Foliar application of melatonin induces tolerance to drought stress in Moldavian balm plants (*Dracocephalum moldavica*) through regulating the antioxidant system

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ABSTRACT

Melatonin, as an indoleamine molecule, regulates a wide range of physiological functions during the growth, morphogenesis and response of plants to biotic and abiotic stresses. In this research, the effect of exogenous application of melatonin (0 (distilled water), 50, 100 and 150 µM) to the leaves of Moldavian balm plants grown under different levels of drought stress (100% (control), 80%, 60% and 40% of field water capacity) was investigated. The results indicate that plants which were treated with 100 µM melatonin showed the greatest leaf surface area, lateral branching, flower length and activities of antioxidant enzymes (superoxide dismutase, guaiacol peroxidase and ascorbate peroxidase). Foliar application of 100 µM melatonin had no significant difference in catalase activity in comparison with the control and other concentrations of melatonin under normal, moderate and severe drought stress conditions. The lowest 

\( \text{H}_2\text{O}_2 \) content and lipid peroxidation (electrolyte leakage, concentrations of malondialdehyde and other aldehydes) were obtained at the concentration of 100 µM melatonin under severe drought stress. This concentration also significantly increased the chlorophyll content and enhanced the relative water content; however, foliar application of 100 µM melatonin had no significant effect on leaf length and proline content compared with the control under normal and stress conditions. The obtained results suggested that foliar application of 100 µM melatonin was more effective than the concentrations of 50 and 150 µM melatonin in reducing the adverse effects of moderate and severe drought stress.

Keywords: antioxidant enzymes, dragonhead, lipid peroxidation, melatonin, oxidative damage

Abbreviations:


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INTRODUCTION

Drought stress is one of the most serious and widespread problems which limits plant productivity because it has a negative effect on plant physiology. The effects of drought stress strongly depend on its duration, intensity, and the phonomological phase of growth and genetic tolerance capacity of plants that could reduce the growth of plants, so that caused the changes of the morphological and physiological structures and the distribution pattern of biomass or even death (Gamze et al., 2005). Drought stress causes oxidative damage by the accumulation of reactive oxygen species (ROS) that inhibit photosynthesis, stomatal closure and alter the activities of enzymes (Maksup et al., 2014). ROS formation is considered a threat to the cell because it leads to electron leakage, lipid peroxidation and subsequent membrane damage, as well as damage to nucleic acids and proteins (Polle, 2001). To reduce these damages, plants have enhanced enzymatic (including superoxide dismutase (SOD), ascorbate peroxidase (APX), catalase (CAT) and guaiacol peroxidase (GPX)) and non-enzymatic (such as ascorbic acid, glutathione, carotenoids and α-tocopherol) defence mechanisms, osmotic adjustment and stomata regulation (Shi et al., 2007). One of the objectives of advanced plant physiology is to distinguish efficient, economic and inexpensive compounds for enhancing plant tolerance to biotic and abiotic stresses. The results achieved in recent years strongly assert that melatonin could be a promising compound for the reduction of environmental stresses, especially water deficit (Arnao and Hernandez-Ruiz, 2014; Arnao and Hernandez-Ruiz, 2015; Nawaz et al., 2016; Shi et al. 2016).

Melatonin (N-acetyl-5-methoxytryptamine) is a low-molecular-weight substance with an indole ring in its structure. Tryptophan is a precursor to melatonin biosynthesis, and it exists in all living organisms, from bacteria to mammals. The level of melatonin in plants varies from one species to another, and also among varieties of a species (Arnao, 2014; Arnao and Hernandez-Ruiz, 2015). Previous studies had demonstrated the protective impacts of melatonin in mitigating biotic and abiotic stresses such as pathogen invasion, chilling stress, extreme temperature, extra copper, intense light, salinity, drought and senescence (Arnao and Hernandez-Ruiz, 2014; Manchester et al., 2000; Wang et al., 2013a). Melatonin has both hydrophilic and lipophilic nature and can pass through morphophysiological barriers easily, so it is quickly transported into cells (Galano et al., 2011; Tan et al., 2012). The major functions of melatonin are as a growth promoter, the first line of defence against oxidative damage, and as an antioxidant molecule (Arnao and Hernandez-Ruiz, 2014; Hernandez-Ruiz et al. 2004, Hernandez-Ruiz and Arnao 2008). Melatonin protects biological tissues through direct ROS scavenging or by regulating the activity and biosynthesis of enzymes and non-enzymes antioxidant, so it alleviates oxidative damage to lipids, proteins and nucleic acids (Li et al., 2015). The positive effects of melatonin on reducing oxidative damage induced by water deficit have been proved in Cucumis sativus (Zhang et al., 2014), Malus species (Li et al., 2015; Wang et al., 2013a), Glycine max (Wei et al., 2015), Lupinus albus (Arnao and Hernandez-Ruiz, 2007) and Vitis vinifera (Meng et al., 2014). Application of exogenous melatonin enables plants to remain alive under abiotic stresses through improvement of the recovery potential (Tan et al., 2012).

