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## Effect of NaCl Stress on Chlorophyll Biosynthesis, Proline, Lipid Peroxidation and Antioxidative Enzymes in Leaves of Salt-Tolerant and Salt-Sensitive Barley Cultivars

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### ABSTRACT

In this research, the salt tolerance of salt-tolerant (Avcı-2002) and salt-sensitive (Tokak 157/37) barley cultivars were analyzed. The seedlings were grown in a hydroponic culture containing 0, 100, 200 and 300 mM NaCl. The decrease in dry weight was more significant in Tokak 157/37 than in Avcı-2002. On the other hand, no significant correlations were found between the altered  $\delta$ -aminolevulinic acid dehydratase (ALAD) activity and the chlorophyll content. The chlorophyll and carotenoid contents in leaves of Avcı-2002 decreased significantly at only 300 mM NaCl, whereas the pigment contents of Tokak 157/37 decreased at all NaCl concentrations. The increase in proline content was significantly higher in Avcı-2002 than in Tokak 157/37 at 300 mM NaCl. The lipid peroxidation level measured in terms of malondialdehyde (MDA) content was higher in Tokak 157/37 than in Avcı-2002. Salt stress caused significant increases in superoxide dismutase (SOD), guaiacol peroxidase (POD), ascorbate peroxidase (APX) and catalase (CAT) activity. However, Avcı-2002 had relatively lower SOD and CAT activities and higher POD and APX activities than Tokak 157/37. As a result, dry weight, pigments content, proline content, lipid peroxidation and activities of POD and APX results are in good correlation with supporting Avcı-2002 being relatively salt-tolerant.

Keywords: Barley; Chlorophyll biosynthesis; Proline; Lipid peroxidation; Antioxidant enzymes; Salt tolerance

## Tuza-Toleranslı ve Tuza-Hassas Arpa Çeşitlerinin Yapraklarında Klorofil Biyosentezi, Prolin, Lipit Peroksidasyonu ve Antioksidant Enzimler Üzerine NaCl Stresinin Etkisi

### ESER BİLGİSİ

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**ÖZET**

Bu araştırmada, tuza toleranslı (Avcı-2002) ve tuza hassas (Tokak 157/37) arpa çeşitlerinin tuza toleransları analiz edilmiştir. Fideler 0, 100, 200 ve 300 mM NaCl içeren hidroponik kültür ortamında büyütülmüştür. Kuru ağırlıktaki azalma Avcı-2002'ye göre Tokak 157/37 çeşidinde daha önemli bulunmuştur. Diğer taraftan,  $\delta$ -aminolevülinik asit dehidrataz (ALAD) aktivitesi ve klorofil içeriği arasında önemli korelasyon bulunmamıştır. Avcı-2002 çeşidinin yapraklarında klorofil ve karotenoid içeriği yalnızca 300 mM NaCl'de önemli düzeyde azalırken, Tokak 157/37 çeşidinin pigment içeriği tüm NaCl konsantrasyonlarında azalmıştır. 300 mM NaCl konsantrasyonunda, prolin içeriğindeki artış Tokak 157/37 çeşidine göre Avcı-2002 çeşidinde önemli düzeyde daha yüksek bulunmuştur. Malondialdehit (MDA) içeriği ile belirlenmiş lipit peroksidasyon seviyesi Avcı-2002'ye göre Tokak 157/37 çeşidinde daha yüksek bulunmuştur. Tuz stresi süperoksit dismutaz (SOD), guaiakol peroksidaz (POD), askorbat peroksidaz (APX) ve katalaz (CAT) aktivitesinde önemli artışa neden olmuştur. Bununla birlikte, Avcı-2002 çeşidinin Tokak 157/37 çeşidinde göre nispeten daha düşük SOD ve CAT aktivitelerine ve daha yüksek POD ve APX aktivitelerine sahip olduğu belirlenmiştir. Sonuç olarak, kuru ağırlık, pigment içeriği, prolin içeriği, lipit peroksidasyonu ile POD ve APX aktivite sonuçları nispeten tuza toleranslı Avcı-2002 ile iyi bir korelasyon göstermiştir.

