EFFECT OF HALOPRIMING ON THE INDUCTION OF NACL SALT TOLERANCE IN DIFFERENT WHEAT GENOTYPES

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Abstract

Salinity is a major environmental stress limiting plant growth and productivity of wide range of crops with impairing effects on germination and yield. The present study was conducted to assess the induction of salt tolerance in seven wheat genotypes (*Bakhtawar-92, Bhakar-2002, Fakhar-e-Sarhad, Khyber-87, Nasir-2000, Pirsabak-2005, and Uqab-2000*) at germination and seedling stage through halo-priming with NaCl. Seeds of each wheat genotype were halo-primed separately. Halo-primed seeds of each wheat genotype were subjected to 0.02 (control), 2, 4, 6 and 8 dS/m NaCl salinity under laboratory conditions. Germination % age varied significantly among various wheat genotypes; however, differences between different salt concentrations were non-significant. All the seedling growth characters (germination, plumule growth, fresh and dry weight of seedling and moisture contents) exhibited significant differences among wheat genotypes as well as under the applied salt concentration except for radicle growth which varied non-significantly under salt stress. Interaction between various wheat genotypes and salt concentration was also significant for all the seedling growth characters, while it was non-significant for germination % age. It is concluded that NaCl proved to be effective priming agents in inducing salt tolerance in the tested wheat genotypes.

Key words: Priming; Salinity tolerance; Wheat; Germination; Seedling growth.

Introduction

World population is increasing at alarming rate and expected to reach about 9.2 billion by the end of year 2050 (Anon., 2009). On the other hand food productivity is decreasing due to the various biotic and abiotic stresses. Therefore, minimizing these losses is a major concern for all the nations to cope with the increasing food requirement. Salinity is a major environmental stress limiting plant growth and productivity of wide range of crops (Athar et al., 2008). Salinity impairs seed germination, retards plant growth and reduces crop yield. Out of 25 million hectares of cultivated land in Pakistan, 10 million hectares are affected with salinity (Anon., 2008). Agriculture is the largest sector in Pakistan economy and account for about 25% of gross domestic product and about 75% of the population depends on agriculture (Anon., 2013). Wheat (Triticum aestivum L.) is one of the most important crops among the cultivated plants with respect to human nutrition. It is a moderately salt tolerant crop and fulfills 95% of the food requirements of our country (Anon., 2013). The significance of soil salinity for agricultural yield is enormous as it affects the establishment, growth and development of plants leading to huge losses in productivity (Mathur et al., 2007).

Various strategies are applied to overcome the deleterious effects of salinity on plants. Seed priming seems to be a promising technique to raise successful crop in arid and semiarid tropics. Various seed priming techniques have been developed; each has advantages and disadvantages and may have varying effects depending upon plant species, stage of plant development, concentration/dose of priming agent, and incubation period (Ashraf & Foolad, 2005). Among the various strategies, pre-sowing soaking of seeds in salt solutions enhance germination and seedling emergence uniformly under

adverse environmental conditions. It is the most important, low cost, low risk and effective approaches for induction of salt tolerance in wheat (Khalil et al., 2010). Seed priming enhances speed and uniformity of germination, induces several biochemical changes in the seed which are required to start the germination process (Asgedom & Becker, 2001; Khalil et al., 2010), resulting in improved stand establishment that can increase the drought and salt tolerance and crop yield (Yari et al., 2011). From the review of literature it is clear that few workers have under taken studies regarding salinity tolerance in Triticum aestivum (Basra et al., 2005; Iqbal & Ashraf, 2005; Afzal et al., 2006 a, b; Iqbal et al., 2006; Khan et al., 2006; Xu et al., 2006; Afzal et al., 2007 a, b; Gurmani et al., 2007; Hajihashemi & Kiarostami, 2007; Iqbal & Ashraf, 2007; Wahid et al., 2007; Afzal et al., 2008; Farooq et al., 2008; Hamid et al., 2008; Nawab & Bakht, 2008; Sakr & El-Metwally, 2009; Abbasdokht et al., 2010; Murungu & Madanzi, 2010; and Yari et al., 2010, 2011).