Moldavian balm or dragonhead (Dracocephalum moldavica) is a herb of the family Lamiaceae (Hussein et al., 2006). The effective substances of its vegetative organs are sedative and appetizing. Its essence is antibacterial and is consumed for curing stomachache, liver disorders, headache and flatulence, as well as being used by food industries, soda manufacturing and health and make-up industries (Hussein et al., 2006). Although the effects of water stress on crops have been comprehensively studied, there have been limited investigations on the behaviour of aromatic and medicinal herbs under drought stress conditions. Various environmental factors, such as water stress, affect the growth of pharmaceutical plants (Letchamo and Gosselin, 1996). Moldavian balm, with its high susceptibility to drought stress (Alaei et al., 2013), is an important medicinal and economic plant. Considering the lack of rainfall and the occurrence of drought stress in most regions of Iran, the application of plant growth promoter (such as melatonin) to help plants survive and produce optimal yields can be a promising approach which will allow the cultivation of medicinal plants in arid and low-water areas. The main objective of this research was to investigate the effect of foliar application of melatonin on the morphophysiological parameters of the Moldavian balm plant under drought stress. In this regard, we studied different aspects of plant responses, including the antioxidant defence system as well as some biochemicals such as proline and photosynthetic pigments besides
the morphological traits of plants. The obtained results can be beneficial in learning the physiological and biochemical mechanisms of melatonin action in plants that enable them to cope with drought stress.

MATERIAL AND METHODS

Plant material
This study was carried out at the research greenhouse of the Bardsir College of Agriculture, Kerman, Iran (29°55'39"N, 56°34'20"E) on April 26, 2016. The experiment was carried out as a completely randomized design in a factorial arrangement with 16 treatments and four replications. Seeds of Dracocephalum moldavica (Isfahan Seedlings and Seeds Co.) grown in plastic pots containing loamy-sandy soil under the temperature regime of 25/22°C (day/night) and the light/dark period of 14/10 h were irrigated daily with water under a relative humidity of 60%. After five weeks of growth under normal conditions, three healthy and uniform plants per pot were selected for foliar application of melatonin and four watering regimes. After optimizing melatonin concentrations, experimental treatments including 0, 50, 100 and 150 µM of melatonin in distilled water and drought stress at four levels of 100, 80, 60 and 40% of field capacity (FC) were applied to plants, and Tween-20 was used as a surfactant. Approximately 30 ml of melatonin was sprayed on each plant. The melatonin treatments were applied based on the design plan two times a week. To reach the desired watering regimes, the pots were weighed every 2 days until the water content dropped to 80%, 60% and 40% of field capacity (FC) were applied to plants, and Tween-20 was used as a surfactant.

Estimation of total chlorophyll (Chl. T)
The Lichtenenthaler method (1987) was used to determine the chlorophyll content. 0.1 g fresh leaves of Moldavian balm was extracted in 80% acetone. After filtration, its absorption was read with a UV-Visible Spectrophotometer (Cary 50; Varian Instruments, Walnut Creek, USA) at wavelengths of 646.8, 663.2 and 470 nm.

\[
\text{Chl.a} = (12.25A_{663.2} - 2.79A_{646.8}) \\
\text{Chl.b} = (21.21A_{646.8} - 5.1A_{663.2}) \\ 
\text{Chl. T} = \text{Chl.a} + \text{Chl.b}
\]

Leaf relative water content (RWC)
RWC was calculated as follows:

\[
\text{RWC} = \frac{\text{[fresh weight} - \text{dry weight}] }{\text{[saturated weight} - \text{dry weight}]} \times 100 \quad \text{(Wheatherley, 1950)}.
\]

Electrolyte leakage
To measure membrane damage, electrolyte leakage was calculated by the Ben Hamed et al. (2007) method. After washing with distilled water to clean the probable ions from the surface of the leaves, 0.2 g fresh leaves of the plant in 10 mL of double distilled water were incubated in a bain-marie at 32°C for 2 h. The primary electrical conductivity of the samples (EC₁) was measured with an EC meter (model Metrohm, Germany). In order to release all the electrolytes, samples were autoclaved at 121°C for 20 min. After cooling the test tubes to 25°C, the secondary electrical conductivity of the samples (EC₂) was re-measured and the following formula was used to calculate electrolyte leakage:

\[
\text{EL} = \left( \frac{\text{EC}_1}{\text{EC}_2} \right) \times 100
\]

Hydrogen peroxide content
\[\text{H}_2\text{O}_2\] content was calculated via the method of Velikova et al. (2000). Leaf samples were extracted with 5 ml of 0.1% trichloroacetic acid (TCA) and then centrifuged at 12,000 × g for 15 min. 0.5 ml of supernatant was combined with 1 ml of 1 M potassium iodide and 0.5 ml of 10 mM phosphate buffer (pH 7.0), and sample absorption was read at 390 nm. The \[\text{H}_2\text{O}_2\] content was estimated using the extinction coefficient (ε) 0.28 mM cm⁻¹ and expressed as µM g⁻¹ DW.

