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Acta Biológica Colombiana, vol. 20, núm. 2, mayo-agosto, 2015, pp. 223-235
Universidad Nacional de Colombia Sede Bogotá
Bogotá, Colombia

Available in: http://www.redalyc.org/articulo.oa?id=319038639022
ALLEVIGATION OF SALINITY STRESS ON Vicia faba L. PLANTS VIA SEED PRIMING WITH MELATONIN

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SUMMARY

Melatonin is an environmentally friendly-molecule with a potent free radical scavenger and antioxidant capacity. Two pot experiments were conducted during two successive winter seasons (2011/2012 and 2012/2013) at the wire-house of the National Research Centre, Dokki, Cairo, Egypt to study the potentiality of melatonin (100 μM and 500 μM) in alleviating the harmful effect of diluted seawater at a relatively low and high concentrations (3.85 dS/m and 7.69 dS/m, respectively) on the performance of faba bean plants. The results revealed that irrigation of faba bean plants with diluted seawater reduced growth parameters (plant height, leaves number/plant, fresh and dry weights of plant), relative water content (RWC), photosynthetic pigments (chlorophylls a, b and carotenoids), indole acetic acid, total carbohydrate, K+/Ca2+, as well as the ratios of K+/Na+ and Ca2+/Na+. This was accompanied by significant increases in phenolic content, compatible solutes (total soluble carbohydrate, free amino acids, proline), Na+ and Cl relative to the control plants (untreated plants). On the other hand, melatonin treatments improved growth parameters, RWC, photosynthetic pigments, total carbohydrate, total phenolic content, indole acetic acid, K+,Ca2+ as well as K+/Na+ and Ca2+/Na+ ratios, either in the plants irrigated with tap water or with diluted seawater, as compared with corresponding controls. Meanwhile, melatonin treatments reduced the levels of compatible solutes, as well as Na+ and Cl contents, relative to those of corresponding controls. Salinity stress and/or melatonin treatments induced the production of new protein bands that did not occur in the control plants. Melatonin at 500 μM had a more pronounced effect in alleviating the adverse effects of the two salinity levels under study on the performance of faba bean plants than 100 μM melatonin.

Keywords: indole acetic acid, legumes, minerals, N-acetyl-5-methoxytryptamine, protein patterns, seawater.

RESUMEN

La melatonina es una molécula ambientalmente amigable con una potente capacidad antioxidante y de trampa de radicales libres. Dos experimentos en maderas fueron realizados en dos inviernos consecutivos (2011/2012 y 2012/2013) en instalaciones del Centro Nacional de Investigaciones, Dokki, Cairo, Egipto, para estudiar el potencial de la melatonina (100 μM y 500 μM) para disminuir los efectos nocivos del agua de mar diluida a concentraciones relativamente bajas y altas (3,85 dS/m y 7,69 dS/m, respectivamente). Los resultados mostraron que la irrigación de plantas de haba con agua de mar diluida reduce los parámetros de crecimiento (altura de la planta, número de hojas/planta, peso fresco y seco de la planta), el contenido relativo de agua (RWC), los pigmentos fotosintéticos (clorofilas a, b y carotenoides), el ácido indo lacético, los carbohidratos totales, K+, Ca2+, al igual que las relaciones K+/Na+ y Ca2+/Na+. Esto fue acompañado por un incremento significativo en el contenido de fenoles, solutos compatibles (carbohidratos solubles totales, aminoácidos libres, prolin), Na+ y Cl en comparación con las plantas control (plantas no tratadas). De otro lado, los tratamientos con melatonina mejoraron los parámetros de crecimiento, RWC, los pigmentos fotosintéticos, carbohidratos totales, contenido fenólico total, ácido indo acético, K+,Ca2+ al igual que las relaciones K+/Na+ y Ca2+/Na+, tanto en las plantas irrigadas con agua dulce de la llave como en las irrigadas con agua de mar diluida en comparación con los controles correspondientes. De otro lado, los tratamientos con melatonina redujeron los niveles de solutos compatibles, al igual que los contenidos de Na+ y Cl, en comparación con los controles. El estrés por salinidad y/o los tratamientos con melatonina indujeron...
la producción de nuevas bandas de proteínas que no estuvieron presentes en las plantas control. El tratamiento de melatonina 500 
µM tuvo un efecto más pronunciado que el tratamiento de 100 µM en disminuir los efectos adversos de los dos niveles de salinidad
estudiados sobre el comportamiento de las plantas de haba.

**Palabras clave:** ácido indol acético, agua de mar, leguminosas, minerales, N-acetyl-5-methoxytryptamina, patrones de proteínas.

**INTRODUCTION**

Faba bean (*Vicia faba* L.) is one of the most important crops cultivated in the developing countries due to the richness of
seed protein content. The importance of faba bean in Egypt lies not only as human food, animal fodder and green manure
but also due to its importance in crop rotation via fixing atmospheric nitrogen that enriches the soil with nitrogen and
organic matter as well as improving the water use efficiency of the cropping system (Khalafallah et al., 2008).

