Effect of Salinity Stress on Physiological Changes in Winter and Spring Wheat

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Abstract: Salinity is a leading threat to crop growth throughout the world. Salt stress induces altered physiological processes and several inhibitory effects on the growth of cereals, including wheat (Triticum aestivum L.). In this study, we determined the effects of salinity on five spring and five winter wheat genotypes seedlings. We evaluated the salt stress on root and shoot growth attributes, i.e., root length (RL), shoot length (SL), the relative growth rate of root length (RGR-RL), and shoot length (RGR-SL). The ionic content of the leaves was also measured. Physiological traits were also assessed, including stomatal conductance (gs), chlorophyll content index (CCI), and light-adapted leaf chlorophyll fluorescence, i.e., the quantum yield of photosystem II (Fv’/Fm’), and instantaneous chlorophyll fluorescence (Ft). Physiological and growth performance under salt stress (0, 100, and 200 mol/L) were explored at the seedling stage. The analysis showed that spring wheat accumulated low Na+ and high K+ in leaf blades compared with winter wheat. Among the genotypes, Sakha 8, S-24, W4909, and W4910 performed better and had improved physiological attributes (gs, Fv’/Fm’, and Ft) and seedling growth traits (RL, SL, RGR-SL, and RGR-RL), which were strongly linked with proper Na+ and K+ discrimination in leaves and the CCI in leaves. The identified genotypes could represent valuable resources for genetic improvement programs to provide a greater understanding of plant tolerance to salt stress.

Keywords: chlorophyll fluorescence; Na+ efflux; salt tolerant; stomatal conductance

1. Introduction

Plants respond to environmental changes by altering metabolism, growth, and development. When climate changes are rapid, plants perceive them as stress. Among the abiotic stresses, those that most affect the productivity of agricultural crops are: extreme temperatures, osmotic stress, drought, and salinity. Many environmental conditions can lead to water stress in plants. For example, the high concentrations of salt in saline habitats
make it difficult for roots to absorb water from the soil. Drought reduces morphological traits such as the reduction of leaf size and vegetative growth, and physiological traits such as the reduction of photosynthesis and stomatal conductance, and alters the anatomical characteristic of the stem [1,2]. The results of these stresses induce an excessive production of reactive oxygen species (ROS), which cause extensive cell damage and inhibition of photosynthesis [3,4]. To these stresses are added the deficits of inorganic nutrients, the residues of chemicals such as herbicides and pesticides used in normal agricultural practice and heavy metals present in the earth’s crust that emanate from industrial activities [5–8].

Salinity is a major threat to agriculture, among other abiotic stresses; currently, more than 20% of agricultural land is affected by salinity, which is expanding day by day and already affects almost 954 million hectares of the world’s total land area [5,9,10]. Pakistan is located in arid and semi-arid regions where precipitation is scarce; therefore, the salt concentration accumulates in the rootzone [11]. The presence of high salt levels in the soil and water used for irrigation is one of the worrying factors for agriculture. For this reason, it is important to develop effective strategies to improve yield through salt tolerance. Salinity affects plant growth due to the toxicity of Na⁺ and decreases the uptake of essential nutrients such as calcium (Ca⁺²) and potassium (K⁺) [12]. A high salt concentration causes both osmotic and ionic stresses, which damages the photosynthetic apparatus and physiology, e.g., closes the stomata and reduces the leaf expansion rate [13,14]. Due to the high concentration of Na⁺ in saline soil, it causes water-stressed conditions that lead to decreased yield production worldwide [15,16]. Plant responses vary in salinity tolerance, as reflected in their different growth and physiological responses [6,17]. Indeed, there is potential for improving salt tolerance in cultivated species through selection and breeding. Among cereals, rice is the most salt-sensitive, and barley is the most salt-tolerant, whereas bread wheat is a moderately salt-tolerant crop.