Thiobarbituric acid reactive substance (TBARS)
The amount of lipid peroxidation products was measured according to the method of Heath and Packer (1969). 100 mg of Moldavian balm leaf tissues were homogenized in 0.1% TCA (W/V), then centrifuged at 10,000 × g for 15 min. 1 ml of supernatant was combined with 5 ml of 20% TCA (W/V) solution containing 0.5% 2-thiobarbituric acid (TBA) (W/V), and the mixture was heated for 30 min. at 90°C. The samples were quickly immersed in ice for 5 min. and then re-centrifuged for 10 min. at 10,000 × g. For MDA measurement, the absorbance of the supernatant was read at
Effect of exogenous melatonin on antioxidant system and alleviation of drought stress in *D. moldavica*

532 nm and correction for non-specific pigments was performed by deducting the absorbance of the same at 600 nm. The extinction coefficient (ɛ) of 1.55 × 10^5 M⁻¹ cm⁻¹ was used to determine the MDA concentration (Heath and Packer, 1969). The extinction coefficient of 0.457 × 10^5 M⁻¹ cm⁻¹ was used to calculate the concentration of other aldehydes (Meirs et al., 1992).

**Enzyme extraction and activity determination**

0.5 g of leaf samples was homogenized in 50 mM potassium phosphate buffer (pH 7.0) containing 1 mM ethylene diamine tetra acetic acid (EDTA), 1% soluble polyvinyl pyrrolidone (PVP) and 1 mM phenylmethylsulfonyl fluoride (PMSF), with the addition of 10 mM ascorbic acid that is related to the APX assay. All extraction steps were carried out in ice at 4°C. The mixture was centrifuged at 20,000 × g for 20 min., and the supernatant was used for estimating the activity of antioxidant enzymes and protein content.

**Superoxide dismutase (SOD) (EC 1.15.1.1)**

SOD activity was determined according to the method of Giannopolitis and Ries (1977) by monitoring its ability to inhibit the photochemical decrease of nitroblue tetrazolium (NBT). One unit of SOD activity was determined as the amount of enzyme needed to cause 50% inhibition of the reduction in NBT as monitored at 560 nm.

**Catalase (CAT) activity (EC 1.11.1.6)**

Measurement of catalase activity was performed by calculating the reduction in H₂O₂ absorption at 240 nm and using the Dhindsa et al. (1981) method. The reaction mixture contained 50 mM potassium phosphate buffer (pH 7) and 15 mM hydrogen peroxide. The reaction was started by adding 100 µl of the enzyme extract to the reaction mixture. The amount of H₂O₂ in the reaction mixture was estimated after 1 minute by using the extinction coefficient of 40 mM⁻¹ cm⁻¹, which indicated the activity of the catalase enzyme.

**Guaiacol peroxidase (GPX) activity (EC 1.11.1.7)**

Guaiacol was used as a substrate to evaluate the activity of GPX. The reaction mixture consisted of 50 mM phosphorus buffer (pH7), hydrogen peroxide (1%) and guaiacol (4%). The reaction was started by adding 20 µl of enzyme extract to the reaction mixture at 25°C. The increase in absorbance at 470 nm due to guaiacol oxidation was recorded for 3 min. (Zhang et al., 2005)

**Ascorbate peroxidase (APX) activity (EC 1.11.1.11)**

Ascorbate peroxidase was measured according to the oxidation of ascorbate (ASA). The reaction mixture contained 50 mM potassium phosphate buffer (pH 7.0), 0.5 mM ascorbate, 0.1 mM H₂O₂ and 150 µl enzyme extract. The amount of ascorbate oxidation was recorded after 2 minutes by measuring the decrease in absorbance at 290 nm and using the extinction coefficient (ɛ) of 2.8 mM⁻¹ cm⁻¹ (Nakano and Asada, 1981).

**Total soluble proteins**

Protein content was calculated by the method of Bradford (1976), in which Bovine serum albumin was used as a standard.

**Proline determination**

500 mg of leaf tissue was homogenized in 7.5 ml of sulfosalicylic acid (3%) and then centrifuged. 2 ml of extract was mixed with 2 ml of glacial acetic acid and 2 ml of ninhydrin reagent. The reaction mixture was heated at 100°C for 1 h. After this time, the tubes containing the mixture were placed in a cold-water bath to stop all reactions. Then, 4 ml of toluene was added to the mixture and the tubes were well vortexed. Then, the estimation of proline content was performed based on the method of Bates

### Table 1. Mean squares (MS) for morphological traits, including leaf length, leaf surface area, number of lateral branches and flower length of Dracocephalum moldavica

<table>
<thead>
<tr>
<th>Sources of variance</th>
<th>Degrees of freedom</th>
<th>Leaf length (cm)</th>
<th>Leaf surface area (mm² per plant)</th>
<th>Number of lateral branches</th>
<th>Flower length (cm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Melatonin</td>
<td>3</td>
<td>0.091**</td>
<td>2904.29**</td>
<td>43.712**</td>
<td>7.67**</td>
</tr>
<tr>
<td>Drought stress</td>
<td>3</td>
<td>0.753**</td>
<td>32245.45**</td>
<td>511.41**</td>
<td>125.27**</td>
</tr>
<tr>
<td>Melatonin × Drought stress</td>
<td>9</td>
<td>0.033**</td>
<td>527.05**</td>
<td>8.214**</td>
<td>1.08*</td>
</tr>
<tr>
<td>Error</td>
<td>48</td>
<td>0.065</td>
<td>512.94</td>
<td>0.369</td>
<td>0.411</td>
</tr>
<tr>
<td>CV%</td>
<td>6.19</td>
<td>7.71</td>
<td>4.03</td>
<td>5.83</td>
<td></td>
</tr>
</tbody>
</table>

**, * and ns denote significant differences at 0.01, 0.05% levels, and not significant, respectively
et al. (1973), and the absorbance was determined at 520 nm.