The present study was therefore, planned to assess the effect of priming on the germination and growth improvement of wheat genotypes under different levels of salinity. The aim was also to determine the genotypic variability in their tolerance to salinity both at the germination and seedling stage.

Materials and Methods

Seeds of seven wheat genotypes viz. *Bakhtawar-92*, *Bhakar-2002*, *Fakhar-e-Sarhad*, *Khyber-87*, *Nasir-2000*, *Pirsabak-2005 and Uqab-2000* were obtained from Agricultural Research Institute (ARI) Pirsabak, Nowshera. Seed priming was done by keeping 1:5 ratios of seed and solution. Seeds of each genotype (500) were exposed to halo-priming with 4 dS/m NaCl solution for 12 hours at 25°C in the dark. Seeds were rinsed thrice and re-dried up to original weight under shade (Basra et al., 2005).

The halo-primed seeds of each genotype were grown in 2, 4, 6 and 8 dS/m saline solution of NaCl. Dry seeds of each wheat genotype were used in the control. Seeds were placed on twice folded Whatman # 1 filter paper as seed beds in petri dishes. Each petri dish in the saline treatments was provided with 8 ml of the respective salt concentration. Ten seeds of each wheat genotype were sown using 10 replicates. While each replicate, in the non-saline environment was provided with the same amount of distilled water. The glassware was thoroughly washed with tap water followed by rinsing with distilled water. The dried glassware was then sterilized at 170°C for 4 hours before use. The experiment was laid out in completely randomized design (CRD) with two factorial arrangements in an incubator at 25°C. The germination, plumule and radicle length were recorded after 72 hours. Five seedlings from each replicate of each treatment were randomly selected for fresh weight and then kept in oven at 65°C for 72 hours for dry weight determination. Moisture contents were measured following Hussain (1989). The collected data was subjected to two ways Analysis of Variance (ANOVA). The means were compared by Least Significant Difference (LSD) test at 5 % level of probability (Steel &Torrie, 1980).

Results

Germination % age: Analysis of variance (ANOVA) revealed that germination % age of wheat genotypes was significantly affected by halo-priming with NaCl under saline conditions. However, salinity and their interaction with genotypes were non-significant (Table 1). Genotypic means showed that *Nasir-2000* exhibited maximum germination followed by *Khyber-87*. While, minimum germination % age was observed in *Bakkar-2002* which is statistically similar to *Bakhtawar-92*. Concentrations means showed that halo-primed seeds performed better than control even at 8 dS/m salinity. Among the interaction *Nasir-2000* showed the highest germination % age at 8 dS/m salt concentration; however, *Bhakkar-2002* showed least germination % age at 6dS/m NaCl (Table 2).

 Table 1. Means squares of the analysis of variance for germination (%), plumule and radicle growth (mm), seedling fresh and dry weight (mg), moisture contents (%).

Source	d.f	Germination (%)	Plumule growth (mm)	Radicle growth (mm)	Fresh weight (mg)	Dry weight (mg)	Moisture contents (%)
Genotypes (G)	6	10528.556^{*}	335.493*	2472.470^{*}	8790.36*	1499.830^{*}	25746.603^{*}
Concentration (C)	4	27.737 ^{NS}	107.134^{*}	108.855 ^{NS}	3220.352^{*}	34.215*	17107.334*
GXC	24	133.737 ^{NS}	56.968^{*}	183.508^{*}	373.796*	32.703^{*}	3567.157*
Error	140	172.920	10.515	73.467	161.715	15.803	1649.213
Coefficient of variation (%)		20.81	16.53	22.23	11.00	10.35	19.67

