Intraspecific genetic variation within and between improved cultivars and landraces of durum wheat in germination and root architectural traits under saline conditions

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Abstract

Although the effect of salinity on plants is well established, yet evaluating genetic variation in yield-related traits under salinity stress remains useful. This study was conducted to investigate the genetic variation within and between two subgroups of durum wheat in Jordan, i.e., cultivars and landraces, in germination and root architectural traits at three salinity concentrations (50, 100 and 150 mM NaCl). The results indicate that salinity significantly impacted most traits investigated. A multivariate discriminant analysis (DA) revealed large variations (~85%) between genotypes, with significant ranking in maximum root length, total root length, seminal root length, coleoptile length, germination percentage, and total and seminal root number over the three salinities. Within the cultivar subgroup, salinity significantly influenced the germination percentage and most root traits of durum genotypes, with variable magnitudes depending on NaCl concentration and within-variety intraspecific genetic variation (ANOVARs; P<0.05). However, within the landrace subgroup, only a salinity of 150 mM NaCl significantly affected the studied traits, and the effect of salinity on germination percentage was highly genotype dependent. Additionally, the durum genotypes in the landrace subgroup were more affected by salinity and showed more genetic variation than those in the cultivar subgroup.

Introduction

Durum wheat (Triticum turgidum L. spp. durum Desf.) is a tetraploid grass with nutritional value for humans. This crop was originally domesticated in the Fertile Crescent and then spread, over several generations, to diverse areas worldwide, where it has been subjected to natural and artificial selection processes. Domestication and later breeding have resulted in the development of varied local landraces and modern cultivars. Intraspecific genetic variation in crop plants is believed to affect many phenotypic traits related to growth and development.1 Therefore, understanding the genetic variation underlying the varied phenotypes of a crop under environmental stress is important.

Durum wheat is grown in Jordan as it is a major constituent of various traditional dishes including Kibbeh, Bulgur, Maftoul, and Tabbouleh. However, growers of durum wheat in Jordan, as in many arid and semi-arid regions worldwide, suffer from reduced productivity in most cereal crops due to the prevalence of soil salinity.2 Soil salinity limits crop productivity by influencing key growth and developmental processes, such as the germination process; by damaging living cells through ionic toxicity, chiefly sodium (Na+) and chloride (Cl-); and by reducing water availability due to osmotic effects.3,4

Seed germination is an important determinant of the successful establishment of a crop and can be greatly inhibited by stress factors that delay germination or retard seed viability.5,6 In addition to germination, salinity may affect other yield-related traits of crops, including the architectural traits of the root system, such as total root length, number of seminal roots, and coleoptile length.7,8-10 Germination and root traits are believed to play an important role in shaping the tolerance of a crop to a stressful environment and thus can be utilized as suitable traits for selection in crop improvement.9,11,12

Although the response of plants to NaCl stress is known, evaluating intraspecific genetic variation in yield-related traits in response to stress factors remains important, as it can be useful in screening for genotypes with improved tolerance to saline fields.13,14 However, selecting salt-tolerant genotypes by evaluating genotypic variation in the field has been proven to be rather misleading: first, because the salinity effect is difficult to disentangle from the many other effects of the biotic and abiotic factors encountered, and second, because of the required labor and resources, i.e., time and consumables, especially in long-term experiments with a large number of genotypes.15 Alternatively, short-term greenhouse experiments of a few days to a week in duration have been favored to identify salt-tolerant genotypes by comparing differential responses in controlled saline and non-saline conditions.

In this study, the intraspecific genetic variation in 13 yield-related traits of two groups (3 cultivar genotypes and 3 landrace genotypes) of durum wheat that are mainly grown in Jordan is evaluated to advise local durum wheat growers of possible salt-tolerant genotypes and to better understand the genetic influences underlying these yield-related traits in saline conditions.