**Statistical analysis**

Statistical analyses were accomplished by one-way ANOVA using the LSD test to determine whether the means were significantly different, taking \( p < 0.05 \) as significant. The statistical analyses and computations were performed using SAS and MSTATC.

**RESULTS AND DISCUSSION**

Foliar application of 100 µM melatonin was more effective compared with the concentrations of 50 and 150 µM melatonin, especially under moderate and severe drought stress. Therefore, based on the obtained results, we focused on the best concentration of melatonin (100 µM).

The results obtained from the variance analysis of data are given in tables 1 and 2. The results showed that drought stress had a significant \( (p \leq 0.01) \) effect on leaf length and area, the number of lateral branches and flower length (Tab. 1). The interaction between melatonin and drought stress was significant for the number of lateral branches \( (p \leq 0.01) \) and flower length \( (p \leq 0.05) \) (Tab. 1). Application of melatonin and drought stress had a significant effect on Chl. T, RWC, EL, \( \text{H}_2\text{O}_2 \), MDA, other aldehydes, GPX and APX activities \( (p \leq 0.01) \) (Tab. 2). Water stress \( (p \leq 0.01) \) and foliar application of melatonin \( (p \leq 0.05) \) significantly affected proline content, SOD and CAT activities (Tab. 2).

The highest level of drought stress reduced Chl. T by approximately 55.57% compared to the control (Tab. 3). Foliar application of 50 and 100 µM melatonin caused an increment of 7.65% and 18.8% in Chl. T, respectively, compared to the control under 40% FC drought stress (Tab. 3). Chl. T decreased approximately 3.73% at the concentration of 150 µM melatonin compared with plants which were sprayed with distilled water under the highest level of drought stress (Tab. 3).

Several studies have highlighted the dual function of melatonin in plants as a growth regulator improving growth and development (Arnao and Hernandez-Ruiz, 2014) and as a protector against biotic and abiotic stresses at low-dose (Li et al., 2012; Manchester et al., 2000). The natural antioxidant capacity of melatonin makes it a beneficial candidate for use as a biostimulant for agricultural goals and enhances the plant’s ability to cope with environmental stresses (Arnao, Table 2.

<table>
<thead>
<tr>
<th>Sources of variance</th>
<th>MS</th>
<th>Degrees of freedom</th>
<th>Mean squares (MS)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Chl. T</td>
<td>RWC</td>
<td>EL</td>
</tr>
<tr>
<td>Melatonin</td>
<td>3</td>
<td>11.48**</td>
<td>174.72**</td>
</tr>
<tr>
<td>Drought stress</td>
<td>3</td>
<td>545.42**</td>
<td>2476.53**</td>
</tr>
<tr>
<td>Melatonin × Drought stress</td>
<td>9</td>
<td>1.34**</td>
<td>20.24**</td>
</tr>
<tr>
<td>Error</td>
<td>48</td>
<td>0.71</td>
<td>18.31</td>
</tr>
<tr>
<td>CV%</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Table 2.** Mean squares (MS) for physiological parameters, including total chlorophyll (Chl. T), relative water content (RWC), electrolyte leakage (EL), concentrations of proline, hydrogen peroxide (H.O.), malondialdehyde (MDA), other aldehydes, and activities of antioxidant enzymes: superoxide dismutase (SOD), catalase (CAT), guaiacol peroxidase (GPX), and ascorbate peroxidase (APX) in *Dracocephalum moldavica*.
Table 3. The effect of foliar application of melatonin on total chlorophyll (Chl. T), relative water content (RWC), electrolyte leakage (EL), proline content, hydrogen peroxide content ($H_2O_2$), malondialdehyde (MDA) and other aldehydes in *Dracocephalum moldavica* under different levels of drought stress