Anahtar Kelimeler: Arpa; Klorofil biyosentezi; Prolin; Lipit peroksidasyonu; Antioksidant enzimler; Tuza tolerans

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**1. Introduction**

Salinity affects almost every aspect of the physiology and biochemistry of plants and significantly reduces yield. Salt stress leads to a decreased efficiency of photosynthesis and is known to influence the chlorophyll content of plant leaves (Meloni et al 2003). The decrease of chlorophyll synthesis may be due to a decrease of  $\delta$ -aminolevulinic acid dehydratase (ALAD) activity under environmental stress (Vajpayee et al 2000), but there is no clear information on the behavior of this enzyme in leaves under salt stress conditions. Proline accumulation is one of the adaptations of plants to salinity. It has also been widely advocated that proline accumulation uses as parameter of selection for salt stress tolerance (Ramanjulu & Sudhakar 2001).

Salinity stress induce enhanced production of the reactive oxygen species (ROS) such as superoxide radicals ( $O_2^{\cdot-}$ ), singlet oxygen ( $^1O_2$ ), hydroxyl radicals ( $\cdot OH$ ) and hydrogen peroxide ( $H_2O_2$ ) in plants (Misra & Gupta 2006; Kholová et al 2010). Lipid peroxidation, induced by ROS, is also important in membrane deterioration (Demiral & Turkan 2005). Thus, cell membrane stability has widely been utilized to differentiate salt-tolerant and salt-sensitive cultivars (Meloni et al 2003). Salinity promotes oxidative stress and plants

with high constitutive and induced antioxidant enzymes have better tolerance to damage (Parida & Das 2005). The generation of ROS is limited or scavenged by antioxidant enzymes such as superoxide dismutase (SOD), guaiacol peroxidase (POD), ascorbate peroxidase (APX) and catalase (CAT) (Foyer & Noctor 2003). Comparison of the responses of cultivars and/or related species that exhibit differential sensitivity to salt stress showed a correlation between salt tolerance and increased activity of the antioxidant system (Neto et al 2006; Athar et al 2008).

Barley is considered highly salt tolerant and can grow in the areas with elevated salt contents (Royo & Aragüés 1999). Existence of differences in salt tolerance not only amongst different species, but also within certain species (Munns & Tester 2008) offers an opportunity for identifying and developing salt-tolerant genotypes. The aim of this study is to evaluate the comparative effects of different concentrations of NaCl salinity on growth, chlorophyll biosynthesis, proline content, lipid peroxidation and SOD, POD, APX and CAT activity of salt-tolerant (Avcı-2002) and salt-sensitive (Tokak 157/37) barley cultivars at early seedling stage. Comparison of these parameters in the present barley cultivars grown in Turkey may

be helpful in developing a better understanding and provide additional information on the mechanisms of salt tolerance.

## 2. Material and Methods

### 2.1. Plant materials, growth and treatment conditions

Based on our preliminary studies, two barley cultivars (relatively salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37) were chosen from twelve barley cultivars for their different salt tolerance at early seedling stage (Yıldız & Terzi 2011). Healthy uniform seeds of barley cultivars were sterilized in 2% (w/v) sodium hypochlorite (NaOCl) solution for 20 min. Seeds were germinated at 25°C in dark on two filter papers in germination cups wetted with distilled water for three days. Three-day-old etiolated seedlings were grown in a jar including half strength Hewitt's nutrient solution at a constant temperature regime of 25±1°C, 60% humidity and at 13.000 lux light intensity, 12 h photoperiod for 24 h. Four-day-old seedlings were grown hydroponically in half strength Hewitt's nutrient solutions without NaCl or with 100, 200 and 300 mM NaCl for six days. At the end of 10<sup>th</sup> day, the first leaves of control and NaCl-treated barley seedlings were sampled for physiological and biochemical analyses.