Materials and Methods

Plant material and growth conditions

The seeds of the studied genotypes of durum wheat were obtained from the National Center for Agricultural Research and Extension (NCARE) of Jordan. These genotypes were of two groups: cultivars (Deir alla 6, Acsad 65, and Sham 1) and local landraces (Um-qais, Hourani 27, and Sham 1) and their countries of origin and pedigrees (Table 1). Initially, the seeds were treated with 1% (w/v) sodium hypochlorite solution for sterilization, and then thoroughly washed with distilled water. The sterilized seeds were then placed into Petri dishes (100 mm in diameter) containing two pieces of filter paper and then moistened with distilled water or with one of the NaCl solutions (50, 100 and 200 mM) with electrical conductivities of 4.2 dS m⁻¹, 8.3 dS m⁻¹ and 12.1 dS m⁻¹, respectively. The dishes of treated seeds were sealed with Parafilm and arranged randomly in a growth chamber at 150 mm x 150 mm x 50 mm at a temperature of 27±2°C and a relative humidity of 65±2% under natural photoperiod conditions.

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25°C. Germination was evaluated daily and was judged to occur if a radicle length of approximately 2 mm was observed. Germination percentage (G%) was calculated as follows: G% = 100 (GS/TS), where GS = germinated seeds and TS = total seeds incubated. To measure changes in root traits, digital photographs of the root architecture were taken for total root length, maximum root length, seminal root length, and coleoptile length, and the images were converted to high-contrast black-and-white pictures using Adobe Photoshop software. These images were then used to measure the root lengths with the help of ImageJ (https://imagej.nih.gov/ij/). Seminal root numbers of the studied genotypes were counted for 10 randomly sampled germinated seeds. For the estimation of chlorophyll content, approximately 0.2 g of the youngest leaf tissues was ground in 10 ml of 80% acetone with a mortar and pestle, and then the extract was transferred into Falcon tubes and centrifuged for 15 minutes at 15,000 rpm. The supernatant of the centrifuged solution was then transferred into new tubes and kept on ice. The absorbance of the supernatant was measured at OD_{645} and OD_{663} nm using a Spectrophotometer (Biochrom Ltd, Cambridge, UK). The measured absorbance were then used as inputs to estimate the concentrations of total chlorophyll, chlorophyll a, and chlorophyll b. Stomatal conductance was measured using a Delta-T AP4 cycling diffusion porometer (Delta-T Devices Ltd, Burwell, UK) using the uppermost fully expanded leaves.

Data analysis

Data analysis for germination and morphological traits was based on the means of four replicates. The measured traits were first subjected to modeling with a multivariate discriminant analysis (DA) to identify the traits most affected by salinity and to summarize the resulting responses to salinity treatments. In the DA model, salinity concentrations of 50, 100 and 150 mM NaCl were used as predefined groups, and the trait estimates were entered into the model simultaneously in a standard procedure. The ranking of the studied traits according to their relative importance by genotype was based upon F and P values. Moreover, the effect of each salinity level on the studied traits was tested with a one-way analysis of variance (ANOVA), and two- and three-way ANOVAs were used to examine the possible interaction effects of salinity according to within- and between-group intraspecific variation. Data were analyzed using IBM SPSS (SPSS Inc., Chicago, USA).

Table 1. Name, origin and pedigree of studied wheat genotypes.

| Genotype identification name | Country of origin | Pedigree |
|-----------------------------|------------------|----------|
| Um-qais                     | Jordan           | Landrace |
| Noorseh                     | Jordan           | Landrace |
| Hourani 27                  | Jordan           | Landrace |
| Deir alla 6                 | Jordan           | YEMEN X Cr's'X Plc's'7 Gly's CM 13857-D65-D11 |
| Ascad 65                    | ICARDA/Syria     | GERARDO-VZ-469/3/JORI-1/ND-61-130/LEEDS |
| Sham 1                      | ICARDA/Syria     | PELICANO/RUFF/GAVOTA/ROLETTE; PELICANO(SIB)/(SIB)RUFF/GAVOTA (SIB)/(SIB)ROLETTE |

*ICARDA, International Center for Agricultural Research in the Dry Areas.