<table>
<thead>
<tr>
<th>Melatonin × Drought</th>
<th>Chl. T (mg g$^{-1}$ DW)</th>
<th>RWC (%)</th>
<th>EL (%)</th>
<th>Proline content (µM g$^{-1}$ DW)</th>
<th>$H_2O_2$ content (µM g$^{-1}$ DW)</th>
<th>MDA (µM g$^{-1}$ DW)</th>
<th>Other aldehydes (µM g$^{-1}$ DW)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>100% FC</td>
<td>27.7 ± 0.8</td>
<td>89.0 ± 3.6</td>
<td>26.4 ± 0.2</td>
<td>3.9 ± 0.2</td>
<td>124.0 ± 7.0</td>
<td>0.18 ± 0.01</td>
<td>1.35 ± 0.02</td>
</tr>
<tr>
<td>80% FC</td>
<td>25.4 ± 0.4</td>
<td>77.1 ± 2.5</td>
<td>47.2 ± 1.5</td>
<td>7.7 ± 0.4</td>
<td>247.0 ± 9.6</td>
<td>0.29 ± 0.02</td>
<td>2.18 ± 0.3</td>
</tr>
<tr>
<td>60% FC</td>
<td>16.6 ± 0.2</td>
<td>64.1 ± 2.5</td>
<td>64.3 ± 0.9</td>
<td>14.3 ± 0.4</td>
<td>358.0 ± 11.5</td>
<td>0.49 ± 0.01</td>
<td>3.60 ± 0.1</td>
</tr>
<tr>
<td>40% FC</td>
<td>12.3 ± 0.1</td>
<td>52.3 ± 2.1</td>
<td>74.2 ± 1.1</td>
<td>19.2 ± 0.5</td>
<td>459.0 ± 5.7</td>
<td>0.67 ± 0.01</td>
<td>5.03 ± 0.2</td>
</tr>
<tr>
<td>50 µM Melatonin</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>100% FC</td>
<td>27.3 ± 0.1</td>
<td>89.6 ± 1.4</td>
<td>26.0 ± 0.8</td>
<td>3.7 ± 0.2</td>
<td>135.0 ± 2.5</td>
<td>0.21 ± 0.01</td>
<td>1.58 ± 0.05</td>
</tr>
<tr>
<td>80% FC</td>
<td>26.5 ± 0.5</td>
<td>79.2 ± 2.5</td>
<td>45.3 ± 0.9</td>
<td>7.3 ± 0.3</td>
<td>229.0 ± 9.6</td>
<td>0.27 ± 0.02</td>
<td>2.08 ± 0.04</td>
</tr>
<tr>
<td>60% FC</td>
<td>18.8 ± 0.3</td>
<td>66.3 ± 1.6</td>
<td>60.3 ± 0.7</td>
<td>14.9 ± 0.1</td>
<td>321.0 ± 6.4</td>
<td>0.48 ± 0.03</td>
<td>3.68 ± 0.1</td>
</tr>
<tr>
<td>40% FC</td>
<td>13.3 ± 0.2</td>
<td>57.2 ± 1.4</td>
<td>72.4 ± 0.9</td>
<td>19.9 ± 0.2</td>
<td>431.0 ± 13.5</td>
<td>0.64 ± 0.01</td>
<td>4.70 ± 0.1</td>
</tr>
<tr>
<td>100 µM Melatonin</td>
<td></td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>100% FC</td>
<td>27.5 ± 0.5</td>
<td>88.6 ± 2.1</td>
<td>26.4 ± 0.4</td>
<td>4.0 ± 0.1</td>
<td>136.0 ± 5.8</td>
<td>0.19 ± 0.01</td>
<td>1.43 ± 0.01</td>
</tr>
<tr>
<td>80% FC</td>
<td>26.6 ± 0.7</td>
<td>85.7 ± 2.9</td>
<td>42.5 ± 2.5</td>
<td>8.6 ± 0.3</td>
<td>195.7 ± 1.0</td>
<td>0.24 ± 0.02</td>
<td>1.87 ± 0.1</td>
</tr>
<tr>
<td>60% FC</td>
<td>19.5 ± 0.3</td>
<td>74.2 ± 0.4</td>
<td>57.9 ± 0.5</td>
<td>15.4 ± 0.2</td>
<td>288.0 ± 6.9</td>
<td>0.37 ± 0.01</td>
<td>2.77 ± 0.01</td>
</tr>
<tr>
<td>40% FC</td>
<td>15.2 ± 0.2</td>
<td>63.2 ± 1.0</td>
<td>67.9 ± 0.7</td>
<td>20.1 ± 0.2</td>
<td>397.0 ± 5.8</td>
<td>0.58 ± 0.01</td>
<td>4.35 ± 0.03</td>
</tr>
<tr>
<td>150 µM Melatonin</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>100% FC</td>
<td>27.4 ± 0.5</td>
<td>88.2 ± 3.7</td>
<td>27.1 ± 0.1</td>
<td>3.8 ± 0.1</td>
<td>129.0 ± 3.4</td>
<td>0.22 ± 0.01</td>
<td>1.65 ± 0.05</td>
</tr>
<tr>
<td>80% FC</td>
<td>24.6 ± 0.5</td>
<td>75.1 ± 1.9</td>
<td>49.2 ± 0.5</td>
<td>7.6 ± 0.2</td>
<td>241.0 ± 12.0</td>
<td>0.31 ± 0.01</td>
<td>2.33 ± 0.2</td>
</tr>
<tr>
<td>60% FC</td>
<td>16.2 ± 0.1</td>
<td>62.4 ± 1.2</td>
<td>69.3 ± 1.0</td>
<td>14.5 ± 0.3</td>
<td>355.0 ± 3.6</td>
<td>0.46 ± 0.03</td>
<td>3.45 ± 0.04</td>
</tr>
<tr>
<td>40% FC</td>
<td>11.9 ± 0.4</td>
<td>51.2 ± 0.7</td>
<td>75.7 ± 1.1</td>
<td>19.3 ± 0.5</td>
<td>451.0 ± 4.5</td>
<td>0.62 ± 0.03</td>
<td>4.80 ± 0.05</td>
</tr>
</tbody>
</table>

Means (± SE) followed by the same letter(s) in each column are not significantly different at the 5% level.
melatonin to one-year-old *Malus sp.* trees under drought stress conditions delayed leaf senescence, with a significant reduction in chlorophyll degradation, through the enhancement of ROS scavenging enzyme activities (Wang et al., 2013a, 2013b). Exogenous melatonin postponed induced senescence in *Hordeum vulgare* leaves and delayed the loss of chlorophylls compared with untreated plants. The cited authors inferred that those results could have been related to the interactive effect of melatonin and other plant growth regulators such as ABA and kinetin (a synthetic cytokinin with plant hormone activity) on foliar senescence (Arnao and Hernandez-Ruiz, 2015, 2017b).