Wheat (Triticum aestivum L.) is the most cultivated cereal in the world, after maize. Wheat is a staple food for more than one-third of the world’s people, supplying about 20% of total protein and daily calories [18,19]. Globally, wheat is cultivated on non-saline and saline soils, covering an area of approximately 214.79 million hectares [19,20]. Primary salinity refers to weathering of natural materials, while secondary salinity may occur due to anthropogenic activities. In low rainfall arid and semi-arid areas, improper irrigation practices are the main sources of secondary salinization [21]. Initial exposure to salinity leads to osmotic stress, which negatively affects plant growth due to the change in water content between cells which inhibits cell expansion and division with decreased stomatal opening and transpiration [10,22]. Long-term exposure to salinity causes plants to undergo ionic stress mainly due to the increase in sodium chloride concentration, which induces premature senescence, chlorosis, and necrosis in older leaves. Such changes negatively affect protein synthesis and photosynthetic activity [10]. In addition, the growth of the shoots is strongly reduced than that of the roots. This is because the decrease in leaf expansion compared with root growth reduces the water supply to the plant. Consequently, the soil moisture is conserved by avoiding an escalation of the salt concentration in the soil itself. Furthermore, high concentration levels of Na⁺ and/or Cl⁻ are toxic in the cell, negatively affecting the photosynthetic capacity. Consequently, there is a lower contribution of carbohydrates to young leaves, with a reduction in the growth rate of shoots [23]. Indeed, studies conducted on plants subjected to salt stress have shown different responses attributable to the plasticity of the plant genome being stress-related. These responses involve specific epigenetic modifications such as activation of transcription factors that modulate gene expression, which can influence the physiological processes of plants, compromise growth, and development [3,24,25]. The selection of salinity-tolerant plants could allow farmers to identify the genotypes best suited to salinized soils. However, progress in developing salt tolerance is limited by the genetic complexity of wheat. It is necessary to understand the mechanism of salinity tolerance in wheat genotypes. Many researchers and scientists have focused on exploring physiological mechanisms for developing the salt-tolerant
germplasm in wheat. Since wheat (*Triticum* spp.) is a major food crop, the development and identification of salt-tolerant wheat cultivars is an important research purpose.

Fluorescence of chlorophyll (e.g., \(F_{v}^{'}/F_{m}^{'}\), \(F_t\)) and gas exchange have been considered important physiological indicators for screening the tolerance of different cultures. There are two important exit photosystems (PSI and PSII) in the plant. PSII is found to be more prone to the hazardous effects of salinity [26]. Measuring chlorophyll fluorescence is a good indicator of salt effects in the photosynthetic apparatus [27]. Consequently, it is important to evaluate the relationship between the efficiency of PSII and CO\(_2\) assimilation in the leaves as the measurement of fluorescence detects the differences in the response of plants to abiotic stresses by evaluating their tolerance. The use of morphological traits, along with physiological tolerance and their relationship with salinity tolerance indices, are applicable and considered sufficient to be exploited as selection criteria in the breeding of salt-tolerant germplasm [27].

Wheat is also called a salt excluder, which means it mitigates salinity stress by excluding Na\(^+\) from the shoot as much as possible [28,29]. The capacity of bread wheat to exclude Na\(^+\) is much stronger than that of durum wheat genotypes [10]. Moreover, salt tolerance is a polygenic trait, and its expression can be influenced by genetic, environmental, and physiological factors. In fact, in the same species it is possible to select a salt-tolerant genotype [30], suggesting that this potential may be improved through conventional breeding approaches. Furthermore, little work has been carried out to examine physiological differences in spring and winter genotypes under salt stress. In the present study, we used five different spring wheat and five durum wheat genotypes to test the different responses and adaptations to salt stress. Some of these, such as S-24, were selected for their well-known salt-tolerance; therefore, it was used as reference one [31]. The genotypes used were evaluated for the effect of salinity on some key physiological and morphological traits. The identified genotypes could represent valuable resources for genetic improvement programs to provide greater understanding of plant tolerance to salt stress, supporting agricultural production on salinized soils irrigated with brackish water.

2. Results

2.1. Response of Wheat Genotypes against Different Salinity Levels

Significant differences \((p \leq 0.05)\) were recorded among the spring wheat genotypes (G), winter wheat genotypes, and the total wheat genotypes (Table 1) in terms of their interactions \((G+S)\) at different NaCl salt stress levels (S) for ionic, physiological, and seedling growth traits (Table 1).
Table 1. Mean square values ($p < 0.05$) for root length (RL), shoot length (SL), the relative growth rate of root length (RGR-RL), the relative growth rate of shoot length (RGR-SL), leaf Na$^+$ and K$^+$ concentration (mg g$^{-1}$ dry wt), K$^+$/Na$^+$ ratio, stomatal conductance $g_s$ (mol/m$^2$ s), maximum quantum efficiency of PSII ($Fv'/Fm'$), instantaneous chlorophyll fluorescence ($F_t$), chlorophyll content index (CCI) of spring wheat genotypes, spring wheat genotypes, and ten wheat genotypes grown under various NaCl stress levels.