### 2.2. Growth parameter

NaCl phytotoxicity was assessed in the terms of shoot and root dry weights for the treated and untreated control seedlings. A total of 10 plants from each treatment were sampled randomly at the seventh day of NaCl treatment, and shoots and roots were separated. The shoot and root tissues were dried in an oven at 80°C for 48 h. The dried samples were weighed immediately after the removal from the oven until biomass became stable. The dry weights were expressed as percentage of control.

### 2.3. Estimation of ALAD activity

The extraction of  $\delta$ -aminolevulinic acid dehydratase (ALAD) from the excised leaf tissue was carried

out according to Naito et al (1980) as described in Vajpayee et al (2000). Fresh leaf samples (1 g) from control and NaCl-treated seedlings were homogenized in 0.05 M Tris-HCl buffer (pH 8.2) containing 0.1 mM dithioeritol (DTT, 0.5 mL g<sup>-1</sup> FW) in a pre-chilled mortar. The homogenate was filtered through four layers of cheese cloth. The filtrate was centrifuged at 10000 ×g for 1 h at 4°C and the resultant supernatant was assayed for ALAD activity according to the method of Schneider (1970). One mL of supernatant was mixed with 0.27 mL of 1 mg mL<sup>-1</sup>  $\delta$ -aminolevulinic acid (ALA) and 1.35 mL of 0.05 M Tris-HCl buffer (pH 8.2) containing 0.1 mM DTT and 0.08 mL of 0.2 M MgCl<sub>2</sub>. The reaction mixture was incubated at 37°C for 2.5 h. The reaction was terminated by addition of 0.3 mL of 3.0 M trichloroacetic acid. After cooling, samples were centrifuged at 2000 ×g for 10 min. One mL of supernatant was mixed with an equal amount of Ehrlich's reagent, and absorbance of the mixture was determined at 555 nm after 10 min against zero time blank. The concentration of porphobilinogen (PBG) was calculated according to Mauzerall and Granick (1956) and the ALAD activity is expressed as nmol PBG 2.5 h<sup>-1</sup> mg<sup>-1</sup> protein at 37°C. In addition, protein concentration of leaf crude extract was determined according to the method of Bradford (1976).

### 2.4. Chlorophyll and carotenoid pigments content

Total chlorophyll (Chl *a* + *b*) and carotenoid pigments in the leaf tissues (100 mg) of control and NaCl-treated seedlings were extracted in methanol and determined by the method of Wellburn (1994).

### 2.5. Proline content

Proline content of leaf tissues was estimated spectrophotometrically following the ninhydrin method described by Bates et al (1973) with minor modification. Powdered frozen tissues (200 mg) were homogenized in 3% of sulphosalicylic acid to precipitate protein. Samples were mixed, centrifuged at 8000 ×g for 15 min, and the supernatants were treated with glacial acetic acid and ninhydrin reagent [3% (w/v) ninhydrin in 60%

(v/v) 6 M phosphoric acid] for 1 h at 90°C. After ice-cooling, the products were extracted with toluene by vortex mixing, and then the absorbance of the upper (toluene) phase was determined at 520 nm against a zero time blank. Proline concentrations were calculated using proline standards (0-100 µg mL<sup>-1</sup>) in identical manner.

### 2.6. Determination of lipid peroxidation

The level of lipid peroxidation was measured by estimating malondialdehyde (MDA), a decomposition product of peroxidized polyunsaturated fatty acid component of membrane lipids, using thiobarbituric acid (TBA) as the reactive material (Heath & Packer 1968). The leaf tissues (250 mg) were homogenized with 5% (w/v) trichloroacetic acid (TCA) followed by centrifugation at 15000 × g for 5 min. For each 250 mL aliquot, 1 mL of 20% (w/v) TCA containing 0.5% (w/v) TBA was added. The mixture was heated at 95°C for 30 min and then cooled in an ice bath to stop the reaction. The tubes were centrifuged at 10000 × g for 10 min, and the absorbance of the supernatant at 532 and 600 nm was recorded. After subtracting the nonspecific turbidity at 600 nm, the MDA concentration was determined by its molar extinction coefficient (155 mM<sup>-1</sup> cm<sup>-1</sup>) (Heath and Packer 1968).

