

INFLUENCE OF NaCl SALINITY ON GERMINATION AND EARLY SEEDLING GROWTH IN WHEAT (*Triticum aestivum*, L.).

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ABSTRACT

Some wheat genotypes were evaluated to salt stress during the germination and early seedling growth. Salinity, in general decreased germination percentage and seedling growth characters representing by root and shoot lengths as well as their fresh and dry weight in most studied genotypes depending on the genotypes and the level of NaCl salinity. However, genotypes Gemmeiza 7 and Gemmeiza 10 x C.B.255 showed a stimulation response to NaCl salinity up to 6000 mg/l. The stimulating effect of salinity on the two mentioned genotypes was discussed.

On the other hand, total sugars and proline concentrations were significantly increased in the shoots of all genotypes grown under salt stress. Moreover, Na⁺ concentration was increased whereas, that of K⁺ decreased in the shoots.

The data concluded that, cv(s) Gemmeiza 7 and Gemmeiza 10x C.B.255 of wheat were more tolerant to salt stress up to 6000 mg NaCl/l and the former cv proved to be more tolerant as compared to all studied genotypes.

Keywords: Wheat, salinity, germination, seedling growth.

INTRODUCTION

Wheat (*Triticum aestivum*, L. Poaceae) is one of the most important cereal crops in Egypt. Its production is still insufficient to overcome the consumption demand due to the limited area and restricted resources of the fresh water. Salinity is one of the main limiting factor for its cultivation, seedling growth and productivity. The common approach in breeding for salt stress tolerance is to select for its components. Breeders assume that, there is considerable variability in abiotic stress tolerance in wheat germplasm. It is closely related characteristics that influenced by genetic (Varietals), environmental conditions, seed processing, physiological treatment effects and others (Helaly *et al.*, 2009-b). However, several of these components are difficult to measure, indirect selection showed be applied (Visser, 1994).

Studied on abiotic stress (salinity and drought) tolerance in wheat have been undertaken for the identification of environmental and physiological factors. Germination percentage of wheat measured under standard conditions correlated with the seedling establishment in the field under stress condition (Sadeghian and Yavari, 2004). Unfavourable germinations such as salinity have serious impacts on the results of vigour tests and plant establishment of wheat and delayed seedling emergence. Small differences in the NaCl levels did not change number of germinated grains but greatly affected water uptake and seedling growth (Bray, 2000).

Sadeghian and Yavari (2004) reported a similar germination rate for primed saline and primed seeds in plots with low salinity level. This practice is certain to become a seed enhancing requirement for crop production under stressed conditions. Tugnoli and Bettin (2001) found that, productivity and germination are linked to key properties of varieties used and tend to decide the ultimate crop yield. Several investigators (Sadeghian and Yavari, 2004 ; Helaly *et al.*, 2009-b) found that, seed germination and early seedling growth are the stages most sensitive to salt stress.

On the other hand, breeding programs should be explicitly direct towards development of salt –tolerant varieties due to tissue culture technique. Foolad *et al.*, (2003) studied the germination response of tomato accessions in Petri dishes containing agar media treated with zero mM NaCl (non-stress) and cold stress treatments or 150 mM NaCl +15 mM CaCl₂ for salt stress treatments. They concluded that, some of the same genetic and physiological parameters contributing to rapid seed germination under non-stress conditions could also facilitate rapid seed germination under stress.

Therefore, the present investigation was performed for the evaluation of genetic trials and to gain some information on the effects of NaCl salinity on germination and certain morphological parameters of some wheat genotypes at the early seedling stage. These genotypes exhibiting contrasting levels of salt resistance.

MATERIALS AND METHODS

The present investigation was carried out under controlled conditions at the Labs of Department of Agric. Botany, Faculty of Agriculture , Mansoura University, Egypt during the growing season of 2007/2008.

Plant material:

Grains of eight wheat genotypes (*Triticum aestivum*, L. Poaceae) with different backgrounds were obtained from the Field Crops Research Institute, Gemmeiza Research Station, ARC, Egypt. The characteristics of these genotypes are presented in Table 1.