| Source, SL | Df | RL  | SL  | RGR-RL | RGR-SL | Na$^+$ | K$^+$ | K$^+$/Na$^+$ | $Fv'/Fm'$ | $g_s$ | $F_t$ | Leaf Senescence | CCI  |
|-----------|----|-----|-----|--------|--------|--------|-------|-------------|-----------|-------|------|----------------|------|
| G         | 4  | 46.6*** |2.1* |0.399*** |0.02 n.s.|10.71** |12.41 n.s.|1.49 n.s.|0.002 n.s.|15.38*** |353972** |36.24 n.s.|126.39 ***|
| SL        | 1  | 136.1** |130.2*** |0.943*** |1.26*** |1292.7*** |1023.0*** |158.14*** |0.621*** |989.45*** |2.441E+1*** |7022.2*** |1886.48**|
| G*SL      | 3  | 10.1*** |3.5*** |0.034*** |0.033** |6.63 n.s.|15.42 n.s.|1.26 n.s.|0.004 n.s.|15.81*** |310460*** |20.7 n.s.|23.00***|
| G         | 4  | 111.5** |61.9*** |1.233*** |0.39*** |47.74*** |10.91 n.s.|2.66* |0.039*** |56.78*** |616640*** |236.8** |368.36***|
| SL        | 1  | 172.2*** |162.3*** |1.624*** |1.27*** |589.1*** |678.2*** |142.7*** |0.749*** |538.58*** |1.500E+1*** |4806.9*** |1252.4***|
| G*SL      | 3  | 2.96 n.s.|3.9 n.s.|0.075 n.s.|0.028 n.s.|15.6*** |30.62** |2.00 n.s.|0.008*** |18.63*** |196887*** |83.61 n.s.|37.55***|

Among the Five Winter Wheat Genotypes

| Source, SL | Df | RL  | SL  | RGR-RL | RGR-SL | Na$^+$ | K$^+$ | K$^+$/Na$^+$ | $Fv'/Fm'$ | $g_s$ | $F_t$ | Leaf Senescence | CCI  |
|-----------|----|-----|-----|--------|--------|--------|-------|-------------|-----------|-------|------|----------------|------|
| G         | 4  | 119.3** |58.9*** |0.984*** |0.36*** |41.46*** |74.6*** |2.86* |0.053*** |36.98*** |614174*** |131.5 n.s.|233.68***|
| SL        | 1  | 301.6** |291*** |2.511*** |2.53*** |1812.7*** |1678.4*** |300.26*** |1.77*** |1483.1*** |3.853E+1*** |11578*** |3081.1***|
| G*SL      | 3  | 6.6*** |3.5** |0.054*** |0.027 n.s.|17.58*** |23.0*** |1.52 n.s.|0.0085*** |20.29*** |322609*** |74.3 n.s.|33.33***|

G = genotypes, SL = salinity level; * = significant (<0.05), ** = more significant (<0.01), *** = highly significant (<0.00), n.s. = non-significant.
2.2. Value of % Control

Significant variation ($p \leq 0.05$) was observed between the spring wheat and winter wheat for ionic, seedling growth, and physiological traits (Figure 1a–l). The value of % control of ionic, physiological, and growth traits was decreased by increasing the salt stress (Figure 1a–l). Spring wheat showed a maximum value of % control for ionic ($\text{Na}^+$, $\text{K}^+$ and $\text{K}^+/$$\text{Na}^+$ ratio; Figure 1a–c), growth (RGR-SL), and physiological traits ($\text{gs}$, $Ft$, $QY$, and CCI; Figure 1i–l) compared with winter wheat.

![Figure 1](https://doi.org/10.3390/xxxxx)

**Figure 1.** Effect of salt stress on winter and spring wheat. Diagrams are based on value of % control (salt-treated/control=100) of $\text{Na}^+$ in leaves (a), $\text{K}^+$ in leaves (b), $\text{K}^+/$$\text{Na}^+$ ratio (c), leaf senescence % (d), shoot length (SL) (e), root length (RL) (f), relative growth rate of root length (RGR-RL) (g), relative growth rate of shoot length (RGR-SL) (h), stomatal conductance ($\text{gs}$) (i), instantaneous chlorophyll fluorescence ($Ft$; j), quantum yield of photosystem II ($Fv′/Fm′$; k), and chlorophyll content index (CCI; l).

