

## Seed Enhancement Technique to Alleviate the Effect of Salinity Stress in Okra (*Abelmoschus esculentus*)

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**ABSTRACT:** Study was conducted to evaluate the effect of seed enhancement technique on seed quality parameters under salt stress condition. Salt stress is the major abiotic stress which cause the reduction in the growth and yield of agriculture crops. Salt stress can be induced due to in appropriate use of fertilizers, low precipitation, lack of proper irrigation & drainage system and irrigating with salt water. In this laboratory experiment, factorial combination of 3 different salt stress conditions induced by using NaCl concentration (0, 100mM and 150 mM) viz S<sub>0</sub>, S<sub>1</sub> and S<sub>2</sub>. Okra seed var. *Arka Anamika* were primed with treatments consists of Panchagavya 3% (T<sub>1</sub>) and 5% (T<sub>2</sub>), Coconut water 8% (T<sub>3</sub>) and 12% (T<sub>4</sub>), Neem leaf extract 3% (T<sub>5</sub>) and 5% (T<sub>6</sub>), KNO<sub>3</sub> (3g/litre) (T<sub>7</sub>), KCL (3g/litre) (T<sub>8</sub>) for 8 hours, with one untreated dry seed serve as control (T<sub>0</sub>) were laid out in CRD with 4 replications. Data collected on germination percentage, root length, shoot length, seedling length, fresh weight, dry weight, seed vigour I, seed vigour II and root-shoot ratio were evaluated. Results revealed that mean values of all characters varied significantly along the salinity gradient. The mean values of all the seed quality parameters are maximum in S<sub>0</sub>(control) And were minimum in S<sub>2</sub>(150mM NaCl) were clearly shows the impact of salt stress is the reason for the reduction. Seed primed with KNO<sub>3</sub> (3g/litre) performed superior in all seed quality parameters under salt stress conditions. So, KNO<sub>3</sub> (3g/litre) is the best seed enhancement technique for improving okra seed quality parameters under saline stress condition.

**Keywords:** Germination, okra, potassium priming, salt-stress, seed vigour, stress adaptability, seed enhancement, seed quality parameters.

### INTRODUCTION

Salinity stress or salt stress is the major abiotic stress in arid, semiarid and coastal regions. Salinization occurs mostly in irrigated lands due to lack of proper irrigation and drainage management, low precipitation, high evaporation and saline water irrigation (Munns and Tester, 2008). In worldwide about 34 million hectares of irrigated land are affected by salt. Thus, 50% of cultivable lands will be lost by the middle of the 21st century (Wang *et al.*, 2003). Abiotic stress causes the delay or inhibit the Seed germination (Jamil *et al.*, 2005; Fazlali *et al.*, 2013). In this respect, Jamil *et al.*, (2005), Patade *et al.*, (2011) and Rouhi *et al.*, (2011) stated that increase in the salt stress will eventually reduce the germination percentage and increase germination time.

Okra (*Abelmoschus esculentus* L.) is commonly known as Bhindi or lady's finger. It belongs to the family Malvaceae and is one of the most widely grown vegetables from tropical to subtropical and warmer parts of the temperate zone of the country. Its Sanjay *et al.*,

chromosome number is  $2n = 130$ . It is an annual vegetable crop in tropical and subtropical parts of the world. It is one of the most important nutritious vegetable crops grown around the year in India. Okra is an important fruit vegetable of high commercial and food values. It is primarily valued for its tender, immature green pods in fresh form; however, its curry, soups, and edible young leaves are also popular. To a limited extent, it finds use in canned, dehydrated, or frozen forms for off-season consumption by the army at high altitudes and export (Sharma *et al.*, 2014). The major problem in okra cultivation is when okra seeds are sown in the early spring season, the seeds don't germinate properly as the temperature is below the optimum range of 25-35°C also the hard seed coat of okra seeds restricts the imbibition of water and uniform growth and embryo development which reduces the uniform field stand and rapid germination. Young seedlings which germinated are extremely sensitive to salt stress, especially when they accumulate soluble salts from surface soils as a consequences of