### 2.7. Extraction and assay of antioxidative enzymes

The leaf tissues (200 mg) were homogenized in 2 mL of 100 mM potassium phosphate buffer, pH 6.8 containing 1 mM of EDTA and 1% (w/v) polyvinylpyrrolidone (PVP). The extract was then centrifuged at 4°C for 15 min at 12.000 × g in a cooled centrifuge. This supernatant was used to measure the activities of superoxide dismutase (SOD), guaiacol peroxidase (POD), ascorbate peroxidase (APX) and catalase (CAT). The protein concentration of leaf crude extract was determined according to Bradford (1976). Activity of SOD was assayed by using the photochemical nitro blue tetrazolium (NBT) method. The assay was performed in terms of SOD's ability to inhibit reduction of NBT to formazan by superoxide radical as described by

Beauchamp and Fridovich (1971). Activity of POD was determined at 25°C with guaiacol (Bergmeyer 1974). Activity of APX was measured by following the rate of hydrogen peroxide-dependent oxidation of ascorbic acid (Nakano & Asada 1981). Finally, the activity of CAT was assayed following H<sub>2</sub>O<sub>2</sub> decomposition according to Aebi (1984).

### 2.8. Statistical analyses

All the experiments were repeated at twice with three replicates (*n* = 6). Statistical analyses were performed with the statistical software package SPSS, version 16.0. The means were compared by the Duncan's test at a significance level of *P* < 0.05.

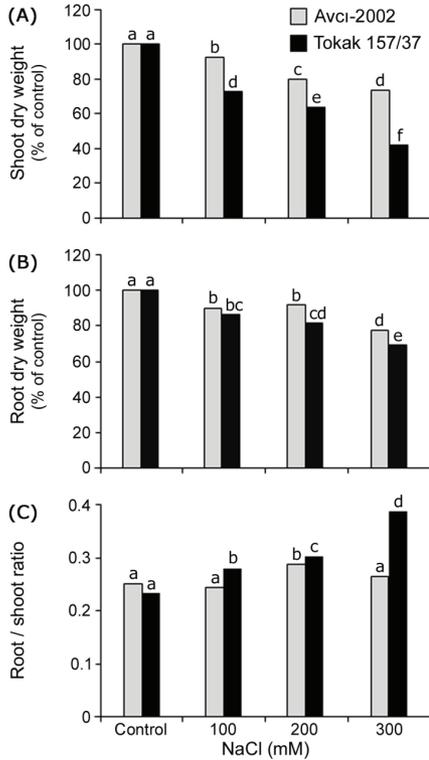
## 3. Results and Discussion

### 3.1. Effect of NaCl on growth

In many crop species, salt tolerance is reported to be a developmental regulated and stage specific phenomenon (Ashraf & Harris 2004). Shoot and root growth inhibition are the most important indices of salt tolerance (Munns 2002). In order to evaluate salt tolerance of salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37 barley cultivars at early seedling stage, dry weights of shoots and roots (percent of control) were tested under the effect of NaCl salinity. Salinity stress significantly reduced shoot and root dry weights of both barley cultivars (Figure 1A & B).