Germination experiment:

Grains of each genotype (8 genotypes) were surface sterilized by soaking in 0.001 HgCl₂ for one min., washed with distilled water and germinated between discs of filter papers (Whatman No. 1) in covered glass Petri dishes (12 cm). The grain beds were moistened with 10 ml of half Hoagland strength nutrient solution (Hewitt, 1952) , supplemented with NaCl at four levels denoted 0 (control), 4000, 6000 and 8000 mg/l for each genotype. Each treatment was replicated five times (5 dishes) and each replicate contained 10 grains. The germination was took place for two weeks in darkness in an incubator at 28 °C. Germination percentage was recorded, daily, during the incubation period . The relative germination (GR) was calculated (Smith and Dobrenz, 1987) .

$$GR = \frac{\text{Number of germination grains in the stressed medium} \times 100}{\text{Number of germination grains in control medium}}$$

Table (1): The commercial name , identification characteristics and sources of the wheat (*Triticum aestivum*, L.) genotypes evaluated in the present investigation.

No	Pedigree	Origin
Gemmeiza 7	CMH74 A.630/Sx11 Seri. 82/ Agent. C Gm 4611-2 Gm-3 Gm- 1 Gm-0 Gm	Gemmeiza Res. Stat.
Gemmeiza 9	ALD "5"/HUAC // CMH 74 A. 630/Sx C Gm 4583-5 Gm-1 Gm-0 Gm	Gemmeiza Res. Stat.
Gemmeiza 10	Maya 74 "5" /on // 1160 147/31 BB/ Gll/4/ CHAt "5"/5/Crows CGm 5820-3 Gm-1 Gm-2 Gm- 0 Gm	Gemmeiza Res. Stat.
Sakha 94	OPAtA/Rayon // KAUZ. CUBW 90 Y 3180-OTO PM- 10 M-015 Y- 0Y- 0AP-05.	Gemmeiza Res. Stat.
In addition there are four hybrids namely:		
Gemmeiza 9xC.B.255	C.B.255: C 182.24/ C 168.3/ 3/ Cno* 2/7C//Cc /Tob SWM 6828- 6AP- 2AP- 2AP -IAP-2 AP- IAP- OAP	Mexico
Gemmeiza 9xC.B.254	C.B.254: BT 2549/ Fath T 78- 156-4 BJ- 1 BJ-13 BJ- 8 BJ- 1 BJ- 0 BJ	Mexico
Gemmeiza 10xC.B.254		Gemmeiza Res. Stat.
Gemmeiza 10xC.B.255		Gemmeiza Res. Stat.

After germination, seedlings were allowed to grow for further two weeks from germination. At the end of experiment, the following characteristics were studied:

Root length (cm), shoot length (cm), root fresh and dry weights (mg), shoot fresh and dry weights (mg).

Seedlings were taken and separated into their roots and shoots. Fresh weight was immediately determined to avoid any loss of water content. Thereafter, roots and shoots were dried overnight in an electric oven at 70 °C until a constant weight and attained.

Total sugar was estimated using the method of Sadasivam and Manikam (1996) as well as proline (Bates *et al.*, 1973). Moreover, 50 mg dried materials from the roots or 100 mg from the shoots were taken for digestion in a mixture of sulphoric acid, salicylic acid and hydrogen peroxidase (Hanaa Fatouh, 2003). Na⁺ and K⁺ were determined by an atomic absorption Spectrophotometers (GBC, 932 AA).

RESULTS AND DISCUSSION

Decreased germination under salt stress may be measured as delaying of emergence, reduction of the ultimate germination percentage or both. Table (2) shows that, the reduction in germination percentage and the increase in time required for wheat grains to germinate due to salt stress were observed in all genotypes under investigation. However, genotypes Gemmeiza 7 and Gemmeiza 10 x C.B. 255 were more tolerant to saline condition particularly at the lowest salinity levels (4000 and 6000 mg NaCl/l) as compared to all other ones since they were less affected by salinity.

