yield constraints. Abiotic factors are considered to be the main cause of yield reduction up to 71% [3]. Salinity is one of the major abiotic stresses which adversely affect the crop yield. It is known to exercise depressive effects on metabolic pathway and energy generating processes in seeds under saline conditions.

Tomatoes, one of the most important and widespread crops

in the world, are classified as moderately salt tolerant [4]. But high concentration of salt causes hyper osmotic and ionic stresses which in turn generate secondary stresses such as oxidative stress, ionic imbalance and ultimately cell death. [5].

Pretreatment of seed has been used not only to increase salt tolerance during germination and early growth stages [6] but also can have effects during fruiting. Plants from tomato seeds primed in 1 M NaCl for 36 h produced a greater fruit yield at low (35 mM NaCl) and moderate (70 mM NaCl) salt levels in the irrigation water than non-primed seed [7]. The positive effect of seed priming on yield was not clear at 140 mM since the negative effect of high salinity during the growing period dominates the positive effect of seed-priming [7]. This adaptation could be due to the sum of the adaptation induced by priming the seed with salt, plus adaptation induced by salinity during radicle emergence, as fruit yield also increased in plants primed at germination [8]. The faster growth of tomato plants from primed seeds seems to be the result of a higher capacity for osmotic adjustment because plants from primed seeds have more Na⁺ and Cl⁻ in roots and more organic acids and sugars in leaves than plants from non-primed seeds [9].

A large number of studies confirmed that various seed pretreatments triggering the so-called “pre-germinative metabolism” [10] have been used to improve seeds germination [11, 12], seedling establishment and increase plant vigor and yield [12].

The advantage of seed priming in reducing the germination time and improving emergence uniformity is well established under laboratory conditions. However, a very few detailed studies have been conducted on performance of primed seeds under field conditions. Earlier works showed that the success of seed priming is influenced by the complex Combined of factors including plant species, water potentiality of the priming agent, duration of priming, temperature, seed vigour and storage conditions of primed seeds [13]. Limited attempts are made to compare the crop specific efficacy of all the popular methods of priming viz., hydro priming (soaking in water), halo priming (soaking in inorganic salt solutions), osmo priming (soaking in solutions of different organic osmotica), thermo priming (treatment of seed with low or high temperatures), solid matrix priming (treatment of seed with solid matrices) and biopriming (hydration using biological compounds) on seedling vigour and plant performance. Since, seed priming is found to be a useful technology there is a need to standardize this technology in every crop species particularly in vegetables and floriculture crops. Being low cost and simple technique, this on-farm seed priming represents good insurance for risk-averse to resource poor farmers and can be promoted as low cost, low risk technology that would be appropriate for all farmers irrespective of their socio-economic status. Hence, the present study was taken up to induce salinity tolerant in tomato through seed priming, determine the effect of seed priming on the growth, yield and quality of tomato. And find out the suitable combination of seed priming and variety for

ensuring the higher yield of tomato under saline condition.

2. Materials and Methods

2.1. Experimental Design

The soil salinity level was fixed to 8 ds/m for all the pots. The experiment was set up in a two factor completely randomized design with five replications. Thus 60 experimental pots were placed in ambient air at the net house premises of “Field Laboratory of Plant Stress Management lab.” The salinity in irrigation water was developed by adding required amounts of NaCl salt in irrigation water as per the procedure of Michael [14] and Ponnampuruma [15].

2.2. Seed Priming Procedure

A quantity of 3g seeds from each tomato variety was superficially sterilized with sodium hypochlorite solution (1%) for 3 minutes and then thoroughly washed for 5 minutes with distilled water. After that, seeds of each varieties were primed with 200 millimoles (mM) of KNO₃ solution for 8 hrs. [16].) and For NaCl priming, seeds of each varieties were primed with 50 mM of NaCl solution for 24 h. For hydropriming, seeds were soaked in distilled water for the same duration [17]. Few seeds were untreated and seed priming was done in Horticulture and Biotechnological Lab, M. A. Wajed Miah Research Laboratory, SAU.

20.22g of KNO₃ was dissolved in 1000 ml of distilled water and 2.93g of NaCl was dissolved in 1000 ml of distilled water.

2.3. Imposition of Salinity Treatments

First application of salt water in the soil was applied 30 days after seedling transplanting. After that two application of salt water was applied in 15 days interval. The developed irrigation water salinity and pot soil were measured by using an electrical conductivity meter (HANNA HI 993310, Direct Salinity Meter). which was expressed in dS/m.