2.3. Leaf $\text{Na}^+$ and $\text{K}^+$ Contents

Significant variations in $\text{Na}^+$ and $\text{K}^+$ content were identified between the leaves of genotypes treated with salinity stress (Table 1; Figure 2a,b). $\text{Na}^+$ concentration in leaves was increased in all wheat genotypes by increasing the salinity stress (Figure 2a,b). At 100 mol/L salinity stress, the minimum $\text{Na}^+$ content in leaves was found in W4909 (8.51 mg $g^{-1}$ DW; SW) followed by S-24 (8.59 mg $g^{-1}$ DW; SW), W4910 (9.09 mg $g^{-1}$ DW; SW), and Sakha 8 (11.03 mg $g^{-1}$ DW; SW), respectively (Figure 2a). Genotypes W4910 (11.40 mg $g^{-1}$ DW; SW), W4909 (15.48 mg $g^{-1}$ DW; SW), Sakha 8 (15.58 mg $g^{-1}$ DW; SW), and S-24 (11.40 mg $g^{-1}$ DW; SW) performed better at accumulating low leaf $\text{Na}^+$ at 200 mol/L salt stress and were considered to be salt tolerant. $\text{K}^+$ concentrations in leaves were decreased in all wheat genotypes by increasing the salinity stress (Figure 2b). The maximum $\text{K}^+$ concentration and $\text{K}^+/$$\text{Na}^+$ ratio were observed in Sakha 8 (23.67 mg $g^{-1}$ DW; SW), followed by S-24 (21.46 mg $g^{-1}$ DW; SW), W4909 (21.17 mg $g^{-1}$ DW; SW), and W4910 (19.31 mg $g^{-1}$ DW; SW), respectively (Figure 2b), at 100 mol/L. At the high salinity level (200 mol/L), W4910 (14.13 mg $g^{-1}$ DW; SW), W4909 (14.12 mg $g^{-1}$ DW; SW), Sakha 8 (213.94 mg $g^{-1}$ DW; SW), and S-24 (12.12 mg $g^{-1}$ DW; SW) genotypes performed better by accumulating more leaf $\text{K}^+$ (Figure 2b). Furthermore, the $\text{K}^+/$$\text{Na}^+$ ratio was also decreased by increasing the salt stress in both wheat genotypes (Figure 2c).
24 (12.12 mg g$^{-1}$ DW; SW) genotypes performed better by accumulating more leaf K$^+$ (Figure 2b). Furthermore, the K$^+$/Na$^+$ ratio was also decreased by increasing the salt stress in both wheat genotypes (Figure 2c).

**Figure 2.** Influence of salt stress on (a) Na$^+$ concentration in leaves, (b) K$^+$ concentration in leaves, and (c) K$^+$/Na$^+$ in leaves of wheat genotypes. Error bars indicate S.E. ($n = 3$). Different letters above the bars represent significant differences at the $p < 0.05$.

### 2.4. Shoot and Root Growth

Significant responses ($p < 0.05$) were observed for growth-related attributes among the genotypes for salt tolerance. Seedlings SL and RL were decreased by increasing the salt stress in all wheat genotypes (Figure 3a,b). Maximum SL were recorded in Sakha 8 (25.656 cm; SW), followed by S-24 (24.611 cm; SW), TXIID 3127 (22.611 cm; WW), W4909 (21.211 cm; SW), and W4910 (20.089 cm; SW), respectively (Figure 3a). The maximum RL was found in S-24 (18.917 cm; SW), followed by Sakha 8 (17.206 cm; SW), W4910 (15.741 cm; SW), and W4909 (15.663 cm; SW), respectively (Figure 3b).
Genotypes also showed significant variation in the RGR-SL and RGR-RL against salt stress level (Figures 3c and 4a, respectively). Among the wheat genotypes, the maximum value of RGR-SL values was found in S-24 (1.20; SW), followed by Sakha 8 (1.08; SW), W4909 (0.79; SW), TXIID 3127 (0.78; WW), and W4910 (0.66; SW; Figure 3c). Meanwhile, the maximum RGR-RL values were recorded in S-24 (0.85; SW), followed by Sakha 8 (0.68; SW), W4910 (0.60; SW), and W4909 (0.53; SW), respectively (Figure 4a).

Figure 3. Influence of salt stress on (a) shoot length, (b) root length, and (c) relative growth rate of shoot length (RGR-SL) on wheat genotypes. Error bars indicate S.E. (n = 3). Different letters above the bars represent significant differences at the p < 0.05.
Figure 4. Influence of salt stress on (a) relative growth rate of root length (RGR-RL), (b) chlorophyll content index (CCI), and (c) senescence of leaves of wheat genotypes. Error bars indicate S.E. (n = 3). Different letters above the bars represent significant differences at the p < 0.05.

2.5. Physiological Traits

The CCI in all wheat genotypes decreased with increasing levels of salt stress (Figure 4b). Among the genotypes, the maximum CCI values were observed in S-24 (29.22; SW), followed by Sakha 8 (24.74; SW), TX110D 2265 (24.25; WW), W4909 (19.84; SW), and TX11D 3134 (19.73; WW), respectively (Figure 4b).

A significant physiological response was observed among the genotypes for gs (Figure 5a). Genotype S-24 (15.98) showed the highest value of gs, followed by Sakha 8 (15.54), TX110D 2265 (13.91), W4909 (12.74), and W4910 (12.71; Figure 5a), whereas the minimum gs value was observed in PI 94341 (9.78) genotypes (Figure 5a).

The senescence of the leaves also increased in wheat genotypes by increasing the salinity stress, as shown in Figure 4c. The TX12M 4713 (26.22; WW) genotype showed the maximum leaf senescence, followed by PI 94341 (23.67; SW), TX110D 2265 (22.85; WW), TX11D 3134 (20.86; WW), and TXIID 3127 (20.21; WW), respectively (Figure 4c).
Figure 5. Influence of salt stress on (a) stomatal conductance ($g_s$), (b) efficiency of photosystem II (PSII), and (c) maximum quantum yield of photosystem II ($F_{v'}/F_{m'}$) of wheat genotypes. Error bars indicate S.E. ($n = 3$). Different letters above the bars represent significant differences at the $p < 0.05$.