evaporation and capillary rise of water (Mahmoudi *et al.*, 2012). Seed priming prevents seeds from entering stage of hydration (growth) by extending and holding seeds within activation phase when seeds are under dry stage (Heydecker and Coolbear, 1977). Seed priming is an efficient method for synchronization of germination and increasing seed vigor as well as allowing the growth of seedlings under stressful conditions (Bajehbaj, 2010). Seed priming increases the effect of salinity stress by promoting  $K^+$  and  $Ca^{2+}$  accumulation on germination and seedling growth and decreases  $Na^+$  and  $Cl^-$  accumulation in the seedlings (Afzal *et al.*, 2008). The main objective of priming treatments is to improve the seed germination and field emergence rate. Priming shows reduce in ion leakage through the membranes which could allows greater membrane integrity in the seed embryo (Merreddy *et al.*, 2000). There are a few priming methods are Hydropriming (seeds are soaked in water) osmopriming (seeds are soaked in osmotic solution), halopriming (soaking of seeds in salt solution). Even though the mechanism behind the techniques of priming the seeds are still not fully understood, but certain physiological and beneficial changes had been observed that were associated with priming of seeds (Hu *et al.*, 2006). The aim of the present study is to evaluate the effect of the seed enhancement on seed quality parameters under salinity stress in Okra var. *Arka Anamika* and to assess the best seed enhancement technique to alleviate the effect of salinity stress.

## MATERIAL AND METHODS

Experiments were carried out at Department of Seed Science and Technology, Sam Higginbottom University of Agriculture, Technology and Sciences, Prayagraj (Uttar Pradesh), 2019-2020. The study was conducted in order to evaluate the effect of priming on seed quality parameters of Okra under salt stress condition. A factorial experiment was conducted based on Completely Randomized Design with four replications. The Salt stress is induced by using NaCl at different concentration (0, 100 and 150 mM) viz  $S_0$ ,  $S_1$  and  $S_2$ . Okra seed var. *Arka Anamika* were primed with treatments consists of Panchagavya 3% ( $T_1$ ) and 5% ( $T_2$ ), Coconut water 8% ( $T_3$ ) and 12% ( $T_4$ ), Neem leaf extract 3% ( $T_5$ ) and 5% ( $T_6$ ),  $KNO_3$  (3g/litre) ( $T_7$ ), KCL

(3g/litre) ( $T_8$ ) for 8 hours, with one untreated dry seed serve as control ( $T_0$ ) were laid out in CRD with 4 replications and 100 seeds per replication. To assess the seed quality parameters, seeds need to bring back to its original moisture content by remove them from the solutions and shade dried at room temperature. The between paper method of germination test were conducted in order to assess salt stress effect on seed germination. Germination paper was pre-soaked in three different concentrations of NaCl solution ( $S_0, S_1$  &  $S_2$ ) used for conducting germination test. Seeds were germinated in a seed germinator maintained at  $25 \pm 2^\circ C$  temperature and  $95 \pm 3\%$  RH. The seedlings were evaluated at the final count of okra seed i.e., 21 days, the normal seedlings were counted and expressed in percentage (ISTA, 2011). The root and shoot length were measured by randomly selecting the ten normal seedlings from each replication and the values were expressed in centimeter. These seedlings are placed in a paper cover and dried for 24 h in the shade and then kept in an oven maintained at  $85 \pm 2^\circ C$  for 24 h. The dried seedlings weight was measured and the mean values were expressed in mg/10 seedlings. Root-shoot ratio is calculated by the ratio between the root length of the seedlings and the shoot length of the seedlings. The formulae given by Abdul-Baki and Anderson (1973) used to calculate Vigour index. The data obtained from the experiments were subjected to an analysis of variance and treatment differences tested for significance ( $P=0.05$ ).

## RESULTS AND DISCUSSION

The present study investigated the seed enhancement technique to alleviate the effect of salinity stress in okra. In general, increase in salt stress, reduce the germination percentage, root length, shoot length, seedling length, fresh weight, dry weight, seed vigour I, seed vigour II and root-shoot ratio. Seed-priming had a significant difference on germination percentage of okra seeds under salinity stress compared to control. Results revealed that mean values of all characters varied significantly along the salinity gradient. The mean values of all the parameters are maximum in  $S_0$  (control) and were minimum in  $S_2$  (150mM NaCl) (Table 1).