The harmful effects of salt stress on germination may be due mainly to either colloidal inhibition of water occurred by grains and /or imbalanced osmotic water uptake occurred by germinated seeds (Helaly *et al.*, 2009-a). Moreover, altering the hormonal balances and decreasing endogenous cytokinins biosynthesis and auxin production were reported under stress by Schmidt (2005). The latter author added that, stress decreased the accumulation of reducing sugars within the plant tissues which decreased wilting resistance. Moreover, it was found that salinity stress did not induce an increase of ascorbic acid which not only quenches reactive oxygen but also regenerates α -tocopherol (Gadalla, 2009).

The harmful effect of stress on germination and seedling growth represented by the dry matter accumulation seemed to be due to the suppression of plant metabolism under such condition (Demir *et al.*, 2003). Salt stress adversely affect the physico-chemical properties of the protoplasm and cell membranes (Tajdoost *et al.*, 2007). Inhibition of cytokinin biosynthesis and hormonal unbalances, reducing water content and some plant nutrients uptake as well as biosynthesis of α -tocopherol , ascorbic acid and net photosynthetic rate accompanied with high respiration rate were also reported under stress conditions (Tripathi *et al.*, 2007).

Table (2): Effect of NaCl salinity on germination percentage of wheat genotypes.

Genotypes \ Salinity (mg/l)	0	4000	6000	8000	Mean
Gemmeiza 7	99.7	98.7	98.0	97.3	98.3
Gemmeiza 10xC.B. 255	99.3	98.3	95.3	90.2	95.8
Gemmeiza 10	98.8	91.5	86.4	71.5	87.0
Gemmeiza 9	98.6	95.5	86.6	75.1	88.9
Gemmeiza 9 x C.B. 255	98.4	96.6	86.7	71.9	88.4
Gemmeiza 9 x C.B. 254	98.3	96.0	87.7	78.3	90.1
Gemmeiza 10xC.B. 254	98.2	93.3	86.8	71.7	87.5
Sakha 94	98.2	87.1	77.8	70.3	83.3
Mean	98.7	94.6	77.3	78.3	89.9
LSD at 5% for:					
Genotypes	0.11				
Salinity	0.08				
Gen. x Salinity	0.21				

Root and shoot length of the seedlings:

Table (3) shows that, all investigated wheat genotypes except that of Gemmeiza 10x C.B.255 showed a decrease in their root and shoot lengths of their seedlings with an increase in NaCl salinity levels. However, both genotypes Gemmeiza 7 and Gemmeiza 10x C.B.255 showed an increase in the length of their roots and shoots due to an increase in salinity levels up to 6000 mg/l thereafter decreased. The stimulating effects of low salinity levels (4000 and 6000 mg/l) were previously reported by Chen and Murata (2002) on wheat plant who attributed this increase to the beneficial effects of salinity on both cell division and cell enlargement . The inhibitive effects of high salt level may be due to its effect on apical growth as well as hormonal balances within the plant tissues (Abo Shama and Hegazy, 2009) .

Table (3): Effect of NaCl salinity on root and shoot length (cm) of wheat genotypes at four weeks from sowing.

Treatments Genotypes	Root length (cm)					Shoot length (cm)				
	Salinity (mg/l)									
	0	4000	6000	8000	Mean	0	4000	6000	8000	Mean
Gemmeiza 7	15	19	18	7	15	18	20	18	9	16
Gemmeiza 10xC.B. 255	16	17	19	8	15	18	19	20	11	17
Gemmeiza 10	17	15	10	5	12	15	16	12	8	13
Gemmeiza 9	18	16	9	6	12	20	18	12	8	14
Gemmeiza 9 x C.B. 255	16	15	13	12	14	18	16	13	14	15
Gemmeiza 9 x C.B. 254	18	16	13	12	15	15	12	9	8	11
Gemmeiza 10xC.B. 254	17	16	12	11	14	15	14	12	9	12
Sakha 94	17	15	13	12	14	16	12	11	9	12
Mean	16	16	13	9	13	17	16	13	9	14
LSD at 5% for:										
Genotypes	1.8					1.9				
Salinity	1.0					0.9				
Gen. x Salinity	3.5					3.7				

Seedlings fresh and dry weights:

Data presented in Tables (4 and 5) show that, salinity decreased fresh and dry weights of the roots and shoot systems. This reduction was found to be a concentration and genotype dependant. However, root fresh and dry weights of all investigated genotypes were decreased under NaCl salinity except that of genotypes Gemmeiza 7 and Gemmeiza 10 x C.B.255 which showed an increase at NaCl salinity up to 6000 mg/l thereafter decreased. The variation between the genotypes may be due to the genetic action (Esmail, 2005). The same trend was recorded for fresh and dry weight of the shoot system. In this context, Chen and Murata (2002) found that, plant growth was stimulated at the low salinity level whereas inhibited as the concentration of salinity was raised. Several investigators recorded an inhibition of growth caused by salinity due to its effects on hormonal balance (Schmidt, 2005), insufficient supply of ions or other solutes to the growing region (Gadalla, 2009), insufficient osmotic solutes to generate turgor (Hatung, 2004).

Table (4): Effect of NaCl salinity on shoot fresh and dry weights (mg/plant) of wheat genotypes at four weeks from sowing .

Treatments Genotypes	Shoot fresh weight (mg)					Shoot dry weight (mg)				
	Salinity (mg/l)									
	0	4000	6000	8000	Mean	0	4000	6000	8000	Mean
Gemmeiza 7	195	202	145	74	154	18	19	17	9	16
Gemmeiza 10xC.B. 255	199	213	200	73	171	19	21	18	11	17
Gemmeiza 10	160	142	105	74	120	16	14	12	9	13
Gemmeiza 9	240	210	199	95	186	21	20	18	11	17
Gemmeiza 9 x C.B. 255	199	175	140	125	160	19	17	16	13	16
Gemmeiza 9 x C.B. 254	180	165	85	50	120	19	17	11	7	13
Gemmeiza 10xC.B. 254	165	160	122	95	135	16	15	14	10	14
Sakha 94	250	190	155	120	179	22	20	13	11	16
Mean	198	182	144	88	153	19	18	15	10	15
LSD at 5% for:										
Genotypes	6					1.8				
Salinity	4					0.9				
Gen. x Salinity	12					3.7				

Table (5): Effect of NaCl salinity on root fresh and dry weights (mg/plant) of wheat genotypes at four weeks from sowing.

Genotypes	Root fresh weight (mg)					Root dry weight (mg)				
	Salinity (mg/l)									
	0	4000	6000	8000	Mean	0	4000	6000	8000	Mean
Gemmeiza 7	99	105	87	30	80	5.5	5.7	4.2	2.1	4.4
Gemmeiza 10xC.B. 255	100	115	120	50	96	6.1	6.5	7.6	3.9	6.0
Gemmeiza 10	102	99	44	25	67	6.8	6.1	3.5	2.8	4.8
Gemmeiza 9	120	101	86	54	90	7.7	7.5	4.8	2.8	5.7
Gemmeiza 9 x C.B. 255	133	120	115	99	12	7.9	7.2	7.0	6.8	7.2
Gemmeiza 9 x C.B. 254	105	95	42	18	65	6.0	5.1	3.2	2.0	4.1
Gemmeiza 10xC.B. 254	115	96	66	45	80	7.0	6.8	6.2	6.0	6.5
Sakha 94	136	120	77	56	97	7.0	6.1	4.2	3.1	5.1
Mean	114	106	80	47	86	6.7	6.4	5.1	3.7	5.5
LSD at 5% for:										
Genotypes	6					0.25				
Salinity	4					0.18				
Gen. x Salinity	12					0.50				

Organic Solutes Accumulation:

1- Free proline:

Data presented in Table (6) reveal that, proline concentration was increased in the shoots of all studied genotypes due to saline treatment up to the highest level (8000 mg NaCl/l). The obtained results confirmed the well-documented results recorded in the majority of cases that the main feature of increasing salinity concentration is the accumulation of proline in the tissues as did with sugars, it is the striking sequence of salt stress. Consequently, increasing salinity level caused an increased accumulation of proline in the tissues of stressed wheat seedling.

Table (6): Effect of NaCl salinity on proline concentration (mg/g F.Wt.) in the shoots of wheat genotypes at four weeks from sowing.