2.4. Preparation and Application of Salt Solution

Saline water was synthesized by using Emplura Sodium Chloride. 4.68 g of salt was dissolved in 1-liter tap water to prepare the salt solution. The salinity level of the salt solution was 8 dS/m. Saline water was applied to the plant as irrigation. 500 ml of well-prepared saline water was applied to every pot.

2.5. SPAD Value

Leaf chlorophyll content as SPAD values were measured from the youngest fully-expanded leaf in the third position from the tip by a portable chlorophyll meter (SPAD-502, Konica Minolta, Japan). The SPAD-502 chlorophyll meter can estimate total chlorophyll amounts in the leaves of a variety of species with a high degree of accuracy and is a

nondestructive method [18]. SPAD was recorded at flowering stage and 30 days after flowering.

2.6. Measurement of Vitamin-C

Oxidation Reduction Titration method was used for determination of vitamin-c in ripen tomato juice. Extract of tomato fruit juice was used for determination of Vitamin-C content in per 100g of tomato sample. It has expressed as mg Vitamin-C per 100mg of tomato. Tomato juice was prepared

$$\text{Vitamin-C content or L-ascorbic acid content} = \frac{0.5 \times \text{mean value of unknown solution reading} \times 100}{\text{Mean value of Known solution reading} \times 5} \text{ mg of L-ascorbic acid}$$

2.7. Determination of Proline and Carotenoid Content

For proline determination, 0.1g of leaf sample was collected from each tomato variety which was developed from different seed priming treatment and chopped into small pieces. The chopped sample was then transferred to a test tube containing 15 mL of 80% ethanol. The test tubes were incubated in 60°C water bath for 30 minutes (Troll and Lindley 1955). The incubated sample was filtered using Whatman No. 1 filter paper. 2 mL of filtered extract was added to a test tube containing 2 mL of ninhydrin acid and 2 mL of glacial acetic acid. The test tubes were incubated in 100°C water bath for 1 hour. Incubated test tubes were placed in cold water for termination of reaction. 4 mL of toluene was added into each test tube and vortexed vigorously to form two distinct layers. The aqueous layer was used for proline determination and absorbance reading was recorded at $A_{520\text{nm}}$ [19] by using SP-UV 500DB Series UV-Vis Spectrophotometer and data were computed based on the proline standard curve.

For determination of carotenoid, 0.1g of ripen fruit sample was collected from each tomato variety and ground with 1g of Calcium Carbonate (CaCO_3). Total of 25 mL of 80% acetone was added to the power and mixed evenly. The mixture was filtered by Whatman No. 1 filter paper and filtrate was collected to determine carotenoid [20]. The absorbance was measured at 440nm by using SP-UV 500DB Series UV-Vis Spectrophotometer to determine carotenoid content and calculated by using following this equation:

$$\text{Carotenoid Content } (\mu\text{g/g}) = \frac{A \times V(\text{mL}) \times 10^4}{A_{1\text{cm}}^{1\%} \times P(\text{g})}$$

2.8. Measurement of Total Soluble Solids (TSS)

Refract meter (Model RHB 32 ATC) was used to measure TSS. One tomato sample was collected from each of the treatment. Tomato samples were cut with the sharp knife and inside was squeeze with the needle for sample juice. A drop of tomato fruit juice was placed on the transparent glass and it was covered by the upper glass. Brix refract meter was directly showed the TSS as percentage.

by blender and the volume was made with meta phosphoric acid up to 100 ml. 5 ml of standard L-ascorbic solution was taken in a conical flask. Then it was titrated with 2, 6 dichlorophenol indophenol taken in a burette. The end point was reached when the pink color lasts 10 seconds. Similarly, 5 ml of tomato juice was titrated with dye. It was measured in "Biotechnological and Horticultural Stress Management Lab," M. A. Wajed Miah Research Centre, SAU.

Calculation:

2.9. Analysis of Data

The data in respect of growth, yield contributing characters and growth & fruit quality parameters were statistically analyzed to find out the statistical significance of the experimental results. The analyses were done following the software SPSS. The significance of the difference among the means was evaluated by the Least Significant Difference Test (LSD) at 5% level of probability.

3. Result and Discussion

The experimental results obtained in this study are presented and discussed under following heads.