However, the decrease in shoot and root dry weights of Tokak 157/37 was significantly higher than that of Avcı-2002 under salt stress. Biomass differences among plant cultivars under saline conditions are important in determining tolerance (Bağcı et al 2003). Similar to our results dry weights of shoots and roots were less affected in many salt-tolerant cultivars such as wheat (Zheng et al 2008) and canola (Ashraf & Ali 2008). In the present study, shoot dry weights of both cultivars was affected more than root dry weights by salinity and this led to higher root/shoot ratio (Figure 1C). In this sense, no the root/shoot ratio generally changed in Avcı-2002, while it increased in Tokak

157/37 with increasing NaCl concentration. It is suggested that the root/shoot dry weight ratio could be correlated to differences in salt tolerance of barley cultivars. Similar result was reported by Dasgan et al (2002).



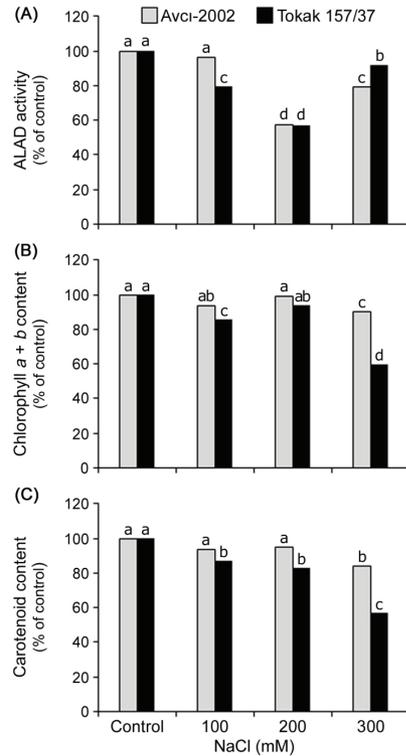
**Figure 1- Effect of NaCl salinity on shoot (A) and root (B) dry weights and root/shoot ratios (C) of salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37 barley cultivars. Values without a common letter are statistically different according to Duncan's multiple range test ( $P < 0.05$ )**

Şekil 1- Tuza-toleranslı Avcı-2002 ve tuza-hassas Tokak 157/37 arpa çeşitlerinin gövde (A) ve kök (B) kuru ağırlıkları ve kök/gövde oranları (C) üzerine NaCl tuzluluğunun etkisi. Ortak harfe sahip olmayan değerler Duncan çoklu karşılaştırma testine göre istatistik olarak önemlidir ( $P < 0.05$ )

### 3.2. Effect of NaCl on ALAD and pigments content

To study the effect of NaCl on intermediate step of chlorophyll biosynthesis, we analyzed ALAD

activity. Although ALAD activity of both cultivars decreased at all NaCl concentrations studied (except for 100 mM NaCl in Avcı-2002); the activity was increased at 300 mM NaCl compared to other NaCl concentrations (Figure 2A).



**Figure 2- Effect of NaCl salinity on ALAD activity (A), total chlorophyll (B) and carotenoid (C) contents in leaf tissues of salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37 barley cultivars. Values without a common letter are statistically different according to Duncan's multiple range test ( $P < 0.05$ )**

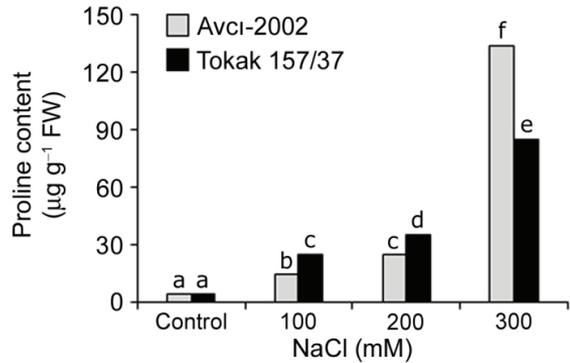
Şekil 2- Tuza-toleranslı Avcı-2002 ve tuza-hassas Tokak 157/37 arpa çeşitlerinin yaprak dokularında ALAD aktivitesi (A), toplam klorofil (B) ve karotenoid (C) içerikleri üzerine NaCl tuzluluğunun etkisi. Ortak harfe sahip olmayan değerler Duncan çoklu karşılaştırma testine göre istatistik olarak önemlidir ( $P < 0.05$ )