d.f. = Degree of freedom; NS= Non-significant; *= Significant

Wheat genotypes		Genotypic				
	Control	2 dS/m	4 dS/m	6 dS/m	8 dS/m	means
Bakhtawar-92	40.0	44.0 (110.00)	40.0 (100.00)	40.0 (100.00)	47.0 (117.50)	42.3 ^e
Bhakkar-2002	30.0	36.0 (120.00)	43.0 (143.33)	30.0 (100.00)	40.0 (133.33)	36.0 ^e
Fakhar-e-Sarhad	54.0	52.0 (96.29)	56.0 (103.70)	58.0 (107.00)	50.0 (92.59)	54.0 ^d
Khyber-87	80.0	74.0 (92.50)	86.0 (107.50)	80.0 (100.00)	86.0 (107.50)	81.2 ^b
Nasir-2000	90.0 -	94.0 (104.44)	90.0 (100.00)	92.0 (102.22)	100.0 (111.11)	93.2 ^a
Pirsabak-2005	74.0	66.0 (89.18)	64.0 (86.48)	74.0 (100.00)	62.0 (83.78)	68.0 ^c
Uqab-2000	66.0 -	78.0 (118.18)	66.0 (100.00)	64.0 (96.96)	64.0 (96.96)	67.6 ^c
Concentration Means	62.0	63.5 (102.41)	63.6 (102.58)	62.6 (100.96)	64.2 (103.54)	

LSD value at 0.05 alpha level for genotype means = 7.353

Means in the last column sharing the same letter do not differ significantly from each other at 5% probability level Figures in parenthesis represent % of control

XX71		Genotypic				
Wheat genotypes	Control	2 dS/m	4 dS/m	6 dS/m	8 dS/m	means
Bakhtawar-92	15.9 klmno	20.0 fghij	13.4 ^{nop}	15.0 ^{lmno}	12.1 ^{op}	15.3 ^e
	-	(125.78)	(84.27)	(94.33)	(76.10)	
DI 11 2002	14.5 ^{mno}	19.1 ^{ghijk}	18.2^{ijklm}	23.1 bcdefg	17.9^{ijklm}	10 6 6
Bhakkar-2002	-	(131.72)	(125.51)	(159.31)	(123.44)	18.6 ^c
Fakhar-e-Sarhad	24.2 ^{bcde}	18.1 ^{ijklm}	$19.0^{\text{ hijkl}}$	14.4 ^{mno}	12.7 ^{nop}	17.7 ^{cd}
	-	(74.79)	(78.51)	(59.50)	(52.47)	
Khyber-87	26.5 ^{ab}	25.5 abcd	21.7 defghi	22.2 ^{cdefgh}	20.8 efghi	23.3 ^a
	-	(96.22)	(81.88)	(83.77)	(78.49)	
Nasir-2000	28.8 ^a	25.9 abc	26.1 abc	24.6 bcde	19.3 ghijk	25.0 %
	-	(89.93)	(90.62)	(85.41)	(67.01)	25.0 ^a
Pirsabak-2005	16.5 ^{jklmn}	14.5 mno	10.1 ^p	20.8 efghi	19.1 ^{ghijk}	1 c 2 de
	-	(87.87)	(61.21)	(126.06)	(115.75)	16.2 ^{de}
Uqab-2000	25.4 abcd	23.7 bcdef	22.4 cdefgh	15.6^{klmno}	19.6 ghijk	21.3 ^b
	-	(93.30)	(88.18)	(61.41)	(77.16)	
Concentration	21.7 ^a	21.0 ^a	18.7 ^{bc}	19.4 ^b	17.4 ^c	
Means	-	(96.77)	(86.17)	(89.40)	(80.18)	

Table 3. Effect of halopriming on plumule growth (mm) of different wheat genotypes under NaCl salt stress.