Fresh water resources are becoming limited due to the competition with human and industrial use. Furthermore,
the saline water can vary greatly in quality depending on type and quantity of dissolved salts. In this respect, irrigation
with diluted seawater plays an important role in saving fresh water resources and can be used successfully to grow crops
under certain conditions (Zeid, 2011). Tolerance of plants to salinity stress is very complex at whole plant and cellular
levels and involves changes in their morphology, physiology and metabolism (Ashraf and Harris, 2004). Reduction of
photosynthetic activity, accumulation of organic acids and osmolytes, and changes in carbohydrate metabolism are
physiological and biochemical responses to stress. One of the most important responses of plants to abiotic stresses is
over production of different types of compatible solutes (Ashraf and Harris, 2004). Compatible solutes are low molecular
weight, highly soluble compounds that are usually non-toxic at high cellular concentrations. Generally, such solutes
protect plants from stress through different courses, including contribution to cellular osmotic adjustment, detoxification
of reactive oxygen species, protection of membrane integrity and stabilization of enzymes and proteins (Bohnert and
Jensen, 1996). Furthermore, some of these solutes are commonly referred to as osmoprotectants because they protect cellular
components from dehydration injury. Bartels and Sunkar (2005) mentioned that organic solutes such as soluble
carbohydrates, soluble proteins, total free amino acids and proline have been involved in osmotic regulation in plant
and playing an important role in the tolerance of plants to salinity stress. Different abiotic stresses may cause osmotic stress,
oxidative stress and protein denaturation in plants, which lead to similar cellular adaptive responses such as induction
of stress proteins and acceleration of reactive oxygen species scavenging systems (Zhu, 2002).

Several researchers attempted to enhance the salinity tolerance of different crops through using antioxidants as pre-
sowing seed treatments or exogenous application on plants at different growth stages. Priming (osmo-conditioning)
is one of the physiological methods that improves seed performance and provides faster and synchronized germination
and performs better under adverse conditions (Ashraf and Foolad, 2005).

Melatonin (N-acetyl-5-methoxytryptamine) was discovered in plants during 1995. It is widely present in many
higher plants, dicotyledons and monocotyledons. Melatonin has been detected and quantified in roots, shoots, leaves,
fruits and seeds of a considerable variety of plant species. The levels of melatonin in plant organs vary considerably,
from picograms to micrograms per gram plant material. Generally, seeds and leaves have the highest level of melatonin
while fruits have the lowest (Van Tassel et al., 2001, Tan et al., 2007). Many microorganisms including bacteria and
fungi produce melatonin (Hardeland and Poeggeler, 2003). The decomposition of microorganisms releases melatonin
into the surrounding soil and the rootlets of the plants may absorb this melatonin and recycle it. Melatonin is an indolic
compound (biogenic indoleamine) structurally related with other important substances, such as tryptophan, serotonin,
indole-3-acetic acid (IAA), etc. Melatonin-intermediate products show antioxidant properties and have synergistic
action with other antioxidants, such as ascorbic acid, glutathione, etc. (Arnao and Hernández-Ruiz, 2009).

Melatonin is soluble in both water and lipid so it may act as a universal hydrophilic and hydrophobic antioxidant
(Janas and Posmyk, 2013). Several authors hypothesized that melatonin may possess some auxin- like effects (Kolár
and Machackova, 2005) and may act as a regulatory molecule in plants (Van Tassel et al., 2001). Its antioxidant activity
protects different plant tissues and organs, particularly reproductive tissues, fruit and germ tissues of the seed
from oxidative stress due to environmental stresses, such as drought, salinity, cold, heat, ultraviolet light and ozone
(Van Tassel et al., 2001). Tan et al. (2007) mentioned that elevated levels of melatonin probably protect plants against
water and soil pollutants through acting as a direct free radical scavenger and as an indirect antioxidant. Melatonin
directly detoxifies the hydroxyl radical, hydrogen peroxide, nitric oxide, peroxynitrite anion, peroxynitrous acid, and
hypochlorous acid that are accumulated under stressful environments. Additionally, melatonin caused increases
in the activity of several antioxidant enzymes, especially at harsh environments. One melatonin molecule may scavenge
up to 10 free radicals (Tan et al., 2007), which contrasts with the classic antioxidants that typically detoxify one
radical per molecule. Its antioxidant activity may manifest
itself in several ways: (i) direct free radical scavenging, (ii) elevating the antioxidant enzyme activity, (iii) protecting antioxidant enzymes from oxidative damage, (iv) increasing the efficiency of mitochondrial transport chain and (v) reducing the generation of free radicals (Tan et al., 2010). Arnau and Hernández-Ruiz (2009) showed that melatonin content in roots increased due to stress, reaching up to six times the melatonin content of control roots and such increase probably plays an important role in the defense against stress.