Genotypes	0	8000	Mean
Gemmeiza 7	11.583	17.353	14.468
Gemmeiza 10 x C.B. 255	6.200	10.373	8.287
Gemmeiza 10	2.617	4.033	3.325
Gemmeiza 9	2.510	3.437	2.973
Gemmeiza 9 x C.B. 255	2.000	3.000	2.500
Gemmeiza 9 x C.B. 254	2.667	4.153	3.410
Gemmeiza 10 x C.B. 254	2.477	3.450	2.963
Sakha 94	1.847	3.607	2.727
Mean	3.988	6.176	5.082
LSD at 5% for:			
Genotypes	0.473		
Salinity	0.237		
Gen. x Salinity	0.673		

The considerable accumulation of proline as a striking sequence and main feature of increasing water or salt stress has been previously confirmed by several workers, due to its major physiological functions under such stress

conditions. These functions include osmoregulation; as a compatible cytoplasmic solute it apparently counteracts the osmotic potential of the vacuole salts (Bray, 2000). Higher osmolytes accumulation especially proline and soluble proteins seems to be related to salt tolerance in wheat as shown in the present investigation and not to be a consequence of tissues reaction to salt stress damage. Kholova *et al.*, (2009) reported that, the deleterious effects of salt on plant growth of wheat may be based on the higher magnitude of total free amino acids and proline. Siddiqui *et al.*, (2008) added that, proline and other compatible solutes are believed to cause the minimal inhibition of metabolism. Proline is organic osmolytes solute with an amphiphilic molecule protects the hydrophobic parts of protein which suffer first when water potential is lowered. The proline functions also include enzyme and membrane protection against salt inactivation (Tajdoost *et al.*, 2007) and as an indicator of plant resistance to stress. Chen and Murata, (2002) strongly suggested that the function and the accumulated proline and other compatible solutes that are not toxic under NaCl salinity, compatible solutes are defined as water-soluble organic compounds of a low molecular weight termed as osmoprotectants. Natural osmoprotectant concentration in cytoplasmic compartments are osmotically significant because they have pivotal roles in maintaining cell turgor and the driving gradient for water uptake under stress (Rontein *et al.*, 2002) allowing processes such as stomatal opening, photosynthesis and cell expansion (Serraj and Sinclair, 2002). Proline did not start to accumulate in the leaves until the concentration of total monovalent cations reached a threshold of approximately 200 μ mol/g fresh weight. In this connection, Bray (2000) reported that, in addition to the role of organic solutes accumulation on cell water relations, it may help towards the maintenance of ionic homeostasis and of the C/N ratio removal of free radicals and stabilization of macromolecules and organelles such as proteins, protein complex and membranes. In plants, the major compatible osmoprotectant solutes are glycinebetaine and proline (Misra and Gupta, 2005), are thought to function as osmoprotectant for protein (Bohnert and Jensen, 1996). These solutes provide a protective environment for enzymes and macromolecular structure and function. The contributory role of osmoprotectants (glycinebetaine and protein) to osmotic adjustment under salt stress was confirmed in various plant species (Meloni *et al.*, 2001). Thus, solute accumulation was considered as a suitable screening parameter for salinity tolerance.

2- Total sugar:

Data presented in Table (7) show that salinity increased total sugars in the shoots in all wheat genotypes. These results indicate a positive correlation between salinity and sugar accumulation potential with special referring to the superiority of genotype Gemmeiza 7 which greatly exceeded the other studied genotypes in their sugar concentrations particularly at the highest salinity level. In this context, Larcher (1995) reported that, glycophytes adapt themselves to some wheat saline conditions by lowering osmotic potential through converting starch to sugar. Helaly *et al.*, (2009-a) found that, the photosynthetic activity per unit assimilating area may not be greatly altered by moderate saline level, even some enhancement might

occurred. This may be due , partly, to increased stability of chlorophyll-protein- lipid complex in the chloroplast of plants grown under salinity (Serraj and Sinclair, 2002) . Sharma *et al.*, (1996) concluded that, sugars and proline accumulations were considered as a suitable screening parameter for chloride salinity tolerance and its mediated and concomitant through decline in protein synthesis, N and K concentrations and K/Na ratio. Tajdoost *et al.*, (2007) suggested that, the increment in soluble carbohydrate due to salinity may, in turn , play an important role in increasing the osmotic pressure of the cytoplasm.