3.1. Plant Height

The data on plant height of tomato plant as influenced by different seed priming treatments was presented in the Table 1. Significant variation in the plant height was noticed at 90 DAT among different varieties and seed priming treatments under saline condition. As influenced by different priming treatments a mean plant height (cm) has shown in the (Table 1.) at 90 DAT. It was highest (150.10cm) from NaCl seed priming (50mM) treatment and lowest plant height (82.44 cm) was recorded from control under saline condition (Table 1).

Significant variation was observed by the varietal effect on plant height of tomato (Table 2). Exotic line 1 (Korean) tomato plant showed highest plant height (137.10 cm) due to seed priming under saline condition. Lowest plant height (98.75 cm) was recorded in BARI tomato 14 (Table 2).

The Combined effect of seed priming and varieties showed significant variation on plant height. The experiment findings noticed the variation among all the treatments (Table 3). The highest plant height (187.00 cm) was recorded from V_1P_2 where the lowest plant height (73.67 cm) was achieved from V_3P_0 which was statistically similar with V_2P_3 , V_2P_0 and V_1P_0 .

The increased plant height influenced by seed priming in the present study might be due to rapid cell division in meristematic region, number of cells and increase in cell elongation due to multiplication of various parts of the plant tissue, auxin metabolism, cell wall plasticity and permeability of cell membrane, increasing photosynthates [21]. A progressively increased enhancement with osmo-priming [22].

Table 1. Effect of priming treatments on yield contributing parameters of tomato under saline condition.

Treatments	Plant height (cm)	Number of branches per plant	Leaf area (cm ²)	Number of flower cluster per plant	Number of fruits per plant	Fruit yield per plant (g)
P ₀	82.44 c	8.67 c	158.40 d	9.78 c	22.22 c	373.10 d
P ₁	119.40 b	13.67 b	221.00 c	15.11 b	34.44 b	478.00 c
P ₂	150.10 a	18.56 a	332.60 a	20.44 a	48.11 a	755.80 a
P ₃	118.10 b	15.00 b	258.10 b	16.56 b	34.00 b	562.10 b
CV%	6.83	17.79	7.68	13.10	17.09	18.88

In a column means similar letter (s) are statistically similar and those having dissimilar letter (s) differ significantly.

3.2. Number of Branches Per Plant

Number of branches per plant of tomato was significantly affected by the different seed priming treatment under saline condition at 90 DAT (Table 1). At 90 DAT, where the highest Number of branches per plant (18.56) was found from P₂ and the lowest value (8.67) was found from P₀.

A significant effect of varieties under salinity stress was found on the number of branches per plant of tomato at 90 DAT (Table 2). At 90 DAT, the highest number of branches per plant (17.17) was observed from V₁ and the lowest value (10.58) was found from V₃.

The Combined effect of seed priming and variety on number of branches per plant of tomato exhibited a significant effect at 90 DAT (Table 3). At 90 DAT, the highest number of branches per plant (22.00) was found from V₁P₂ which was statistically similar with V₁P₃ (19.33) V₃P₃ (18.67) and V₃P₁ (16.67). The lowest value (6.67) was found from V₃P₀ which was statistically similar with V₂P₀ (8.00) and V₂P₁ (10.00). The positive influence of seed priming in improving the productive branches and there by increased yield.

Table 2. Effect of different varieties on yield contributing parameters of tomato under saline condition.

Treatments	Plant height (cm)	Number of branches per plant	Leaf area (cm ²)	Number of flower cluster per plant	Number of fruits per plant	Fruit yield per plant (g)
V ₁	137.10 a	17.17 a	289.00 a	15.83 a	40.92 a	585.00 a
V ₂	116.80 b	14.17 b	248.10 b	15.58 a	32.33 b	554.30 b
V ₃	98.75 c	10.58 c	190.40 c	15.00 a	30.83 b	487.40 c
CV%	6.83	17.79	7.68	13.10	17.09	3.58
LSD (0.05)	6.77	2.68	15.68	1.71	5.00	16.35

In a column means similar letter (s) are statistically similar and those having dissimilar letter (s) differ significantly

3.3. Leaf Area

Statistically significant variation was recorded for leaf area due to seed priming treatment at 30 days after flowering. At flowering stage, the maximum leaf area (332.60 cm²) was recorded from P₂ while the minimum leaf area (158.40 cm²) was found from P₀ (Table 1). Seed priming treatment effects were registered on leaf area of tomato. Different varieties varied significantly on leaf area of tomato at flowering stage and 30 days after flowering. At flowering stage, the maximum leaf area (289.00 cm²) was obtained from V₁ whereas the minimum leaf area (190.40 cm²) was found from V₃. (Table 2).