**Table 1: Comparison between the overall mean of different seed quality parameters under different salinity levels.**

Characters	Salinity level		
	$S_0$ (control)	$S_1$ (100mM NaCl)	$S_2$ (150mM NaCl)
Germination%	68.64	61.83	53.58
Shoot Length	20.21	17.03	12.43
Root Length	7.24	6.61	5.41
Seedling Length	27.45	23.64	17.84
Root-Shoot Ratio	0.36	0.392	0.442
Fresh Weight	5.65	4.77	3.49
Dry Weight	0.201	0.168	0.111
Seed Vigour Index I	1,904.94	1,480.38	970.49
Seed Vigour Index II	13.912	10.519	6.016

With increase in salt stress, the germination percentage of okra has significantly decreased which is similar to Ghoulam and Fares (2001) as he reported that salt stress at low level just delays the germination but higher salinity level not only reduce the germination percentage also inhibit the seedling emergence.

However, germination percentage of primed seeds have significantly increased when compared to ( $T_0$ ) control (Table 2) were similar to Hopper *et al.*, (1979) and the highest germination percentage was registered with seeds primed with in all the three salinity stress levels. In saline condition, the significant increase in seed emergence and enhancement of seedling growth with increase in  $K^+$  and  $Ca^{2+}$  concentration in different crop seed (Cramer *et al.*, 1990). The reason for improving the germination and growth of okra seed under salinity stress by  $KNO_3\ 3gl^{-1}(T_7)$  is may due to the ability of the

potassium to maintain the ionic balance in cell and respiration and carbohydrate metabolism of seed by its effect on enzymes like pyruvate kinase (Aisha *et al.*, 2007; Nawaz, *et al.*, 2011).

Similar trend was observed in shoot length, root length, seedling length, fresh weight, dry weight of okra seeds under salinity stress. Shoot length of primed seeds were greater than control and the maximum Shoot length was registered with seeds primed with  $KNO_3\ 3gl^{-1}(T_7)$  in all salinity level (Table 3).

Under saline condition plant growth and survival depends upon adaptation to high concentration of salts and a reduction in shoot length is expected (Wu *et al.*, 2015). Root length of primed seeds were greater than control and the maximum root length was registered with seeds primed with  $KNO_3\ 3gl^{-1}(T_7)$  in all salinity level (Table 2).