LSD value at 0.05 alpha level for genotype means = 1.813, treatment means = 1.533 and interaction = 4.055

Means in the last column/ rows sharing the same letter do not differ significantly from each other at 5% level of probability Figures in parenthesis represent % of control

Plumule growth: ANOVA exhibited significant variation for plumule growth among wheat genotypes and salt concentrations. Interaction between genotypes and concentrations was also significant (Table 1). Genotypic means showed that maximum plumule growth was recorded in Nasir-2000 closely followed by Khyber-87. Plumule growth in Uqab-2000 was significantly lower than the aforementioned genotypes while, higher than Bhakkar-2002 and Fakhar-e-Sarhad, which were significantly greater than Pirsabak-2005. The growth of plumule in Bakhtawar-92 was significantly lowest among all the tested genotypes. Concentration means revealed that the observed maximum plumule growth under control condition was statistically similar with 2dS/m which decreased significantly at 4 dS/m. Average plumule length at 6 dS/m was statistically same to 4 dS/m salt concentration; however it was significantly greater than plumule growth at 8 dS/m salinity (Table 3).

Radicle growth: ANOVA revealed that radicle growth varied significantly among wheat genotypes. Interaction between genotypes and concentrations was also significant. However, differences due to salt concentrations were non-significant (Table 1). Genotypic means show that maximum radicle length was observed in *Nasir-2000* followed by significantly lower and statistically similar radicle growth in *Khyber-*87, *Pirsabak-2005, Uqab-2000* and *Bakhtawar-92*. Radicle growth in *Fakhar-e-Sarhad* was significantly lower than the aforementioned genotypes however; it was significantly higher than radicle growth in *Bhakkar-*

2002. Concentration means revealed that radicle growth of the halo primed seedlings decreased under salt stress except at 6 dS/m level of salt, whereas it increased as compared to control. Interaction between wheat genotypes and salt concentrations showed that radicle growth was maximum in *Nasir-2000* under control condition, which was statistically similar to radicle growth at 2 dS/m and 4 dS/m salt levels in the same genotype. While, the minimum radicle growth was observed in *Pirsabak-2005* at 4 dS/m salinity (Table 4).

Fresh weight: Results indicated that fresh weight varied significantly among wheat genotypes and salt concentrations. Interaction between genotypes and concentrations was also significant (Table 1). Genotypic means revealed maximum fresh weight/ seedling in Uqab-2000, followed by Nasir-2000, Pirsabak-2005, Fakhar-e-Sarhad, Bhakkar-2002, Khyber-87 and Bakhtawar-92. Concentration means showed that maximum fresh weight in the halo-primed seedlings was recorded in control, which was statistically at par with 2 dS/m and 4 dS/m salinity level respectively. Average fresh weight at 6dS/m level of salt was significantly lower than control and higher than 2 dS/m salinity. Interactions means revealed maximum average fresh weight in the halo-primed seedlings of Uqab-2000 under 2 dS/m level of salt which was statistically similar with average fresh weight at control and 4 dS/m level of salt in the same genotype. The minimum average fresh weight was observed in Bakhtawar-92 at the highest concentration of NaCl (Table 5).

Wheat genotypes		Genotypic				
	Control	2 dS/m	4 dS/m	6 dS/m	8 dS/m	means
	33.8 ghijkl	41.7 defghi	35.0 ghijk	41.1 efghi	35.6 ghijk	37.4 ^b
Bakhtawar-92	-	(123.37)	(103.55)	(121.59)	(105.32)	37.4
DI 11 2002	18.4 ^m	18.8 ^m	25.6 klm	42.0 defgh	24.1 ^{lm}	25 0 d
Bhakkar-2002	-	(102.17)	(139.13)	(228.26)	(130.97)	25.8 ^d
	35.0 ghijk	33.4 ghijkl	35.4 ghijk	28.9 ^{jklm}	26.2 klm	21.0 6
Fakhar-e-Sarhad	-	(95.42)	(101.14)	(82.57)	(78.85)	31.8 °
	$40.6 \ ^{\mathrm{fghi}}$	35.9 ghijk	39.2 fghij	44.1 defg	43.6 defgh	40.7 ^b
Khyber-87	-	(88.42)	(96.55)	(108.62)	(107.38)	
N : 2000	68.4 ^a	60.0 ^{ab}	58.2 abc	51.9 ^{bcd}	51.6 ^{bcde}	- 0.03
Nasir-2000	-	(87.71)	(85.08)	(75.87)	(75.43)	58.0 ^a
D: 1 1 0005	33.1 hijkl	35.8 ghijk	31.2 ^{ijkl}	47.6 ^{cdef}	43.8 defgh	ac a b
Pirsabak-2005	-	(108.15)	(94.25)	(143.80)	(132.32)	38.3 ^b
Uqab-2000	$40.8 \ ^{\mathrm{fghi}}$	40.8 fghi	39.4 fghij	35.1 ghijk	33.5 ghijkl	or o h
	-	(100.00)	(96.56)	(86.02)	(82.10)	37.9 ^b
Concentration	38.6	38.1	37.7	41.5	36.9	
Means	-	(98.70)	(97.66)	(107.51)	(95.59)	