Leaf area of tomato showed significant differences due to interaction effect of different seed priming treatment and variety at flowering stage and 30 days after flowering. At flowering stage, the maximum leaf area (437.10 cm²) was attained from V₁P₂ treatment combination and the minimum (139.70 cm²) from V₃P₀ treatment combination which was significantly similar to V₂P₀ (152.30 cm²) (Table 3).

3.4. Number of Flower Cluster Per Plant

Different seed priming treatment under salt stress significantly in terms of number of flower cluster per plant of tomato. Data revealed that the highest number of flower cluster per plant (20.44) was found from P₂ while the lowest

number (9.78) was recorded from P₀ (Table 1).

Varieties showed no significant differences on number of flower cluster per plant of tomato. The highest number of flower cluster per plant (15.83) was recorded from V₁ which was closely followed (15.58 and 15.00) by V₂ and V₃ respectively (Table 2).

Combined effect of different seed priming treatment and variety showed significant differences on number of flower cluster per plant. The highest number of flower cluster per plant (23.00) was observed from V₁P₂ treatment combination while the lowest number (9.00) was attained from V₃P₀ treatment combination (Table 3).

3.5. Number of Fruits Per Plant

Significant variation was recorded in terms of number of fruits per plant of tomato due to different seed priming treatment under salt stress. The highest number of fruits per plant (48.11) was recorded from P₂ whereas the lowest number (22.22) was found from P₀ (Table 1).

Number of fruits per plant of tomato showed statistically significant difference due to different varieties. The highest number of fruits per plant (40.92) was recorded from V₁ and the lowest number (30.83) was recorded from V₃ which was statistically similar to V₂ (32.33) (Table 2).

Combined effect of different seed priming treatment and

varieties observed significant differences on number of fruits per plant. The highest number of fruits per plant (55.00) was observed from V₁P₂ treatment combination, whereas the lowest number (16.33) was attained from V₃P₀ treatment combination which was statistically similar to (20.33) V₂P₀ (Table 3). Similarly, significant positive association of this trait with yield [23, 24].

3.6. Fruit Yield Per Plant

Different seed priming treatment varied significantly in terms of yield per plant of tomato under salt stress. The highest yield per plant (755.80 g) was recorded from P₂ while

the lowest yield (373.10 g) was found from P₀ (Table 1).

Different tomato varieties showed significant differences on yield per plant of tomato. The highest yield per plant (585.00 g) was recorded from V₁ whereas the lowest yield (487.40 g) was observed from V₃ (Table 2).

Yield per plant varied significantly due to the combined effect of different seed priming treatment and different tomato varieties under salt stress. The highest yield per plant (523.70 g) was recorded from V₁P₂ treatment combination and the lowest yield (487.30 g) was observed from V₁P₃ treatment combination (Table 3).

Table 3. Combined effect of different priming treatments and different varieties of tomato on plant yield contributing parameters under saline condition.

Treatment Combination	Plant height (cm)	Number of branches per plant	Leaf area (cm ²)	Number of flowers cluster per plant	Number of fruits per plant	Total fruit weight per plant (gm)
V ₁ P ₀	83.33 fg	11.33 cd	285.40 c	10.00 e	30.00 de	377.00 f
V ₁ P ₁	141.70 b	14.33 bcd	292.70 c	14.67 d	41.00 bc	523.70 d
V ₁ P ₂	187.00 a	22.00 a	437.10 a	23.00 a	55.00 a	829.30 a
V ₁ P ₃	136.30 b	19.33 ab	353.10 b	14.67 d	37.67 bed	487.30 e
V ₂ P ₀	90.33 ef	8.00 de	152.30 gh	10.33 e	20.33 ef	357.00 f
V ₂ P ₁	116.00 cd	10.00 de	194.20 ef	16.00 cd	33.00 cd	388.00 f
V ₂ P ₂	145.30 b	14.33 bcd	183.10 fg	17.00 cd	44.00 b	651.70 c
V ₂ P ₃	115.30 cd	15.00 bcd	250.30 d	16.67 cd	32.00 cd	553.00 d
V ₃ P ₀	73.67 g	6.67 e	139.70 h	9.00 e	16.33 f	385.30 f
V ₃ P ₁	100.70 e	16.67 abc	196.10 ef	14.67 d	29.33 de	522.30 d
V ₃ P ₂	118.00 c	11.33 cd	207.60 ef	21.33 ab	45.33 ab	786.30 b
V ₃ P ₃	102.70 de	18.67 ab	218.40 e	18.33 bc	32.33 cd	646.00 c
CV%	6.83	17.79	7.68	13.10	17.09	25.70
LSD (0.05)	13.53	5.37	21.37	3.42	9.99	3.58

In a column means similar letter (s) are statistically similar and those having dissimilar letter (s) differ significantly.