**Table 2: Overall treatment means for various seed quality parameter under salinity stress.**

Treatment	Salinity	Germination %	Shoot length	Root length	Seedling length	Root-Shoot ratio	Fresh weight	Dry weight	Seed Vigour index 1	Seed Vigour index 11
$T_0$	$S_0$	58.75	16.28	6.13	22.40	0.376	4.30	0.168	1316.00	9.871
	$S_1$	53.00	13.18	5.63	18.80	0.427	3.50	0.141	996.35	7.473
	$S_2$	45.25	08.95	4.58	13.53	0.511	2.38	0.093	612.05	4.209
$T_1$	$S_0$	68.25	22.25	7.10	29.35	0.319	5.75	0.198	2003.25	13.531
	$S_1$	61.25	17.83	6.50	24.33	0.365	5.00	0.167	1489.95	10.199
	$S_2$	53.50	13.03	5.30	18.33	0.407	3.70	0.111	980.45	5.872
$T_2$	$S_0$	76.00	23.15	8.08	31.23	0.349	5.78	0.223	2373.03	16.964
	$S_1$	66.75	19.13	7.15	26.28	0.374	5.00	0.182	1753.95	12.148
	$S_2$	57.75	14.15	5.85	20.00	0.414	3.80	0.12	1155.03	6.929
$T_3$	$S_0$	63.50	18.30	6.63	24.93	0.362	5.48	0.186	1582.70	11.810
	$S_1$	57.50	15.88	6.03	21.90	0.380	4.80	0.156	1259.23	8.985
	$S_2$	49.50	11.30	4.98	16.28	0.441	3.40	0.103	805.58	5.098
$T_4$	$S_0$	61.00	17.10	6.45	23.55	0.378	5.23	0.174	1436.63	10.613
	$S_1$	55.00	13.83	5.88	19.70	0.425	4.20	0.146	1083.58	8.043
	$S_2$	47.50	9.68	4.83	14.50	0.499	2.98	0.097	688.78	4.583
$T_5$	$S_0$	71.75	20.28	7.58	27.85	0.374	6.23	0.211	1998.48	15.138
	$S_1$	64.50	16.83	6.90	23.73	0.410	5.18	0.177	1530.43	11.400
	$S_2$	56.25	12.60	5.68	18.28	0.451	3.90	0.117	1028.38	6.566
$T_6$	$S_0$	65.50	19.25	6.90	26.15	0.359	5.08	0.192	1712.60	12.576
	$S_1$	58.75	15.78	6.30	22.08	0.400	4.13	0.161	1296.80	9.459
	$S_2$	51.00	11.35	5.18	16.53	0.456	3.00	0.106	842.58	5.418
$T_7$	$S_0$	79.25	24.05	8.48	32.53	0.353	6.63	0.236	2577.55	18.705
	$S_1$	71.25	20.93	7.78	28.70	0.372	5.58	0.198	2044.83	14.109
	$S_2$	62.25	15.90	6.30	22.20	0.422	4.20	0.131	1381.88	8.140
$T_8$	$S_0$	73.75	21.25	7.83	29.08	0.368	6.43	0.217	2144.25	16.003
	$S_1$	68.50	19.90	7.38	27.28	0.371	5.53	0.188	1868.35	12.859
	$S_2$	59.25	14.93	6.00	20.93	0.377	4.10	0.124	1239.70	7.330
SE(d)	(T)	58.75	0.147	0.059	0.138	0.006	0.044	0.001	12.426	0.091
	(S)	53	0.085	0.034	0.079	0.003	0.025	0.001	7.174	0.053
	(S)×(T)	45.25	0.254	0.103	0.238	0.01	0.076	0.002	21.523	0.158
SE(m)	(T)	0.262	0.104	0.042	0.097	0.004	0.031	0.001	8.787	0.064
	(S)	0.151	0.06	0.024	0.056	0.002	0.018	0.001	5.073	0.037
	(S)×(T)	0.453	0.18	0.073	0.169	0.007	0.054	0.002	15.219	0.111
CD@5%	(T)	0.737	0.292	0.118	0.274	0.012	0.087	0.002	24.77	0.181
	(S)	0.426	0.169	0.068	0.158	0.007	0.05	0.001	14.301	0.105
	(S)×(T)	1.277	0.506	0.205	0.475	0.021	0.151	0.004	42.904	0.314
F Test	S	s	S	S	S	S	S	S	S	S

This can be the result of physiological drought by the salt deposit in root cells, which could be the reason for the reduction of cell division and reduced root growth (Munns, 1993). Seedling length of okra seedling is the sum of shoot and root length, as maximum shoot length

and root length are registered in seeds primed with  $KNO_3\ 3gl^{-1}(T_7)$  under all salinity level, seedling length also maximum registered with  $KNO_3\ 3gl^{-1}(T_7)$  primed seeds in all salinity levels (Table 2). Significant reduction of fresh weight and dry weight of okra

seedling were observed under salinity levels (Table 1). However, Fresh weight & Dry weight of primed seeds have significantly increased when compared to control ( $T_0$ ) and the highest Fresh weight & Dry weight was registered with seeds primed with  $KNO_3$   $3g\ l^{-1}$  ( $T_7$ ) in all the salinity levels (Table 1). Salinity stress reduced the

Fresh weight and Dry weight of seedling. The contribution of phytohormones in the biosynthesis is decreased due to the reaction of salinity stress in seeds Cuartero *et al.*, (2006); Cicek and Cakirlar (2002) also reported that salinity reduced Fresh and Dry weight of maize seedling.

**Table 3: Analysis of variance for different seedling growth parameters in okra under salinity stress.**

Sr.No.	Character	Mean Sum of Square			
		Treatment (df=8)	Salinity (df=2)	Interaction (df=16)	Error (df=81)
1.	Germination%	481.058*	2,046.29*	3.068*	0.821
2.	Shoot Length	76.95*	550.827*	1.182*	0.129
3.	Root Length	5.888*	31.169*	0.065*	0.021
4.	Seedling Length	123.616*	843.291*	1.549*	0.114
5.	Shoot-Root Ratio	0.301*	2.318*	0.03*	0.009
6.	Fresh Weight	5.442*	42.373*	0.043*	0.011
7.	Dry Weight	0.004*	0.074*	0.001*	0
8.	Vigour Index 1	1425506*	7880625.583*	38678.515*	926.456
9.	Vigour Index 2	57.208*	564.824*	3.002*	0.05

\* Significant @5% Level of Significance.