Bartels and Sunkar (2005) found a strong correlation between sugar accumulation and osmotic tolerance. Hence, improvement of crop performance by increasing osmotic potential-adjusting ability might be more significant in increasing plant growth. The current hypothesis is that sugars acts as osmolytic and /or protect specific macromolecules and contribute to the stabilization of membrane structure (Hatung, 2004).

The accumulation of sugars was the result of an enhanced efficiency in the use of carbon coupled to a reduction in cellular metabolism, that could favour the accumulation of respiratory substrate to support the osmotic adjustment required to survive in saline media (Tajdoost *et al.*, 2007). This accumulation has been attributed to an impaired carbohydrates utilization and reduced respiration rate at high salinity level. However, the significance of sugars as an adjustment is still in debate and varies according to the plant species.

Table (7): Effect of NaCl salinity on total sugars concentration (mg glucose/g D.Wt.) in the shoots of wheat genotype at four weeks from sowing.

Salinity (mg/l)	0	8000	Mean
Genotypes			
Gemmeiza 7	77,8	123,9	90,3
Gemmeiza 10 x C.B. 255	03,4	97,1	70,3
Gemmeiza 10	47,1	97,3	72,1
Gemmeiza 9	47,0	07,7	02,6
Gemmeiza 9 x C.B. 255	01,1	9,1	7,6
Gemmeiza 9 x C.B. 254	46,1	80,0	60,7
Gemmeiza 10 x C.B. 254	40,3	00,6	0,4
Sakha 94	49,2	08,1	03,7
Mean	0,8	83,1	67.1
LSD at 5% for:			
Genotypes		4.7	
Salinity		2.4	
Gen. x Salinity		6.7	

3- Na⁺ and K⁺ as well as their ratio:

Table (8) shows that, sodium concentration in the roots as well as in the shoots of all studied genotypes was increased due to an increase in NaCl whereas, that of potassium was decreased. Na⁺ / K⁺ ratio was also increased due to the increase in Na⁺ influx . The promotion of Na⁺ uptake by salinity was accompanied by a corresponding decrease of K⁺ concentration showing an apparent antagonism between K⁺ and Na⁺.

It is well documented that, nutrient concentration and accumulation were adversely affected under salinity as reported by many investigators (Helaly *et al.*, 2009-a and Gadalla, 2009). Several mechanisms may be responsible by a decline in K⁺ concentration with increasing salinity including the antagonism of sodium at the site of uptake in roots (Hatung, 2004).

Table (8): Effect of NaCl salinity on Na⁺ ,and K⁺ concentration (mg/g dry weight) as well as K⁺/Na⁺ ratio in the shoot of wheat genotypes at four weeks from sowing.

Treatments Genotypes	Na ⁺			K ⁺			K ⁺ /Na ⁺		
	Salinity (mg/l)								
	0	8000	Mean	0	8000	Mean	0	8000	Mean
Gemmeiza 7	2.53	4.60	3.56	31.03	17.20	24.37	12.48	3.76	8.12
Gemmeiza 10xC.B. 255	3.13	5.36	4.24	24.70	14.13	19.37	7.86	1.84	4.80
Gemmeiza 10	3.13	6.90	5.01	22.97	10.00	18.98	7.33	2.18	4.70
Gemmeiza 9	3.10	6.87	4.98	21.00	21.23	21.42	7.94	3.11	0.22
Gemmeiza 9 x C.B. 255	3.33	7.87	5.60	24.40	14.33	19.37	7.35	1.82	4.08
Gemmeiza 9 x C.B. 254	3.20	8.13	5.66	21.77	14.57	18.17	6.81	1.81	4.30
Gemmeiza 10xC.B. 254	2.83	8.50	5.66	21.33	16.50	18.92	7.56	1.94	4.70
Sakha 94	2.90	8.43	5.66	21.20	10.87	18.03	7.33	1.88	4.61
Mean	3.02	7.08	5.05	22.66	16.12	19.89	7.96	2.29	5.12
LSD at 5% for:									
Genotypes	0.44			1.4			0.70		
Salinity	0.21			0.7			0.35		
Gen. x Salinity	0.62			2.0			1.01		