3.7. Chlorophyll and Vitamin-C Content

Significant variation was observed for Chlorophyll content values of tomato plant due to different seed priming treatment under salt stress. At flowering stage, the highest SPAD values (47.92) was obtained from P₂ whereas the lowest SPAD values (22.56) was found from P₀ (Table 4)

SPAD values of tomato at flowering stage varied significantly due to different tomato varieties under salt stress. At flowering stage, the highest SPAD value (41.13) was found from V₁ which was statistically similar (39.97) with V₂, while the lowest SPAD value (21.36) was recorded from V₃. (Table 5).

Combined effect of different seed priming treatment and different tomato variety showed significant differences in terms of chlorophyll content of tomato at flowering stage under salt stress. At flowering stage, the highest SPAD value (51.60) was observed from V₁P₂ treatment combination and the lowest SPAD values (29.87) from V₂P₃ treatment combination. (Table 6). The results were in agreement with results reported by Hajer *et al.*, [25], that chlorophyll content decreased with increasing water salinity and seed priming tend to protect plants from damage caused by salinity.

It was observed from the result of present experiment that different seed priming treatment significantly varied the vitamin-c in ripen tomato fruit under saline condition. The maximum Vitamin-C content (18.06 mg/100g) was found from P₂ while the minimum content of Vitamin-C (10.26 mg/100g) was achieved from P₀. (Table 4)

Vitamin-C content in ripen tomato varied significantly with the different tomato varieties under saline condition. The highest value of Vitamin-C content in ripen fruit was found from V₁ (14.71 mg/100g) and lowest from V₃ (9.23 mg/100g). (Table 5).

Combined effect of the seed priming treatment and different tomato varieties under saline condition varied significantly for the content of Vitamin-C of ripen tomato fruit. The maximum amount of Vitamin-C content (24.78 mg/100g) was attained from V₁P₂ whereas the minimum amount of Vitamin-C content (7.45 mg/100g) was found from V₃P₀ (Table 6).

3.8. Carotenoid Content

It was noticed that from the result of the experiment, different seed priming treatment significantly varied the carotenoid in ripen tomato fruit under saline condition. The maximum carotenoid content (5.13 mg/100g) was found from P₂ while the minimum content of carotenoid (2.12 mg/100g) was achieved from P₀. (Table 4)

Significant effect of different tomato varieties observed in ripen tomato fruit under salt stress. The highest value of carotenoid content (3.98 mg/100g) was found from V₁ and lowest value (2.10mg/100g) was recorded from V₃ (Table 5). The Combined effect between different seed priming treatment and different tomato varieties on carotenoid content of tomato plant was statistically significant. The highest carotenoid content (5.88 mg/100g) was found from V₁P₂. The lowest value (2.20 mg/100g) was found from V₃P₀ (Table 6).

Table 4. Effect of different priming treatments on fruit quality parameters of tomato under saline condition.

Treatments	Chlorophyll content of leaves (%)	Vitamin-C (mg per 100gm)	Total soluble solid (brix%)	Carotenoid (mg per 100g)	Proline (mg/g)
P ₀	22.56 c	10.26 c	2.21 d	2.12 c	2.11 d
P ₁	38.06 b	14.13 b	5.22 c	4.07 b	3.34 c
P ₂	47.92 a	18.06 a	6.82 a	5.13 a	5.45 a
P ₃	40.92 b	14.83 b	5.94 b	4.12 b	3.85 b
CV%	11.98	5.49	11.28	8.83	8.55

In a column means similar letter (s) are statistically similar and those having dissimilar letter (s) differ significantly.

3.9. Proline Content

This experiment exhibited distinct variation in proline content in leaves of tomato under salt stress at different seed priming treatment. Result on changes in proline content have been presented in (Table 4). Among the different seed priming treatment proline content was highest in P₂ (5.45 mg/g) and lowest in P₀ (2.11 mg/g) (Table 4).