Table 4. Effect of halopriming on radicle growth (mm) of different wheat genotypes under NaCl salt stress.

LSD value at 0.05 alpha level for genotype means = 4.793, and interaction = 10.72

Means in the last column/ rows sharing the same letter do not differ significantly from each other at 5% level of probability Figures in parenthesis represent % of control

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Wheat genotypes		Genotypic				
	Control	2 dS/m	4 dS/m	6 dS/m	8 dS/m	means
Bakhtawar-92	85.5 °	92.9 ^{mno}	83.6 °	87.0 ^{no}	62.3 ^p	82.3 ^f
	-	(108.65)	(97.77)	(101.75)	(72.86)	82.3
	110.6 hijkl	122.7 defghi	120.6 efghi	$129.7 \ ^{cdefg}$	86.9 ^{no}	114.1 ^d
Bhakkar-2002	-	(110.94)	(109.04)	(117.26)	(78.57)	114.1
Fakhar-e-Sarhad	136.0 cde	124.3 defghi	123.6 defghi	103.0 ^{jklm}	108.9 ^{ijkl}	119.2 ^{cd}
	-	(91.39)	(90.88)	(75.73)	(80.07)	
V 1 1 07	114.3 ghijk	111.8 ^{hijk}	102.1 klmn	95.5 ^{lmno}	90.4 ^{mno}	102.8 ^e
Khyber-87	-	(97.81)	(89.32)	(83.55)	(79.09)	
Nasir-2000	138.5 bcd	131.7 ^{cdef}	137.2 ^{cd}	$118.9 \ ^{\mathrm{fghij}}$	$110.2^{\text{ hijkl}}$	127.3 ^b
Nastr-2000	-	(95.09)	(99.06)	(85.84)	(79.56)	127.5
Pirsabak-2005	$125.6^{\text{ defgh}}$	123.1 defghi	117.3 fghijk	$127.8 \ ^{defg}$	$118.6 \ ^{\mathrm{fghij}}$	122.5 ^{bc}
Pirsabak-2003	-	(98.00)	(93.39)	(101.75)	(94.42)	122.5
Uqab-2000	154.0 ab	155.1 ^a	145.4 abc	$123.8 \ ^{defghi}$	125.8 defgh	140.8 ^a
	-	(100.71)	(94.41)	(80.38)	(81.68)	
Concentration	123.5 ^a	123.1 ^a	118.6 ^a	112.2 ^b	100.4 ^c	
Means	-	(99.67)	(96.03)	(90.85)	(81.29)	

LSD value at 0.05 alpha level for genotype means = 7.111, treatment means = 6.010 and interaction = 15.90

Means in the last column/ rows sharing the same letter do not differ significantly from each other at 5% level of probability Figures in parenthesis represent % of control