Chlorophyll fluorescence, e.g., $F_{v'}/F_{m'}$ and $F_l$ values, declined under salt stress. There was a significant difference ($p < 0.05$) in chlorophyll fluorescence concerning genotypes, salinity level, and the interaction between genotypes × salinity level (Figure 5b,c).

A significant decrease in $F_l$ was observed in all genotypes by increasing the salt stress level (Figure 5b). The maximum values of $F_l$ were observed in W4909 (2700; SW), W4910 (2554.3; SW), S-24 (2547.3; SW), TX110D 2265 (2486.4; WW), and Sakha 8 (2464.3; SW), respectively (Figure 5b). $F_{v'}/F_{m'}$ decreased in wheat genotypes with increasing levels of salt stress (Figure 5c). The maximum values of $F_{v'}/F_{m'}$ were observed in S-24 (0.6056; SW), followed by W4909 9 (0.5778; SW), W4910 (0.5489; SW), Sakha 8 (0.5433; SW), and PI 94341 (0.4324; SW; Figure 5c).
2.6. Trait Correlations

A highly significant correlation was observed among the various traits under salt stress (Table 2). All traits were positively correlated except Na\(^+\) and the senescence of leaves, which were negatively correlated with all other attributes (Table 2). Na\(^+\) was positively correlated with the senescence of leaves (Table 2).

**Table 2.** Correlations among different ionic, growth, and physiological traits of wheat genotypes grown under various NaCl stress levels.

| Traits                          | Senescence of Leaf | Na\(^+\) in Leaf | K\(^+\) in Leaf | gs  | Ft  | Fv'/Fm' | Root Length |
|--------------------------------|--------------------|------------------|----------------|-----|-----|---------|-------------|
| Na\(^+\) in leaf               | 0.81               |                  |                |     |     |         |             |
| K\(^+\) in leaf                | −0.75              | −0.76 **         |                |     |     |         |             |
| gs                             | −0.76              | −0.73 **         | 0.75 **        |     |     |         |             |
| Ft                             | −0.83              | −0.87 **         | 0.77 **        | 0.82 ** |     |         |             |
| Fv'/Fm'                        | −0.85              | −0.88 **         | 0.79 **        | 0.84 ** | 0.93 ** |         |             |
| Root length                    | −0.75              | −0.74 **         | 0.66 **        | 0.76 ** | 0.75 ** | 0.84 ** |             |
| Shoot length                   | −0.67              | −0.65 **         | 0.58 **        | 0.75 ** | 0.69 ** | 0.70 ** | 0.83 **     |

**Table 3.**Correlations among different ionic, growth, and physiological traits of wheat genotypes grown under various NaCl stress levels.

| Traits                          | Senescence of Leaf | Na\(^+\) in Leaf | K\(^+\) in Leaf | gs  | Ft  | Fv'/Fm' | Root Length |
|--------------------------------|--------------------|------------------|----------------|-----|-----|---------|-------------|
| Na\(^+\) in leaf               | 0.88 **            |                  |                |     |     |         |             |
| K\(^+\) in leaf                | −0.78 **           | −0.85 **         |                |     |     |         |             |
| gs                             | −0.82 **           | −0.85 **         | 0.91 **        |     |     |         |             |
| Ft                             | −0.79 **           | −0.85 **         | 0.89 **        | 0.93 ** |     |         |             |
| Fv'/Fm'                        | −0.86 **           | −0.92 **         | 0.87 **        | 0.88 ** | 0.83 ** |         |             |
| Root length                    | −0.84 **           | −0.86 **         | 0.84 **        | 0.82 ** | 0.84 ** | 0.86 ** |             |
| Shoot length                   | −0.49 **           | −0.47 **         | 0.52 **        | 0.47 ** | 0.55 ** | 0.43 ** | 0.58 **     |

**Table 4.** Correlations among different ionic, growth, and physiological traits of wheat genotypes grown under various NaCl stress levels.

| Traits                          | Senescence of Leaf | Na\(^+\) in Leaf | K\(^+\) in Leaf | gs  | Ft  | Fv'/Fm' | Root Length |
|--------------------------------|--------------------|------------------|----------------|-----|-----|---------|-------------|
| Na\(^+\) in leaf               | 0.85 **            |                  |                |     |     |         |             |
| K\(^+\) in leaf                | −0.74 **           | −0.81 **         |                |     |     |         |             |
| gs                             | −0.79 **           | −0.81 **         | 0.82 **        |     |     |         |             |
| Ft                             | −0.81 **           | −0.86 **         | 0.83 **        | 0.89 ** |     |         |             |
| Fv'/Fm'                        | −0.79 **           | −0.86 **         | 0.85 **        | 0.82 ** | 0.84 ** |         |             |
| Root length                    | −0.70 **           | −0.74 **         | 0.76 **        | 0.72 ** | 0.73 ** | 0.87 ** |             |
| Shoot length                   | −0.54 **           | −0.56 **         | 0.61 **        | 0.59 ** | 0.60 ** | 0.66 ** | 0.80 **     |

\*\* Highly significant correlation.