At 300 mM NaCl, altered ALAD activity did not accompany with reduced chlorophyll content. In light of this knowledge, the ALAD activities of Avcı-

2002 decreased at 100 and 200 mM NaCl compared to the controls, although chlorophyll pigments content did not alter. It is suggested that ALAD was not the limiting factor for the transformation of  $\delta$ -aminolevulinic acid (ALA) to porphobilinogen in the chlorophyll biosynthesis. Chlorophyll contents have been suggested as one of the parameters of salt tolerance in crop plants (Srivastava et al 1988). In our study, salt-sensitive Tokak 157/37 showed significantly higher decline in total chlorophyll content than salt-tolerant Avcı-2002 under NaCl salinity that reflects Avcı-2002 cultivar's tolerance (Figure 2B). Sairam et al (2005) observed higher chlorophyll degradation in salt-sensitive wheat cultivars as compared to tolerant cultivars. The decrease in chlorophyll content may be due to an increase of chlorophyll degradation or to a decrease of chlorophyll biosynthesis (Santos 2004). On the other hand, salt-tolerant Avcı-2002 retained higher carotenoid content than salt-sensitive Tokak 157/37 under salt stress (Figure 2C). Similar results were found in leaves of salt-tolerant cultivars of maize (Kholová et al 2010). Carotenoids are responsible for quenching of singlet oxygen (Knox & Dodge 1985), hence their comparative levels in a genotype may determine its relative salt tolerance (Kholová et al 2010).

### 3.3. Effect of NaCl on proline content

A positive correlation between magnitude of free proline accumulation and salt tolerance has been suggested as an index for determining salt tolerance potentials between cultivars (Misra & Gupta 2005). The magnitude of increase in free proline accumulation was higher in the tolerant cultivars than in the sensitive ones (Misra & Gupta 2005; Kholová et al 2010). However, some researchers reported that proline accumulation cannot be used as a sole criterion for salt tolerance (Moradi & Ismail 2007). In the present study, the proline contents increased significantly in the leaf tissues of Avcı-2002 and Tokak 157/37 barley cultivars under salinity (Figure 3).



**Figure 3- Effect of NaCl salinity on proline content in leaf tissues of salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37 barley cultivars. Values without a common letter are statistically different according to Duncan's multiple range test ( $P < 0.05$ )**

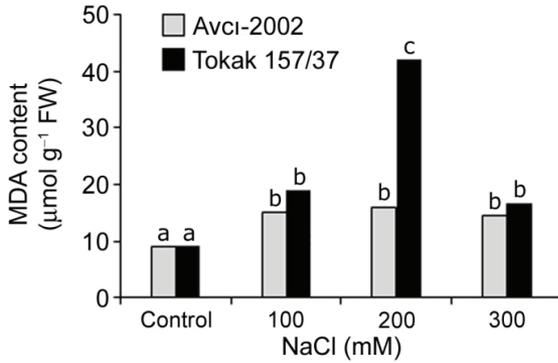
*Şekil 3- Tuza-toleranslı Avcı-2002 ve tuza-hassas Tokak 157/37 arpa çeşitlerinin yaprak dokularında prolin içeriği üzerine NaCl tuzluluğunun etkisi. Ortak harfe sahip olmayan değerler Duncan çoklu karşılaştırma testine göre istatistiki olarak önemlidir ( $P < 0.05$ )*

At 300 mM NaCl, the proline contents in the leaf tissues of salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37 increased 31- and 20-fold, respectively. According to our results, there was a positive correlation between proline accumulation and salt tolerance of barley cultivars. Among various compatible solutes, proline is the only molecule that can act as free radical scavenger and antioxidant activity (Sharma & Dietz 2006). Therefore, proline is able to stabilize proteins, DNA as well as membranes (Matysik et al 2002). It is suggested that the higher proline concentration in salt-tolerant Avcı-2002 under salt stress may help maintain structure and function of cellular macromolecules.