Figure 1. Genotypic variation in germination percentage and root architectural traits in three cultivars of durum wheat at salinities of 100 and 150 mM NaCl. Bars represent the means ± standard error of four individuals per genotype. Different letters indicate significant differences between genotypes at P≤0.05.
Results and Discussion

The results of this study illustrate that, depending on NaCl concentration and on the intraspecific genetic variation of the studied genotypes, salinity negatively affects the germination and root traits of durum wheat. A multivariate DA of all studied traits resulted in two functions with a combined prediction of salinity effect of approximately 60% across the three salinity concentrations, i.e., 50, 100, and 150 mM NaCl (Table 2). In particular, the prediction of the DA model was 84% for the 150 mM NaCl concentration and 54% and 42% for the 50 and 100 mM NaCl concentrations, respectively, implying that genotypic differences in germination and root architectural traits are best assessed at the highest level of salinity, i.e., 150 mM NaCl.17 To determine the ranking of the traits that best discriminated the durum genotypes, two DA canonical axes were obtained; the first explained a total variation of approximately 85% with significant loadings for maximum root length, total root length, seminal root length, coleoptile length, germination percentage, and total and seminal root numbers (Table 2). In DA, a greater magnitude of the discriminant function coefficient (regardless of sign) indicates a greater contribution of that trait to the total observed variation. The results of the DA indicate that the studied traits are suitable metrics for elucidating salinity effects in durum wheat, as they show significant total variation among wheat genotypes, and this variation may suggest their importance in screening for salt-tolerant and salt-sensitive genotypes.14

Furthermore, to see if these traits were similarly affected by different salinity concentrations, one-way ANOVAs were conducted. According to this analysis, a salinity of 150 mM NaCl, relative to control (0 mM NaCl), significantly decreased the total, maximum, and seminal axile root lengths and the coleoptile length in the Deir alla 6 genotype, while salinities of 50 and 100 mM NaCl did not affect any of the traits studied (Table 3). For the genotype Sham 1, a salinity of 150 mM NaCl also resulted in a decrease in germination percentage and a reduction in total and maximum root lengths, while a salinity of 100 mM NaCl reduced total root number and maximum root length. The Acsad 65 genotype was found to be the least affected, as only the coleoptile length was reduced and only by a salinity of 150 mM NaCl (Table 3). The effects of salinity on germination percentage and root traits are best attributed to the reduced water availability caused by osmotic stress rather than to ionic stress.2,9 Additionally, these results reveal differences

| Traits                  | Canonical variable 1 | Canonical variable 2 |
|-------------------------|----------------------|----------------------|
| Germination             | 0.192                | -0.247               |
| Number of roots         | -0.407               | 0.602                |
| Root length             | -0.700               | -0.138               |
| Coleoptile length       | -0.146               | 0.376                |
| Number of seminal roots | 0.189                | 0.524                |
| Seminal root length     | -0.487               | -0.720               |
| Maximum root length     | 2.148                | 0.189                |
| Percentage of variance  | 84.60                | 15.4                 |
| Canonical correlation    | 0.520                | 0.252                |
| Predicted classification |                      |                      |
| 50 mM NaCl              | 54.2%                |                      |
| 100 mM NaCl             | 41.7%                |                      |
| 150 mM NaCl             | 83.3%                |                      |

| Salinity (NaCl mM) | Germination (%) | Total root number | Seminal root number | Maximum root length | Coleoptile length |
|--------------------|-----------------|-------------------|--------------------|---------------------|------------------|
| Un-gae              | 100             | 95.7              | 21.2               | 18.4                | 70               |
| Nourshe             | 50              | 93.5              | 20.7               | 18.1                | 65               |
| Houreni ZF          | 150             | 88.3              | 19.2               | 16.9                | 58               |