Variation was noted due to the differences in different tomato varieties under salt stress. Salinity stress creates a maximum demand for proline during stress conditions in

tolerant varieties to withstand against salinity stress than sensitive cultivars. The maximum proline content found in V₁ (3.87 mg/g) and lowest value from V₃ (2.09 mg/g). (Table 5)

The Combined effect of different seed priming treatment and different tomato varieties under salt stress varied significantly on proline content in tomato leaves. Maximum proline accumulation was seen from V₁P₂ (6.88 mg/g) under salt stress and minimum proline content found from V₂P₃ (2.15 mg/g) which was statistically similar to V₁P₀ (2.50 mg/g) and V₂P₀ (2.31 mg/g) (Table 6).

Table 5. Effect of different varieties on fruit quality parameters of tomato under saline condition.

Treatments	Chlorophyll content of leaves (%)	Vitamin-C (mg per 100gm)	Total soluble solid (brix%)	Carotenoid (mg per 100g)	Proline mg/g
V ₁	41.13 a	14.71 a	4.92 a	3.98 a	3.87 a
V ₂	39.97 a	10.58 b	5.42 a	3.37 b	3.04 b
V ₃	21.36 b	9.23 c	2.18 b	2.10 c	2.09 c
CV%	11.44	5.49	11.28	8.83	8.55
LSD (0.05)	3.98	0.67	0.52	0.29	0.27

In a column means similar letter (s) are statistically similar and those having dissimilar letter (s) differ significantly.

3.10. Total Soluble Solids

Seed priming treatment had significant differences in total soluble solids under salt stress condition. Soluble sugar in ripe tomato fruit increased with NaCl seeds priming treatment. The data showing the changes in soluble solids content at different seed priming treatment. Maximum soluble solids found (6.82%) when seed was treated with 50mM NaCl and lowest value was found (2.21%) from control (P₀). (Table 4)

Total soluble solid (TSS) in tomato fruit varied significantly with the different variety. It was noticed that

highest TSS (4.92%) was found from V₁ which was statistically similar to V₂ (5.42%) under saline condition and the lowest TSS (1.18% from V₃). (Table 5)

Combined effect of different seed priming treatment and different tomato varieties varied nonsignificant on TSS of ripen tomato fruit under salt stress. It was observed that highest value of TSS (7.80%) was found from V₁P₂ which was statistically similar to V₁P₃ (7.50%) while the lowest value of TSS (2.36%) from V₃P₀ (Table 6).

Table 6. Combined effect of priming treatments and different varieties of tomato on fruit quality parameters under saline condition.

Treatment Combination	Chlorophyll content of leaves (%)	Vitamin-C (mg per 100gm)	Total soluble solid (brix%)	Carotenoid (mg per 100g)	Proline mg/g
V ₁ P ₀	42.93 bcd	15.97 cd	5.33 d	3.65 e	2.50 f
V ₁ P ₁	43.53 bcd	15.00 d	6.67 bc	4.45 c	3.05 e
V ₁ P ₂	51.60 a	24.78 a	7.80 a	5.88 a	6.88 a
V ₁ P ₃	46.63 abc	12.40 e	7.50 ab	5.24 b	5.09 b
V ₂ P ₀	32.80 ef	8.61 f	5.00 de	2.23 f	2.31 f
V ₂ P ₁	40.37 cde	18.58 b	6.00 cd	4.18 cde	4.79 bc
V ₂ P ₂	41.57 cd	11.37 e	5.00 de	4.09 cde	3.82 d
V ₂ P ₃	29.87 f	9.82 f	4.00 ef	1.73 f	2.15 f
V ₃ P ₀	22.71 g	7.45 g	2.36 g	1.20 g	1.18 g
V ₃ P ₁	49.87 ab	17.00 c	3.67 f	4.28 cd	4.38 c
V ₃ P ₂	41.53 cd	16.03 cd	5.67 cd	3.82 de	3.14 e
V ₃ P ₃	38.63 de	9.92 f	5.00 de	4.37 cd	4.25 c
CV%	11.44	5.49	11.28	8.83	8.55
LSD (0.05)	7.96	1.33	1.05	0.58	0.54

In a column means similar letter (s) are statistically similar and those having dissimilar letter (s) differ significantly.

4. Conclusion

From the above-mentioned results considering, it may be concluded most of the parameters of tomato under saline condition showed positive relation with seed priming. Among the seed priming treatments, NaCl priming (50 mM) showed the best result than other priming treatment. Exotic line 1 (Korean) tomato showed the highest result in growth, fruit yield and quality parameters under saline condition. Therefore, further study with more seed priming and more crop varieties may provide more conclusive and precise result.

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