### 3.4. Effect of NaCl on MDA content

Lipid peroxidation measured as is produced when polyunsaturated fatty acids in the membrane undergo oxidation by the accumulation of free oxygen radicals. The level of lipid peroxidation (the amount of malondialdehyde, MDA) has been consider an indicator of salt-induced oxidation in cell membranes and a tool for determining salt

tolerance in plants (Hernández & Almansa 2002). The results reported here show that salt-sensitive Tokak 157/37 has a higher MDA content than salt-tolerant Avcı-2002 in all NaCl concentrations, indicating less oxidative damage in Avcı-2002 compared to Tokak 157/37 (Figure 4). At the highest NaCl concentration, we observed the low MDA levels in both barley cultivars, suggested that high proline content probably plays a functional role in mitigating the effects of salt stress on lipid peroxidation. Salt-tolerant barley, wheat and rice (Liang et al 2003; Sairam et al 2005; Moradi & Ismail 2007) had lower levels of lipid peroxidation which is important sign of higher oxidative damage limiting capacity under salinity.



**Figure 4- Effect of NaCl salinity on malondialdehyde (MDA) content in leaf tissues of salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37 barley cultivars. Values without a common letter are statistically different according to Duncan's multiple range test ( $P < 0.05$ )**

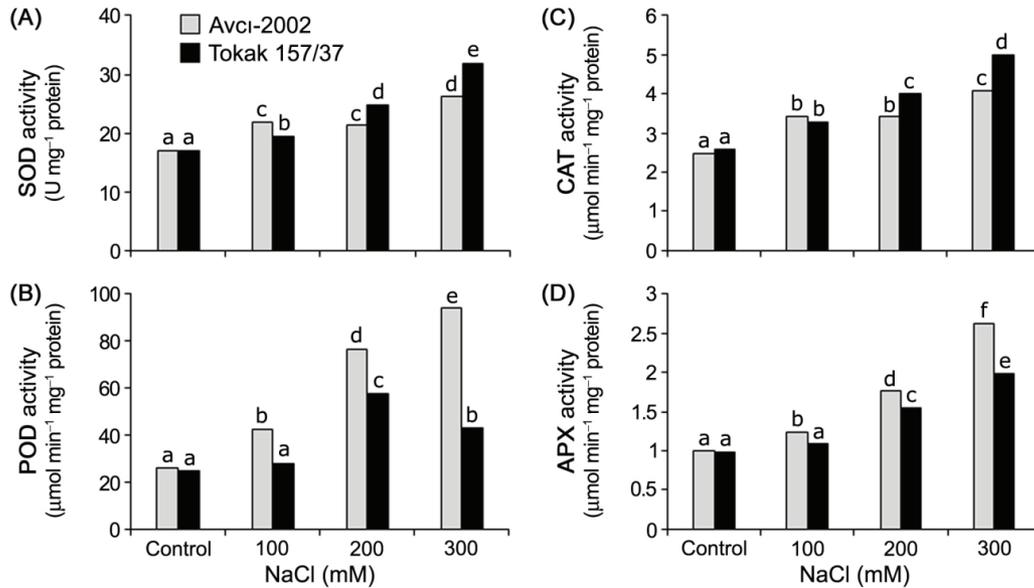
Şekil 4- Tuza-toleranslı Avcı-2002 ve tuza-hassas Tokak 157/37 arpa çeşitlerinin yaprak dokularında malondialdehit (MDA) içeriği üzerine NaCl tuzluluğunun etkisi. Ortak harfe sahip olmayan değerler Duncan çoklu karşılaştırma testine göre istatistik olarak önemlidir ( $P < 0.05$ )

### 3.5. Effect of NaCl on antioxidant enzymes activity

In plants, salinity induces oxidative stress by generating reactive oxygen species (ROS) such as  $H_2O_2$ ,  $O_2^{\cdot-}$ ,  $\cdot OH$  and  $^1O_2$  (Misra & Gupta 2006). Plants possess antioxidant enzymes as well as

antioxidant compounds to scavenge these ROS, and antioxidant capacity of plants is directly related to their salt tolerance. Our results show that salinity stress increased the activities of SOD, POD, APX and CAT in the leaf tissues of salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37 (Fig 5A-D).