3. Discussion

Salt stress induces a number of negative effects including physiological and biochemical changes in plants which manifest as a reduction in plant biomass and crop yield. Different plants have different tolerance levels, as do most cereals, including wheat [17]. In the present study, five genotypes of winter wheat and five genotypes of summer wheat were considered in order to carry out a comparative study on the salt tolerance of new genotypes, with the exception of the S-24 genotype already extensively studied by Ashraf [31], that is better suited to grow on salinized soils. Overall, the results showed different responses between summer and winter genotypes in terms of physiological traits (Figure 1). Under salinity stress, all genotypes accumulated a higher Na\(^+\) content in their leaves compared with non-stress conditions (Figure 2a). However, the sodium uptake in leaves was different in spring and winter wheat (Figure 1a). Winter wheat accumulated more sodium com-
pared with spring wheat (Figure 1a). Most of the salt-excluder genotypes were previously recognized as salt-tolerant by many scientists [32–34], which is further confirmed by this study’s results (Figures 1a and 2a). The salt-tolerant genotypes may possess a better ability to maintain low Na⁺ in their leaves, as reported by Elkelish et al. [35]. Saddiq et al. [32] reported that tolerant genotypes preferred to accumulate low Na and high K in their leaves, which was also observed in this study (Figure 1a,b and Figure 2a,b). Munns and Tester [10] reported that the removal of Na⁺ from the cytoplasm into the apoplast is due to the salt-inducible enzyme Na⁺/H⁺ antiporter located at the plasma membrane. Moreover, Na⁺ accumulation in wheat is controlled by *Nax1* and *Nax2* genes, located on 2A and 5A chromosomes, respectively [36,37], which are being used as molecular marker cultivars in a breeding program. *TNHX1*, *TNHX2*, and *TVP1* (vascular Na⁺/H⁺ antiporter) are responsible for improved seedling shoot growth by generating the pH gradient and facilitating sodium sequestration into values under salt stress [38]. Furthermore, salt-tolerant genotypes could have a sophisticated K⁺ regulation system, such as two-pore K⁺ channels and a shaker type, as described by Shabala and Pottosin [39], and on-selective cation channels, which aid the permeability of K⁺ and transporters (HKT, KUP/HAK/KT, and K⁺/H⁺). An inverse relationship exists between Na⁺ and K⁺ ions due to direct competition for ions in plant absorption [40].

The plants’ growth performance was also decreased under salt stress conditions (Figure 1e–g, Figure 3a–c, and Figure 4a). All genotypes had lower shoot lengths and root lengths in salt-stressed conditions compared with controls (Figures 3a–c and 4a). Nevertheless, spring wheat genotypes improved their RGR-SL compared with winter wheat genotypes (Figure 1g). Janmohammadi et al. [41] reported that winter wheat had a lower root length than spring wheat under abiotic stress (e.g., cold stress), ultimately affecting the wheat’s growth performance. NaCl stress induced a significant reduction in plant height, root length, and dry weight of roots and shoots in winter wheat [42]. Qiong et al. [43] reported that salinity significantly increased Na accumulation in winter wheat, which significantly reduced dry shoot weight and plant height. Na remarkably reduced the accumulation of K⁺, K⁺/Na⁺ ratio, as well soluble proteins and proline. Brestic et al. [44] reported that chlorophyll fluorescence is a more effective method for screening PSII thermostability in winter wheat genotypes. A high concentration of salt in the soil causes water stress, which leads to a significant decrease in the yield of many crops worldwide. Zivcak et al. [45] reported that the photosynthesis efficiency of PSI of winter wheat was decreased by increasing the water stress. Damage caused by salt stress was more prominent at the donor side, rather than the acceptor side of PSII [46]. Munns and Tester [10] reported that the accumulation of Na⁺ at toxic concentrations in the leaf negatively affects the photosynthetic mechanism, resulting in a lower intake of carbohydrates to the young leaf, reducing root and shoot growth. Thus, spring wheat was considered a tolerant crop, with a greater supply of assimilates from leaves to growing parts, e.g., root and shoot length (Figure 3a–b). This might be linked with prolonged retention of chlorophyll in the leaves of spring wheat (Figures 11 and 5b), which could stamp out Na⁺ from leaves, and thereby prevent Na⁺ from reaching toxic levels [41]. Poor performance in terms of growth might be linked with high cell membrane injury and senescence of leaves due to Na⁺ toxicity in growing embryos [4,39,47,48], and this suggestion is supported by the present study (Figures 1d and 4c).