The seed vigour I and seed vigour II of okra seedling observed in this study is significantly decrease with increase in salinity stress. Since Seed vigour I and Seed vigour II is depend on the germination percentage, seedling length and dry weight of okra seedling as seed vigour I is the product of germination percentage and seedling length and seed vigour II is the product of Germination percentage and dry weight of okra seedling. Seed vigour I and seed vigour II of okra Seeds primed with  $KNO_3$   $3g\ l^{-1}$  ( $T_7$ ) is found to be performed significantly better when compared to all other treatments and control under all salinity level. Decrease in the germination and seedling vigour due to NaCl salt stress observed in this study might be due to toxic effects of  $Na^+$  and  $Cl^-$  ions as well as salt solution create an external osmotic potential in seed that inhibit water uptake reported by (Khajeh-Hosseini *et al.*, 2003). It is also evident that the salinity disrupted the cellular expansion and differentiation which cause modifications in metabolic activity (Scialabba and Melati 1990).

In case of Root-Shoot ratio, with increase in salt stress the root-shoot ratio of okra has increase significantly. The Root-shoot ratio of primed seeds also have significant difference among the treatment with maximum root-shoot ratio was registered in control ( $T_0$ ) in salt stress levels but when we compare the Root-shoot ratio of treatments with the seedling length of the treatment it is clear that the maximum Root-shoot ratio of control is due to the lower shoot length (Table 2). When we consider the Root-shoot ratio, it is also important to analyse the shoot and root length of the seedling.

The higher Root-shoot ratio with low root and shoot length doesn't have any utility. Among the treatments with high shoot and root length, seed priming with  $KNO_3$   $3g\ l^{-1}$  ( $T_7$ ) followed by seed priming with  $KCL3g\ l^{-1}$  ( $T_8$ ) having optimum root-shoot ratio. In addition, salinity stress affects the absorption of mineral and

balance of ions in plant. Hence mineral nutrient deficiency can attribute to reduction of shoot and root length. Activation of abiotic stress defence responses that are displayed by primed plants are faster and stronger (Conrath *et al.*, 2006).

From the previous findings in Fenugreek by Soughir *et al.*, (2013) ; Sivritepe *et al.*, (2003) working on melon and Bajehbaj, (2010) working on sunflower shows that decrease in  $K^+$  and  $Ca^{2+}$  in the different part of the seedling due to NaCl salt stress causes increase in  $Na^+$  ion concentration. So, the seed priming with  $KNO_3$   $3g\ l^{-1}$  ( $T_7$ ) followed by seed priming with  $KCL3g\ l^{-1}$  ( $T_8$ ) shows better performances in terms of salinity stress. Cakmak, (2005) also suggested that under environmental stress conditions the improvement of K-nutritional status crucial for the survival of crop plants.

## CONCLUSION

Seed priming is a seed enhancement technique which is low cost, easy technique with minimal or no risk has improved seed quality parameters even under salinity stress. From this present investigation, any seed priming treatment have shown significantly better performance than control in all salinity stress levels. Among the seed priming treatment, seeds primed with  $KNO_3$   $3g\ l^{-1}$  ( $T_7$ ) have shown superior performance followed by  $KCL3g\ l^{-1}$  ( $T_8$ ) in all the seed quality parameters under all salinity stress level. Therefore, seed priming with  $KNO_3$   $3g\ l^{-1}$  ( $T_7$ ) is the best seed enhancement technique to alleviate the effect of salinity stress in okra (*Abelmoschus esculentus*). These results are obtained in a controlled atmosphere of germination chamber, so it needs to get further investigated in field trails. If the results from the field trails will give clarity to the enhancement of seed under salinity stress. After that this method can be utilized for commercial cultivation by farmers.

**Conflict of Interest.** Nil.

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