Figure 3. Variation between the landrace and cultivar subgroups of durum wheat in germination percentage and root architectural traits when grown on media containing 50 mM, 100 mM and 150 mM NaCl. Points represent the means ± standard error of four individuals per genotype. Asterisks (*) indicate significant differences at P≤0.05; (**) at <0.01; and (***) at <0.001.
tial responses between genotypes in the cultivar subgroup in germination and most root architectural traits. Namely, the germination percentage, the root traits and coleoptile length were profoundly affected especially in sham1 followed by Acsad 65 and lessly in Dear allah (Figure 1). Accordingly, the genotypes can be ranked by the number of traits indicating tolerance as salt-tolerant > salt-sensitive as follows: Deir alla 6 > Acsad 65 > Sham 1. Therefore, Deir alla 6 seems to be a salt-tolerant genotype, and Sham 1 seems to be a salt-sensitive genotype. These results suggest the suitability of these traits for classifying the studied genotypes into salt-tolerant and salt-sensitive genotypes.

To discover whether the genetic variation of genotypes in the cultivar subgroup interactively underlies variation in germination and root architectural, two-way ANOVAs for the effects of salinity, genotype, and their interaction were performed (Table 4). The analysis showed that total, seminal, and maximum root lengths and coleoptile length were all associated with intraspecific genetic variation and that the combined effect of genotype and salinity was not significant for any trait (two-way ANOVA; P>0.05).

However, in the landrace subgroup, a one-way ANOVA showed that a salinity of 150 mM NaCl differentially affected all traits studied except for total root number (Table 3). Specifically, the germination percentage and total and maximum root lengths were significantly affected in both Um-qais and Hourani 27, while the number of seminal roots, length of seminal root and coleoptile length were also affected in Um-qais. Based on their differential responses to salinity, the genotypes in the landrace subgroup can be ranked as salt-tolerant > salt-sensitive as follows: Um-qais > Hourani 27 > Noorseh (Figure 2).

Unlike the genotypes in the cultivar subgroup, all studied traits in the landrace subgroup were found to be associated with genetic variation, and the effect of salinity on germination percentage was highly genotype-dependent, as a significant salinity x genotype interaction was found (Table 5). Additionally, durum genotypes in the landrace subgroup were found to be more affected by salinity and showed a larger genetic variation than cultivar genotypes, especially in germination and root traits (Figure 3). This larger genetic variation in landraces compared with improved varieties could be attributed to the fact that landraces are typically more adapted to varied stressful environments.

### Table 3: Mean estimates, based on four independent replicates, of germination percentage and root architectural traits of the studied durum wheat varieties (cultivars and landraces) under three salinity concentrations (50, 100, and 150 mM NaCl) and their variations relative to the control (0 mM NaCl) as measured using a one-way ANOVA.