Wheat genotypes		Genotypic				
	Control	2 dS/m	4 dS/m	6 dS/m	8 dS/m	means
Bakhtawar-92	28.1 lmn	25.9 ^{no}	27.8 ^{lmn}	30.2 klmn	22.3 °	26.9 ^f
	-	(92.16)	(98.93)	(107.47)	(79.35)	
DI 11 2002	45.3 abcd	36.0 ^{ghij}	$40.6^{\text{ defg}}$	41.9 ^{cdef}	34.2 hijk	20.68
Bhakkar-2002	-	(79.47)	(89.62)	(92.49)	(75.49)	39.6 °
Fakhar-e-Sarhad	39.6 efg	42.9 bcde	43.6 ^{b-e}	40.1 efg	46.7 abc	42.6 ^b
	-	(108.33)	(110.10)	(101.26)	(117.92)	
	30.5 klmn	32.3 ^{ijkl}	$31.2^{\ jklm}$	30.2 klmn	26.3 mno	30.1 ^e
Khyber-87	-	(105.90)	(102.29)	(99.01)	(86.22)	
N : 2000	36.0 ^{ghij}	35.7 ^{ghij}	37.8 ^{fgh}	36.9 ^{ghi}	37.0 ^{fghi}	
Nasir-2000	-	(99.16)	(105.00)	(102.50)	(102.77)	36.7 ^d
Dimarkat 2005	49.8 ^a	47.0 ^{ab}	48.7 ^a	45.8 abc	46.3 abc	47.5 ^a
Pirsabak-2005	-	(94.37)	(97.79)	(91.96)	(92.97)	
Uqab-2000	45.3 abcd	45.2 abcd	45.8 abc	45.8 abc	46.2 ^{abc}	45.6 ^a
	-	(99.77)	(101.10)	(101.10)	(101.98)	
Concentration	39.2 ^a	37.9 ^{ab}	39.4 ^a	38.7 ^{ab}	37.0 ^b	
Means	-	(96.68)	(100.51)	(98.72)	(94.38)	

Table 6. Effect of halopriming on dry biomass/ seedling (mg) of different wheat genotypes under NaCl salt stress.

LSD value at 0.05 alpha level for genotype means = 2.223 and interaction = 4.971

Means in the last column/ rows sharing the same letter do not differ significantly from each other at 5% level of probability Figures in parenthesis represent % of control

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Wheat genotypes		Genotypic				
	Control	2 dS/m	4 dS/m	6 dS/m	8 dS/m	means
Bakhtawar-92	214.7 ^{e-k}	285.5 ^a	203.7 ^{f-m}	188.4 ^{h-o}	189.0 ^{h-o}	216.3 ^b
	-	(132.97)	(94.78)	(87.75)	(88.02)	216.3
DL =11 == 2002	152.4 nop	233.1 ^{b-i}	197.1 ^{g-n}	209.2 ^{f-1}	152.9 ^{nop}	189.0 ^{cd}
Bhakkar-2002	-	(152.95)	(129.33)	(137.27)	(100.32)	189.0
	245.2 ^{a-g}	189.4 ^{h-o}	183.2 ^{i-p}	159.4 ^{1-p}	133.6 ^p	182.2 ^d
Fakhar-e-Sarhad	-	(77.24)	(74.71)	(65.00)	(54.48)	182.2
Vlash or 97	273.2 abc	248.8 ^{a-f}	226.5 ^{c-j}	218.9 ^{d-k}	245.5 ^{a-g}	242.6 ^a
Khyber-87	-	(91.06)	(82.90)	(80.12)	(89.86)	
Nasir-2000	280.6 ab	269.5 a-d	263.1 ^{a-e}	222.0 ^{d-k}	197.7 ^{g-n}	246.6 ^a
<i>Nusir-2000</i>	-	(96.04)	(93.76)	(79.11)	(70.45)	240.0
Ding ab ab 2005	154.2 ^{m-p}	162.7 ^{1-p}	141.0 ^{op}	179.3 ^{j-p}	157.3 ^{m-p}	150.0 %
Pirsabak-2005	-	(105.51)	(91.43)	(116.27)	(102.01)	158.9 ^e
Uqab-2000	239.2 ^{a-h}	243.4 ^{a-g}	220.3 ^{d-k}	172.6 ^{k-p}	172.8 ^{k-p}	209.7 ^{bc}
	-	(101.75)	(92.09)	(72.15)	(72.24)	209.7
Concentration	222.8 ab	233.2 ^a	205.0 ^{bc}	192.9 ^{cd}	178.4 ^d	
Means	-	(104.66)	(92.01)	(86.57)	(80.70)	