In addition, salinity appears to affect the distribution pattern of nutrient elements within various plant organs (Yang *et al.*, 2003) on wheat plants. In a saline conditions, plants take up excessive amount of Na⁺ as in halophytes resulting in high Na⁺ /Ca⁺⁺ and lower K⁺/Na⁺ ratios which may impair the selectivity of the root membrane (Gadalla, 2009). The increase in Na⁺ concentration mainly in the vacuole provides an osmotic adjustment of salt affected plants (Sakr *et al.*, 2004). This accumulation might be due to the important role of sodium in increasing osmotic pressure which facilitate absorption of water need for plants to tolerate the harmful effect on growth caused by salinity. The hyper-accumulation of sodium has direct toxic effect, as it interferes with enzyme structure and function. It may also interfere with the function of potassium as a co-factor in various reactions. Many of the deleterious effects of Na⁺, however, seems to be related to the structure change observed in various membranes of salt stressed plants, and the

plasmalemma has been shown to lose its specific permeability (Helmy, 2008). On the other hand, the increase in vacuolar sodium induced with sodium chloride was corresponded by a decrease in K^+ in the cytoplasm, of xylem parenchyma cells, or of the shoot (Yeo *et al.*, 1977).

The reduction of internal potassium concentration could be related to (i) increased potassium efflux into the growth medium and inhibit transport of this ion root and up to the shoot (Cramer *et al.*, 1989). (ii) the antagonism between K^+ and Na^+ cations, which increased considerably as salinity increased (Mozafar and Oertli, 1990). This antagonism may be due to the direct competition between sodium and potassium on the absorption sites of roots leading to a reduced level of internal K^+ at high external NaCl concentrations (Ben-Hayyim *et al.*, 1989). (iii) excess of Na^+ in the root media results in a passive accumulation of this ion in the root and shoot lead to a high Na^+/K^+ ratio and reduced plant growth (Helmy, 2008).

The increase in Na^+/K^+ ratio noticed in the present investigation due to excessive amount of Na^+ and Cl^- in the media may be attributed to low accumulation of Na^+ and Cl^- in the plant organs which, in turn, impairs the selectivity of the root membrane (Misra and Gupta, 2005 and 2006).

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تأثير ملوحة كلوريد الصوديوم على إنبات ونمو بادرات القمح .
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تم تقييم بعض الطرز الوراثة لنبات القمح تحت الإجهاد الملحى أثناء الإنبات والنمو المبكر للبادرة. أدت الملوحة بصفة عامة الى نقص فى نسبة الإنبات وصفات البادرة معبرا عنها بأطوال الجذور والمجموع الخضرى فضلا عن أوزانها الطازجة والجافة فى معظم الطرز تحت الدراسة معتمدا فى ذلك على الطراز الوراثة ومستوى الملوحة. وعلى أية حال, أظهر كلا الطرازين جمييزة ٧ وهجين جمييزة ١٠ x C.B. ٢٥٥ إستجابة محفزة للملوحة حتى مستوى ٦٠٠٠ مليجرام/لتر وتمت مناقشة هذا التأثير المنشط . من جهة أخرى , زادت تركيزات كلا من السكريات الكلية والبرولين فى المجموع الخضرى لجميع الطرز الوراثة محل الدراسة والنامية تحت ظروف الإجهاد الملحى . علاوة على ذلك, زاد تركيز أيونات الصوديوم بينما نقص تركيز أيونات البوتاسيوم فى المجموع الخضرى لجميع الطرز.
تؤكد النتائج أن جمييزة ٧ وهجين جمييزة ١٠ x C.B. ٢٥٥ كانا أكثر مقاومة لظروف الإجهاد حتى مستوى ٦٠٠٠ مليجرام/لتر وكان جمييزة ٧ أكثر مقاومة مقارنة بجميع الطرز تحت الدراسة.

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