Foliar application of 100 µM melatonin markedly alleviated the adverse effects of water stress. The response of RWC to the interaction of melatonin concentrations and drought levels varied (Tab. 3). RWC increased 8.57% and 17.2% at the concentrations of 50 and 100 µM melatonin, respectively, compared with the control plants under severe drought stress, while the concentration of 150 µM melatonin caused a reduction of 2.1% in RWC compared to non-treated plants under drought stress of 40% FC (Tab. 3). The highest level of drought stress (40% FC) caused a reduction of 16.67% and 38.5% in leaf length and leaf surface area, respectively, as compared to the control (Fig. 1). No significant difference was observed in leaf length between any concentration of melatonin and the control under both conditions (Fig. 1A).

The concentration of 100 µM melatonin produced a significant difference in leaf surface area compared with the other levels of melatonin and non-treated plants under moderate (60% FC) and severe (40% FC) drought stress (Fig. 1B). The response of the number of lateral branches to the

Figure 1. Effect of melatonin treatment on leaf length (A) and leaf surface area (B) of Moldavian balm plants under drought stress. The mean comparisons were performed using the LSD method at $p \leq 0.05$ significance level. Means (± SE) followed by the same letter(s) are not significantly different.
demonstrated that the decreases in RWC in plants to this type of stress (Turk et al., 2014). This study regulation of RWC is also related to plant adaptation indices for plant tolerance to drought stress, and the (Maksup et al., 2014). In addition, RWC is one of the first responses of plants to cope with water deficit (Wu et al., 2012). This precautionary approach is one of the lateral branches to prevent water loss (Wu et al., 2012). Consequently, the state of turgor membrane fluidity caused a reduction in water influx (Wu et al., 2013b) and had been reported for (Zhang et al., 2013). Similar results under drought stress (Tab. 3). Our findings showed that drought stress had a negative effect on RWC, and treatment with melatonin alleviated the adverse effect of water stress (Tab. 3). Similar results under drought stress had been reported for Malus hupehensis (Wang et al., 2013b) and Cucumis sativus (Zhang et al., 2013). An increase in water viscosity and decrease in membrane fluidity caused a reduction in water influx through the roots under water stress conditions (Wu et al., 2012). Consequently, the state of turgor gradually decreased in cells, and the cells began to shrink. Simultaneously, plants reduced the surface area of their leaves, leaf length and the number of lateral branches to prevent water loss (Wu et al., 2012). This precautionary approach is one of the first responses of plants to cope with water deficit (Maksup et al., 2014). In addition, RWC is one of the indices for plant tolerance to drought stress, and the regulation of RWC is also related to plant adaptation to this type of stress (Turk et al., 2014). This study demonstrated that the decreases in RWC in plants which were subjected to drought stress were ameliorated by the exogenous application of 100 µM melatonin, and also indicated that this concentration of melatonin could substantially reduce the damage induced by water stress. This ameliorating role of melatonin may be related to its direct and/or indirect effect on inhibiting injury to membranes (Turk et al., 2014). The larger leaf surface area, excess lateral branches and maximum flower length in melatonin-treated plants (especially with the concentration of 100 µM) clearly explain the mitigating effect of melatonin on RWC. Melatonin enhanced drought and salt tolerance in soybean plants was evident from the increased leaf length and leaf surface area, and a reduction of biomass lost when exposed to these stresses (Wei et al., 2015). Our results are in accordance with the report showing that melatonin at low concentrations increased vegetative growth and development of Glycyrrhiza uralensis (Afreen et al., 2006) and Prunus avium (Sarropoulou et al., 2012). Melatonin as a biostimulant has been proven to affect plant metabolites and stimulate biosynthesis of phytohormones, facilitate nutrient absorption, improve plant root water uptake ability, stimulate root and shoot growth, and finally enhance the quality and quantity of production and provide a much greater supply of assimilates to growing tissue (Arnao and Hernandez-Ruiz, 2007; Arnao and Hernandez-Ruiz, 2017a, 2017b; Ye et al., 2016). Exogenously applied melatonin affects developmental processes during both vegetative and reproductive growth. The structure of auxin is similar to that of melatonin, so it seems that the role of melatonin in plants is the same as of this