Table 7. Effect of halopriming on seedling moisture contents (%) of	f different wheat genotypes under salt stress.
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LSD value at 0.05 alpha level for genotype means = 22.71, treatment means = 19.19 and interaction = 50.78

Means in the last column/ rows sharing the same letter do not differ significantly from each other at 5% level of probability Figures in parenthesis represent % of control

Dry weight: Our results revealed significant variation among wheat genotypes for dry weight of seedling. Interaction between genotypes and salt concentrations was also significant. However, differences between salt concentrations were non-significant (Table 1). Genotypic means indicated that maximum average dry weight of seedling was observed in Pirsabak-2005, which was statistically similar with Ugab-2000 followed by Fakhare-Sarhad, Bhakkar-2002, Nasir-2000, Khyber-87 and Bakhtawar-92. Concentrations means revealed that average dry weight of seedling was maximum at 4 dS/m level of salt followed by control, 6 dS/m, 2 dS/m and 8 dS/m levels of salt respectively. Interaction means showed maximum average dry weight in the halo-primed seedlings of Pirsabak-2005 under control condition which was statistically similar to average dry weight at 4 dS/m level of salt in the same genotype. The minimum average dry weight of seedling was observed in Bakhtawar-92 at 8 dS/m concentration of salt (Table 6).

Moisture contents: ANOVA exhibited that moistures contents varied significantly among wheat genotypes and salt concentrations. Interaction between genotypes and concentrations was also significant (Table 1). Genotypic means showed that maximum moisture contents was observed in Nasir-2000, which were statistically similar to moisture contents in Khyber-87 followed by Bakhtawar-92 which were in turn statistically at par with Uqab-2000. Moisture contents in Bhakkar-2002 and Fakhar-e-Sarhad were 189.0% and 182.2%, respectively. The minimum moisture contents were observed in Pirsabak-2005 which was significantly lower among all the genotypes studied. Concentration means reveal that moisture contents increased at 2 dS/m salt level which were statistically similar to moisture contents in control followed by 4 dS/m and 6 dS/m concentration of salt. The least moisture contents were recorded under 8 dS/m level of salt. Interaction means showed that moisture contents were highest in Bakhtawar-92 under 2 dS/m level of salt which were statistically similar to moisture contents in Nasir-2000 under control condition. The moisture contents in Fakhar-e-Sarhad at 8 dS/m level of salt were the lowest at any concentration of salt among the wheat genotypes studied (Table 7).

Discussion

Halo-priming with NaCI induces salt tolerance by diminishing the inhibitory effects of salinity on the germination and seedling growth of wheat (Afzal *et al.*, 2006; Iqbal *et al.*, 2006; Afzal *et al.*, 2008; Farooq *et al.*, 2008). The improved germination of the halo-primed seeds under increasing salinity in the present study is in contrast with Basra *et al.* (2005) and Afzal *et al.* (2007) who concluded that NaCl priming was ineffective in improving germination and seedling vigor of wheat under saline conditions. However, the present findings are in line with those of Iqbal & Ashraf (2007), who stated that priming increased the germination percentage in wheat. It is also reported that halo-priming with NaCl has improved the germination of wheat, sugarcane and sweet sorghum (Afzal *et al.*, 2008; Patade *et al.*, 2009; Patanè *et al.*,

2009), which is also in accordance with the results of this study. The increased germination in halo-primed seeds under salt stress could be due to the faster water absorption occurring in primed seeds as compared to the non-primed (Patanè *et al.*, 2009). Germination % age was maximum in *Bakhtawar-92*, *Fakhar-e-Sarhad*, *Khyber-87* and *Nasir-2000* at higher levels of salt whereas; it was maximum at lower levels of salt in *Bhakkar-2002* and *Uqab-2000*.