The increase in SOD activity of Tokak 157/37 was greater than in Avcı-2002 at 200 and 300 mM NaCl compared to their controls. Higher POD activity was observed in Avcı-2002 than in Tokak 157/37 at all NaCl concentrations. Higher CAT activity was observed in Tokak 157/37 than in Avcı-2002 at 200 and 300 mM NaCl. The increase in APX activity of Avcı-2002 was greater than Tokak 157/37 at all NaCl concentrations compared to the control. As a result, Avcı-2002 had relatively lower SOD and CAT activities and higher POD and APX activities than Tokak 157/37. Previous studies reported that salt-tolerant cultivars generally show higher activity of these antioxidant enzymes as compared to salt-sensitive ones (Zheng et al 2008; Ashraf & Ali 2008; Kholová et al 2010). However, the results reported here show that high POD and APX activities may possibly play a significant role in imparting salt tolerance in Avcı-2002 under increasing salinity stress.  $O_2^{\cdot-}$  and  $H_2O_2$  are toxic ROS, having deleterious effects in plant tissue. SOD catalyzes the dismutation of toxic  $O_2^{\cdot-}$  to  $H_2O_2$  (Salin 1988). The salinity induced enhancement of POD, APX and CAT activities in salt-stressed plants indicates that these cells have a higher capacity for the decomposition of  $H_2O_2$  generated by SOD (Meloni et al 2003). It is noteworthy that salt-induced SOD activity in the leaves was accompanied by a greater increase in POD and APX activities in Avcı-2002 than in Tokak 157/37, indicating that the  $H_2O_2$  scavenging mechanism was more effective in Avcı-2002. Similar results related with antioxidative system activity were also reported by other researchers (Liang et al 2003; Neto et al 2006).



**Figure 5-** Effect of NaCl salinity on SOD (A), POD (B), CAT (C) and APX (D) activity in leaf tissues of salt-tolerant Avcı-2002 and salt-sensitive Tokak 157/37 barley cultivars. Values without a common letter are statistically different according to Duncan's multiple range test ( $P < 0.05$ )

Şekil 5- Tuza-toleranslı Avcı-2002 ve tuza-hassas Tokak 157/37 arpa çeşitlerinin yaprak dokularında SOD (A), POD (B), CAT (C) ve APX (D) aktivitesi üzerine NaCl tuzluluğunun etkisi. Ortak harfe sahip olmayan değerler Duncan çoklu karşılaştırma testine göre istatistiki olarak önemlidir ( $P < 0.05$ )

#### 4. Conclusions

This study showed that the differences in growth, chlorophyll and carotenoid contents, proline content, lipid peroxidation and SOD, POD, APX and CAT activities in the two barley cultivars could be ascribed to the difference in mechanisms underlying oxidative stress injury and subsequent tolerance to salinity. However, ALAD activity was not found to be an effective determinant of salt tolerance between the barley cultivars examined. Avcı-2002, a relatively salt-tolerant cultivar, exhibits a better protection mechanism against oxidative damage and lipid peroxidation by maintaining higher pigments content, proline content and activities of POD and APX than Tokak 157/37, a relatively salt-sensitive cultivar. Higher free radical scavenging capacity and more efficient protection mechanism in Avcı-2002 against salt stress as revealed by the lower lipid peroxidation level. The marked difference in the

activities of POD and APX, and proline contents in tolerant and susceptible genotypes suggest that these play an important role in the overall mechanism of salinity stress tolerance in barley.

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