**Figure 2.** Effect of melatonin treatment on the number of lateral branches of Moldavian balm plants under drought stress. The mean comparisons were performed using the LSD method at \( p \leq 0.05 \) significance level. Means (± SE) followed by the same letter(s) are not significantly different. **Figure 3.** Effect of melatonin treatment on flower length of Moldavian balm plants under drought stress. The mean comparisons were performed using the LSD method at \( p \leq 0.05 \) significance level. Means (± SE) followed by the same letter(s) are not significantly different.
hormone (Janas and Posmyk, 2013). Promotion of plant growth in *Lupinus albus* (Hernandez-Ruiz et al., 2004) and some monocot plants (Hernandez-Ruiz et al., 2005) and regulation of the growth of roots, shoots, and explants, and activation of seed germination and rhizogenesis (Arnao, 2014) have shown that melatonin could have an auxin-like activity (Arnao and Hernandez-Ruiz, 2017b). Electrolyte leakage was measured to evaluate the effect of drought stress on membrane permeability. Drought stress caused an increase in EL to intercellular space so that at 40% FC it caused an increase of 64.5% in EL compared to the control (100% FC). EL decreased by approximately 8.5% in the plants which were sprayed with the concentration of 100 µM melatonin compared to the control under severe drought stress (Tab. 3). The highest level of drought stress (40% FC) caused an increase of 72% in H2O2 content as compared to the control (Tab. 3). The concentration of 100 µM melatonin markedly alleviated the effects of drought stress (Tab. 3). Melatonin at the concentration of 100 µM had a significant difference in comparison with non-treated plants and other levels of melatonin in H2O2 content under all levels of drought stress (Tab. 3). The indicator of lipid peroxidation related to the measurement of MDA and other aldehydes. The highest and the lowest concentration of MDA and other aldehydes were recorded for the control and drought stress of 40% FC treatments, respectively (Tab. 3). At all the concentrations used in this study, melatonin had no significant impact on aldehyde content compared to non-treated plants under normal conditions (Tab. 3). Among the melatonin concentrations, only the concentration of 100 µM melatonin applied through foliar spraying was more effective at both levels (60% and 40% FC) of drought stress (Tab. 3).

In the present investigation, water deficit significantly increased the generation of H2O2, electrolyte leakage, and the concentrations of MDA and other aldehydes in the leaves. Drought stress induced lipid peroxidation by the production of ROS (Shi et al., 2007), thus making the membranes leaky as evidenced by the increased electrolyte leakage. Membrane damage increased with the increment in the intensity and duration of stress. Melatonin alleviated the adverse effect of drought stress in Moldavian balm plants (Tab. 3). Increased concentrations of H2O2 lead to lipid peroxidation, which causes membrane damage and electrolyte leakage (Cui et al., 2017). Several studies have reported that the role of melatonin in the prevention of lipid peroxidation is dependent on its ability to react with lipid peroxyl (LOO•) and lipid alcoxyl (LO•) radicals and interrupt the peroxidation chain (Sarropoulou et al., 2012; Li et al., 2012; Zhang et al., 2013). The comparative rates of lipid peroxidation were evaluated in the leaves of the control and drought-treated Moldavian balm plants through the measurement of the levels of MDA and other aldehydes (Tab. 3). The concentrations of MDA and other aldehydes were similar between the control samples and all the concentrations of melatonin under normal conditions. Upon drought imposition, the levels of MDA and other aldehydes increased in non-melatonin treated plants, showing a decrease in TBARS content (Tab. 3). A protective effect of melatonin on membrane damage has been reported under water deficit (Meng et al., 2014; Zhang et al., 2013; Wang et al., 2013a). The reduction in the level of electrolytes (Tab. 3) and the decrease in TBARS concentration proved the protective role of melatonin in membrane injury induced by water stress. The response of proline content to the interaction of melatonin concentrations and drought levels were similar under normal conditions and the highest level of water stress (Tab. 3). The concentration of 100 µM melatonin without any difference from the control showed a significant difference with the concentrations of 50 and 150 µM melatonin under drought stress of 80% FC (Tab. 3). The difference in proline content was not statistically significant among the different concentrations of melatonin under moderate level of drought stress (60% FC) (Tab. 3).

Metabolic acclimation through the synthesis, accumulation, and temporary increase of compatible solutes (e.g., proline, betaine, glycine, mannitol, soluble sugar, soluble protein) are considered basic strategies for neutralizing drought stress symptoms by either conserving the vital enzymes and cellular macro-molecular structures or detoxifying free radicals. Water stress increased the endogenous levels of sugars, proline and other osmolytes (Meng et al., 2014), referring to its involvement in influencing the process of osmoregulation. Except for the concentration of 100 µM melatonin under moderate drought stress, the other groups of melatonin did not have any significant effect on proline content as compared to the control (Tab. 3). These results contradict previous reports claiming that the proline content increased in melatonin-treated plants (Antoniou et al., 2017; Arnao and Hernandez-Ruiz, 2009, 2014, 2015; Meng et al., 2014; Sarropoulou et al., 2012). Therefore, it
Effect of exogenous melatonin on antioxidant system and alleviation of drought stress in *D. moldavica*

It seems that the protective effect of melatonin on Moldavian balm plants was achieved through other mechanisms.