Salt stress has an adverse impact on photosynthesis by destroying chlorophyll pigments and inhibiting the PSII activity. In this study, photosynthesis efficiency declined in both winter and spring wheat genotypes under salt-stress conditions (Figure 5b,c), but winter wheat was more affected (Figure 1j,k). In fact, under saline stress, stomatal closing results in a reduction in the photosynthetic rate of the plant. CO₂ assimilation in leaves, the efficiency of PSII, and their relationship allow fluorescence to be used to screen salt-tolerant germplasm against abiotic stresses [49]. Kanwal and his coworkers [26] evaluated the effects of salt stress on newly licensed wheat cultivars using gas exchange parameters and
chlorophyll fluorescence. The results reported a smaller reduction in plant biomass in cultivars S-24, Saher-226, and FSD-2008.

Measuring chlorophyll fluorescence is an excellent indicator to quantify salt-induced destruction in the photosynthetic apparatus [50]. Damage to photosystem II has been studied using this technique. Reactive oxygen species (ROS) degrade various proteins (a membrane linker protein, chlorophyll protein) that are necessary for the hooking of phycoobilisomes to thylakoids [35,46]. ROS burst destroys thylakoid membranes, resulting in modulations in membrane protein profiles, which leads to decreased activity of the oxygen-evolving complex (OEC) of PS II and increases the working of PS I. Salt-tolerant plants grown under a salt regime downregulate PS II in order to improve the quantum efficiency of excitation energy \( (Fv'/Fm') \) [50], as found in this study (Figure 1j,k). The maximum quantum yield of PSII, i.e., \( Fv'/Fm' \), is an important parameter to discriminate wheat genotypes. Of the different physiological attributes, stomatal conductance and the chlorophyll content index have been reported to be of prime importance in screening crop plants for salt tolerance. Generally, salt stress is known to cause a marked reduction in stomatal conductance and the chlorophyll content index have been reported to be of prime importance in screening crop plants for salt tolerance. Generally, salt stress is known to cause a marked reduction in stomatal conductance and the chlorophyll content index [35,51], as found in this study (Figure 1l). ROS are considered as the main source of structural damages under abiotic stresses such as drought, salinity, and heat [52]. ROS are highly cytotoxic and can seriously react with vital biomolecules such as lipids, proteins, nucleic acid, and disturb normal metabolic pathways [52,53]. It has been well documented that during salinity stress, somatically stressed plants reduced \( CO_2 \) assimilation due to the closing of stomatal pores, which generate ROS in the plant leaves [54]. In this study, spring wheat exhibited comparatively lower reductions in stomatal conductance and chlorophyll content index compared with winter wheat under salt stress (Figure 1i,l). The reduction in winter wheat might have been due to lower root water potential and the transport of plant hormone ABA from the root into different plant organs, thereby inducing stomatal closure [55]. Compared with spring wheat, winter wheat was more affected (Figure 1), which might be strongly linked with high Na levels in their leaves. The toxic concentration of Na\(^+\) in leaves encourages the reduction of stomatal conducntance in wheat by limiting photosynthesis efficiency [32,47]. Abiotic stress conditions caused by exposure to salinity, drought, heat, and waterlogging cause the stressed plant to produce ROS. The plant also produces antioxidants, flavonoids, and secondary metabolites that detoxify the ROS, thus protecting the plant from abnormal conditions, i.e., abiotic stress [52,53,56]. Therefore, tolerant genotypes prefer to accumulate high K\(^+\) instead of Na\(^+\) [57]. In this study, the influx of K\(^+\) was higher in spring wheat compared with winter wheat, helping to mitigate the salinity stress. Over time, salinity causes Na\(^+\) toxicity in leaves [10]. Therefore, controlling the transport of Na\(^+\) in the plant through the exclusion of Na\(^+\) from mesophyll cells is an important and reliable trait used to improve the salinity tolerance in many crops, i.e., durum wheat [58] and bread wheat [32,59].

4. Materials and Methods

4.1. Germplasm Collection

Seeds of five winter and five spring wheat genotypes (Table 3) were obtained from the USDA-ARS National Small Grains Collection, Aberdeen, ID, USA.