Significantly reduced plumule growth of seeds subjected to halo-priming under salt stress is in contrast to Wahid et al. (2008), Khan et al. (2009) and Patade et al. (2009), who reported that NaCl priming improved shoot or plumule length in sunflower, Capsicum annuum and sugarcane, respectively. Interaction between wheat genotypes and salt levels also elucidate that plumule growth in the halo primed seedlings decreased under increasing level of salt in Bakhtawar-92, Fakhar-e-Sarhad, Khyber-87, Nasir-2000, Uqab-2000, especially at higher concentrations of salt. The 2 dS/m treatment in Bakhtawar-92 and Khyber-87 and higher doses in Pirsabak-2005 enhanced plumule growth however, it increased in Bhakkar-2002 at all levels of applied salt. Non-significant decrease in the radicle growth of the haloprimed seedlings under different concentration of salt in the present study is in line with the findings of Amjad et al. (2007), who reported that seed priming with NaCl had no significant effect on root length of seedlings. On the other hand, it is also reported that priming with NaCl improved root or radicle length in sunflower and Capsicum annuum (Wahid et al., 2008; Khan et al., 2009), which disagree with our results. Radicle growth in the halo primed seedlings of Bakhtawar-92, Bhakkar-2002, Khyber-87 and Pirsabak-2005 increased while, it decreased in Fakhar-e-Sarhad, Nasir-2000 and Uqab-2000 under salt stress. Both the maximum increase and decrease in different genotypes were observed at higher salt concentrations.

Decrease in fresh weight of halo-primed seedling under salt stress in the present investigation is contradictory to the observations of Amjad et al. (2007) and Khan et al. (2009) who reported non-significant effect on fresh weight of seedlings under NaCl priming. Average seedling fresh weight of the halo-primed seeds of wheat genotypes responded in a different way under salt stress. Reduced seedling fresh weight was noted in Fakhar-e-Sarhad, Khyber-87 and Nasir-2000; however, in Pirsabak-2005 it decreased up to 4 dS/m salt level and increased thereafter. Average fresh weight in Uqab-2000 increased at 2 dS/m level of salt which decreased at the subsequent higher concentration. While, in Bhakkar-2002 it increased up to 6 dS/m level of salt and declined at the highest level of salt. However, in Bakhtawar-92 it increased or decreased variously under salt stress. Significant decrease in the dry weight of haloprimed seedling under different concentration of salt in the present study disagrees with the findings of Farhoudi & Sharifzadeh (2006), who reported higher dry weight of seedlings derived from haloprimed seeds of canola. Interaction revealed that dry weight of the haloprimed seedlings of different genotypes increased or decreased variously under salt stress. Dry weight of the haloprimed seedlings of Fakhar-e-Sarhad, Nasir-2000 and Uqab2000 increased whereas, in *Bhakkar-2002* and *Pirsabak-2005* it decreased under salt stress. While, dry weight in *Bakhtawar-92* and *Khyber-87* increased at lower concentrations and decreased at higher concentrations of salt. Moisture contents in the haloprimed seedlings of *Fakhar-e-Sarhad*, *Khyber-87* and *Nasir-2000* showed a decreasing trend under salt stress while it increased in *Bhakkar-2002* and *Pirsabak-2005*. Low dose of salt tended to increase moisture contents in *Bakhtawar-92* and *Uqab-2000* however, higher doses decreased it in both the genotypes.

It is concluded that pre-treated wheat seeds with NaCl can be used to enhance salt resistance in terms of improved germination and seedling growth. Moreover, it is suggested that halo-primed wheat seeds with different concentrations of NaCl salt can help to find a priming medium more suitable for better germination, plant growth and mineral components under control and saline field conditions.

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