Changes in the specific activity of antioxidant enzymes is the consequence of oxidative stress. The effect of drought stress on SOD, CAT, GPX and APX in Moldavian balm leaves, either with or without melatonin treatment, was determined. The results showed that the activity of SOD, CAT, GPX and APX increased in drought-stressed plants (especially at 40% FC) rather than in the control groups (Fig. 4). Treatment of plants with 100 µM melatonin increased the activity of antioxidant enzymes in those plants which were subjected to the severe drought stress. SOD, CAT, GPX and APX activities were increased by approximately 69.9%, 94.9%, 68.6% and 80.2% compared to the control under 40% FC drought stress (Fig. 4). The difference in SOD activity was not statistically significant between 50 and 100 µM melatonin treatments, while the concentration of 100 µM melatonin produced significantly higher SOD activity than in the control and 150 µM melatonin under 40% FC drought stress (Fig. 4A). There was no significant difference in CAT activity between the melatonin treatments and the control under normal conditions and at the levels of 60% and 40% FC drought stress (Fig. 4B). Exogenous application of 100 µM melatonin caused an increment of 27.3% in CAT activity as compared to the control plants under mild drought stress (Fig. 4B). On the basis of these results, increasing drought levels markedly raised GPX activity (Fig. 4C). No significant difference was observed in GPX activity between all the concentrations of melatonin and non-treated plants under normal and mild drought stress conditions, while under 60% and 40% FC drought stress conditions its values for 100 µM melatonin were 6.3 and 8.41 Unit/mg protein, respectively (Fig. 4C). The data indicated that drought treatment at the levels of 60% and 40% FC caused an increment of 71% and 80.2% in APX activity, respectively, compared with the control (Fig. 4D). Treatment of Moldavian balm plants with 100 µM melatonin had a positive effect on the activity of this enzyme under moderate and severe drought conditions (Fig. 4D).

In Moldavian balm plants under drought stress, SOD, CAT, GPX and APX activities were
elevated over those in the controls (Fig. 4). Shi et al. (2007) had reported that tolerant genotypes had higher activities of ROS-scavenging enzymes than susceptible ones, so that explained the important role of the antioxidant system in plant tolerance of environmental stresses. Our results indicated that under water stress, the H$_2$O$_2$ content increased (Tab. 3). In melatonin-treated plants, the H$_2$O$_2$ content declined, which might be related to the activity of antioxidant enzymes (including SOD, CAT, GPX and APX activities). Melatonin and some of its metabolites are known as endogenous free radical scavengers (Sarropoulou et al., 2012; Zhang et al., 2013) and antioxidants (Zhang et al., 2013) which may directly scavenge H$_2$O$_2$ (Cui et al., 2017). One of the main functions of melatonin, along with the activities of CAT, GPX and APX, may be to preserve intracellular H$_2$O$_2$ concentrations at steady-state levels (Cui et al., 2017). Exogenous application of melatonin inhibited H$_2$O$_2$ accumulation through direct ROS-scavenging and enhancement of CAT, GPX and APX activities. These results could be explained by two mechanisms. First, melatonin is a broad-spectrum antioxidant and a receptor-independent free radical scavenger (Arnao and Hernandez-Ruiz, 2014). Second, it could stimulate the activities of antioxidant enzymes or induced other antioxidants to protect plants which were exposed to oxidative stress (Galano et al., 2011; Ye et al., 2016). SOD, GPX and APX showed significant rises in activity following the treatment with 100 µM melatonin. The primary role of SOD in the protection of cells against oxidative damage induced by drought stress have been proved (Sharma and Dubey, 2007). SOD activity increased significantly in Moldavian balm plants which were subjected to 100 µM melatonin under water deficit (Fig. 4A). Zhao et al. (2011) had proposed that melatonin protected *Rhodiola crenulata* cells against oxidative stress during cryo-preservation by increasing SOD and CAT activities. However, the physiological and molecular mechanisms of exogenous melatonin application which ameliorate drought tolerance of plants have not been fully clarified. Considering the positive effects of melatonin on increasing plant tolerance to drought stress, it is expected that the application of melatonin in agriculture will lead to an increase in the quality and quantity of yields and enhance plant development. Since melatonin is an inexpensive and eco-friendly substance, its application as a biostimulant could be a useful and affordable method for agricultural purposes to reduce the adverse effects of stresses and promote plant growth and development under different conditions (Arnao and Hernandez-Ruiz, 2015; Hernandez-Ruiz and Arnao, 2016).

**CONCLUSIONS**

According to the currently available data, exogenous application of melatonin, especially at the concentration of 100 µM, greatly improves the growth of Moldavian balm plants through the inhibition of membrane injury and reduced concentrations of MDA, other aldehydes and H$_2$O$_2$ under moderate and severe drought stress. These effects are probably carried out by regulating the activity of antioxidant enzymes such as SOD, CAT, GPX and APX. Reduced oxidative damage and improved water status enable plants to maintain a higher total chlorophyll content and consequently increase leaf expansion, the number of lateral branches and flower length. The concentration of 100 µM melatonin had no significant effect on proline content (as an osmoregulator) under mild and severe drought stress. Based on our results, it seems that melatonin alleviates drought stress mostly through activating the antioxidant defence system in Moldavian balm plants rather than via other regulatory pathways such as osmoprotection and proline content. Further studies concerning the details of the method and focused on elucidating the influence of exogenous melatonin on changes in the specific molecular and biochemical pathways will provide new information about the direct and indirect mechanisms of melatonin activity in plants.

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**AUTHOR CONTRIBUTIONS**

A.H., H.O. and F.N. – equally designed the experiment; M.N. – provided greenhouse, laboratory and experimental materials; H.O. and M.N. – supervised the laboratory work; R.K. – performed analytical measurements and statistical analyses, and contributed to manuscript writing; Z.T. – helped in statistical analyses; H.O. – revised the manuscript.

**CONFLICT OF INTEREST**

Authors declare no conflict of interest.
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