Table 3. Spring and winter wheat genotypes were used in this study.

| Spring Wheat | Winter Wheat |
|--------------|--------------|
| PI 94341     | TX12M 4713   |
| W4909        | TX11D 3154   |
| W4910        | TX11ID 3127  |
| S-24         | TX110D 2265  |
| Sakha 8      | TX12M 4637   |
4.2. Hydroponic Culture

A germplasm nursery (5 spring and 5 winter wheat genotypes) was raised in November 2015 in a growth chamber by sowing 50 seeds in 8 cm × 6 cm sand-filled polythene bags at 50% relative humidity and a light intensity of 400 mol m$^{-2}$s$^{-1}$. Plants were grown with a 14-h day length and with a 20 °C/17 °C day/night temperature cycle. Fifteen plants per genotype, replicated three times, were transplanted at the two-leaf stage into hydroponic tubs filled with 50 L of aerated half-strength Hoagland solution, which was changed fortnightly [60]. Seedling root length and shoot length were also recorded before being transplanted. The experimental design was a completely randomized design (CRD) factorial with three replications. Subsequently, commercial-grade salt was added in 50 mol/L increments twice daily to create different NaCl salt stress levels (0, 100, and 200 mol/L) to avoid osmotic shock.

4.3. Determination of Leaf Na$^+$ and K$^+$ Concentrations

After applying the salt in hydroponic culture, the expanded leaves that emerged under stress conditions were collected and put into the oven for drying. Leaf dry weight was determined. Dried leaves were put into falcon tubes filled with 25 mL of 1% HNO$_3$ solution for digestion on a hot plate at 85 °C for 4 h. One milliliter was taken from the digested solution, and a volume of 10 mL was prepared to measure the K$^+$ and Na$^+$ concentration in the leaf samples using a flame photometer (Sherwood, U.K., Model 360) [58,61].

4.4. Morphological Traits

After 10 days in a saline environment, the performance of seedlings was assessed based on morphological traits such as seedling root length (RL), shoot length (SL), the relative growth rate of root length (RGR-RL), and the relative growth rate of shoot length (RGR-SL). The relative growth rate was calculated using the formula of Gardener et al. [62].

\[
\text{RGR} = \frac{W_2 - W_1}{T_2 - T_1}
\]

where:
W$_1$ = root/shoot length at first harvest
W$_2$ = root/shoot length at second harvest
T$_2$ − T$_1$ = time interval between two harvests (10 days)

4.5. Chlorophyll Index and Stomatal Conductance

From the seedlings in a saline environment, the topmost fully expanded leaf was used to determine the chlorophyll index using a chlorophyll meter (Model Spad-502) [63]. Stomatal conductance was measured using a leaf photometer (Model Sc$^{-1}$).

4.6. Chlorophyll Fluorescence

The data for the chlorophyll fluorescence were recorded based on Baker [64] and Krame et al. [65] nomenclature. Chlorophyll fluorescence parameters, i.e., instantaneous chlorophyll fluorescence ($F_t$) and quantum yield of photosystem II (QY), were recorded by using the portable fluorescence meter, FluorPen FP 110 (Photon systems instruments, Czech Republic). The FluorPen FP 110 was equipped with a blue LED emitter (470 nm) optically filtered and precisely focused on delivering light intensities of up to 3000 µmol m$^{-2}$ s$^{-1}$ to measure plant tissues. QY is a measure of the Photosystem II efficiency. QY is equivalent to $F_v'/F_m'$ and $F_0$ is equivalent to $F_t$ in a light-adapted leaf. Quantum yield of PSII ($F_v'/F_m'$) was calculated as

\[
\frac{F_0'}{F_m'} = \frac{F_0'}{F_m'} - \frac{F_0'}{F_m'}
\]

where
$F_0'$ or $F_t'$: minimum fluorescence from a light-adapted leaf
$F_m'$: maximum fluorescence from a light-adapted leaf
Fv′: variable fluorescence from a light-adapted leaf (Fv′ = Fm′ − F0′)

4.7. Leaf Senescence

Three random plants in each treatment were tagged. At harvesting time, the total number of leaves and the number of green and senesced leaves were counted. A leaf was considered senesced if less than half of its area remained green.

4.8. Statistical Analysis

Quantitative observations of experiments were uploaded in SAS 9.4 (Texas A&M University, College Station, TX, USA) software to deduce the results in the form of variance analysis (ANOVA) for spring wheat genotypes, winter wheat genotypes, and all wheat genotypes. Data are presented in Table 1 with critical values to compare treatment means using the LSD test at the 5% probability level. The Statistix 8.1 package was also used to find correlations among the spring wheat genotypes, winter wheat genotypes, and all wheat genotypes for various growth, ionic, and physiological attributes (Table 2).

5. Conclusions

In this study, physiological comparisons of wheat genotypes under salt regimes Sakha 8, S-24, W4909, and W4910 performed better compared with PI 94341, TX12M 4713, and TX12M 4637, depicted by improved seedling growth, CCI, which was linked with better physiology traits, i.e., Fv′/Fm′; Ft and gs due to preferential K⁺ uptake and translocation to leaves. The identified plant material can be a source for more deeper insight into determining the genes responsible for enhanced salt tolerance in wheat.

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