| Variety type | Dependent variable | 50 | 100 | 150 |
|--------------|--------------------|----|-----|-----|
| **Cultivars** |                    |    |     |     |
| Deir alla 6  | Germination percentage | 73.00 | 85.00 | 85.00 |
|              | Number of roots     | 4.50 | 5.00 | 5.00 |
|              | Number of seminal roots | 2.00 | 2.25 | 2.75 |
|              | Total root length   | 17.96 | 18.77 | 11.20** |
|              | Seminal root length | 7.08 | 8.16 | 5.97* |
|              | Maximum root length | 5.19 | 5.61 | 3.25* |
|              | Coleoptile length   | 3.04 | 3.27 | 2.28* |
| Acsad 65     | Germination percentage | 76.00 | 77.00 | 81.00 |
|              | Number of roots     | 4.50 | 3.50 | 4.50 |
|              | Number of seminal roots | 2.50 | 2.25 | 2.75 |
|              | Total root length   | 13.55 | 9.96 | 9.47 |
|              | Seminal root length | 7.09 | 5.32 | 4.92 |
|              | Maximum root length | 4.89  | 3.82 | 2.98 |
|              | Coleoptile length   | 3.47 | 3.05 | 2.10*** |
| **Landraces** |                    |    |     |     |
| Sham 1       | Germination percentage | 90.00 | 85.00 | 56.01* |
|              | Number of roots     | 4.25 | 3.50* | 4.00 |
|              | Number of seminal roots | 2.00 | 1.50 | 2.00 |
|              | Total root length   | 11.99 | 9.67 | 4.94* |
|              | Seminal root length | 3.91 | 4.07 | 2.32 |
|              | Maximum root length | 3.38 | 3.30* | 1.76** |
|              | Coleoptile length   | 2.31 | 1.65 | 1.28 |
| Um-qais      | Germination percentage | 88.00 | 87.00 | 79.00** |
|              | Number of roots     | 4.5 | 3.50 | 3.00 |
|              | Number of seminal roots | 2.50 | 2.00* | 2.00* |
|              | Total root length   | 8.41 | 6.77 | 3.12*** |
|              | Seminal root length | 5.53 | 4.58 | 1.85*** |
|              | Maximum root length | 3.11 | 2.51 | 1.27*** |
|              | Coleoptile length   | 1.78 | 1.09 | 0.82** |
| Noorseh      | Germination percentage | 43.00 | 45.00 | 39.00 |
|              | Number of roots     | 4.00 | 2.50 | 2.00 |
|              | Number of seminal roots | 2.25 | 1.75 | 1.00 |
|              | Total root length   | 4.74 | 4.55 | 1.83 |
|              | Seminal root length | 2.31 | 2.80 | 0.88 |
|              | Maximum root length | 1.97 | 1.74 | 0.62 |
|              | Coleoptile length   | 0.95 | 0.70 | 0.48 |
| Hourani 27   | Germination percentage | 78.00 | 57.00 | 21.00** |
|              | Number of roots     | 5.25 | 4.75 | 3.50 |
|              | Number of seminal roots | 2.75 | 2.00 | 1.50 |
|              | Total root length   | 15.19 | 13.12 | 6.11*** |
|              | Seminal root length | 4.77 | 3.97 | 1.70 |
|              | Maximum root length | 4.89 | 3.74 | 2.00* |
|              | Coleoptile length   | 4.15 | 2.74 | 2.12 |

*S Significant at P<0.05; ** significant at P<0.01.

### Table 4: Summary of the P values of two-way ANOVAs for the effects of salinity (S), genotype (G), and their interaction (S + G) on the germination and root architectural traits of durum wheat cultivars.

| Traits / source of variation | S     | G     | S + G |
|------------------------------|-------|-------|-------|
| Germination percentage       | 0.45  | 0.83  | 0.064 |
| Number of roots              | 0.44  | 0.09  | 0.81  |
| Number of seminal roots      | 0.35  | 0.15  | 0.90  |
| Root length                  | 0.04  | 0.01  | 0.78  |
| Seminal root length          | 0.27  | 0.008 | 0.87  |
| Maximum root length          | 0.009 | 0.020 | 0.842 |
| Coleoptile length            | 0.017 | 0.004 | 0.827 |
Conclusions

Because germination and root traits in wheat are positively related to increased grain number and productivity in the field, the results of this study maybe suggest their suitability as target phenotypes for genetic improvement to increase wheat productivity under saline soil conditions. Further research should investigate the genetic components, i.e. identify or clone the genes controlling the salt response of the salt-tolerant genotype and the salt-sensitive genotype of wheat.19

Table 5. Summary of the P values of two-way ANOVAs for the effects of salinity (S), genotype (G), and their interaction (S + G) on the germination and root architectural traits of durum wheat landraces.

| Traits / source of variation | S              | G              | S + G |
|------------------------------|----------------|----------------|-------|
| Germination percentage       | 0.000          | 0.000          | 0.002 |
| Number of roots              | 0.018          | 0.026          | 0.953 |
| Number of seminal roots      | 0.008          | 0.214          | 0.773 |
| Root length                  | 0.000          | 0.000          | 0.283 |
| Seminal root length          | 0.000          | 0.010          | 0.660 |
| Maximum root length          | 0.000          | 0.000          | 0.651 |
| Coleoptile length            | 0.179          | 0.002          | 0.872 |

Figure 2. Genotypic variation in germination percentage and root architectural traits in three landraces of durum wheat at salinities of 100 and 150 mM NaCl. Bars represent the means ± standard error of four individuals per genotype. Different letters indicate significant differences between genotypes at P≤0.05.

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