Several investigators have studied the physiological role of melatonin in plants. These studies suggested that melatonin was involved in many plant functions as delaying flower induction (Kolár et al., 2003); stimulation of hypocotyl, coleoptile and root growth (Chen et al., 2009), and protection against chlorophyll degradation (Arnau and Hernández-Ruiz., 2009). Paredes et al. (2009) reported that melatonin functions in plants can be recognized into three categories: growth promoters as auxins; antioxidants for free radicals and serve as a first-line defense against oxidative stress; and other functions (signal molecules for circadian maintenance, regulation of flower development, or maintenance of developmental stages in fruit tissues).

Thus, this work aimed to study the effect of seed priming with melatonin in ameliorating the harmful effects of salinity (diluted seawater irrigation conditions) on the performance of faba bean plants.

MATERIALS AND METHODS

Materials
Seeds of faba bean (Vicia faba L. cv. Giza 461) were obtained from the Legumes Crops Research Department, Ministry of Agriculture and Land Reclamation, Egypt. Melatonin was purchased from Science Lab Company, 14025 Smith Road, Houston, Texas, USA.

Methods

Growth conditions
Two pot experiments were conducted at the wire-house of the National Research Centre, Dokki, Cairo, Egypt on the middle of November during two growing seasons (2011/2012 and 2012/2013). During this period, temperature ranged from 10–27 °C. Relative humidity ranged from 21–87 %. Healthy faba bean seeds were selected for uniformity by choosing those of equal size and identical color. The selected seeds were washed with distilled water, sterilized with 1 % (v/v) sodium hypochlorite for approximately two min, and then washed thoroughly with distilled water. The seeds were divided into three groups, the first group was soaked with distilled water, while second and third groups were soaked with two concentrations of melatonin at 100 µM and 500 µM, respectively for 12 hours then allowed drying at room temperature (25 °C) for about 1h. Ten air-dried faba bean seeds were sown along a centre row in each pot (30 cm diameter) at a depth of 30 mm, in approx. 7 kg of clay: sand (3:1 v/v) soil. Granular ammonium sulphate (20.5 (w/w) % N) was applied at a rate of 40 kg N ha⁻¹, and single superphosphate (15 % P₂O₅) was added at a rate of 60 kg P₂O₅ ha⁻¹ to each pot. These doses of nitrogen and phosphorous were added and mixed thoroughly into the soil of each pot immediately before sowing.

The experiment consisted of three levels of melatonin namely 0 µM (control), 100 µM and 500 µM considered as ME₀, ME₁, and ME₂ respectively where melatonin was dissolved in distilled water. Irrigation water consisted of two levels of diluted seawater namely 3.85 dS/m and 7.69 dS/m that were referred to as S₁ and S₂, respectively, whereas the control plants were irrigated with tap water (S₀). Treatments were arranged in a factorial manner with six replicates for each treatment. Ten days after sowing, (DAS), faba bean seedlings were thinned leaving four uniform seedlings per pot.

Starting from day 15th, plants were irrigated with the two levels of diluted seawater mentioned above. Irrigation was carried out as follows, 3 times with diluted seawater followed by irrigation with tap water once and so on till the end of experiment.

Data recorded
Plants were sampled during vegetative stage (75 days after sowing) for measurement of some growth parameters (plant height, number of leaves/plant, fresh and dry weights of plant, and relative water content (RWC), moreover, fresh leaves were used for determination of photosynthetic pigments, endogenous indole acetic acid and electrophoretic protein bands. Oven-dried leaves (for 72 h at 70 °C) were ground to a powder and kept in a desiccators to determine total carbohydrates, total soluble carbohydrate, total phenolic contents, total free amino acid, proline, as well as some mineral contents (Na⁺, K⁺, Ca²⁺, Cl⁻). It is imperative to mention that, data of the yield and its components as well as nutritive value of the yielded seeds especially its melatonin content will be reported in another research paper.

Measurements
Relative water content (RWC) was measured in the first fully-expanded leaf (from the top) using the method of Yamasaki and Dillenburg (1999).

\[
\text{RWC} (%) = \frac{\text{Fresh weight} - \text{Dry weight}}{\text{Turgid weight} - \text{Dry weight}} \times 100.
\]

Chlorophyll a, chlorophyll b and carotenoids concentrations were estimated using the method of Moran (1982). Indole acetic acid content was determined according to Larsen et al. (1962). Fresh leaves were subjected to protein analysis according to their molecular weights by denatured sodiumdodecylsulphate (SDS)-PAGE as described by Laemmli (1970). Total soluble carbohydrates
were determined according to Smith et al. (1956). Total carbohydrates were determined according to Dubois et al. (1956). Free amino acid content was determined with the ninhydrin reagent method (Yemm and Cocking, 1955). Proline was estimated according to Bates et al. (1973). Total phenolic compounds were determined according to the method described by Zhang and Wang (2001). Mineral contents of faba bean leaves (Na⁺, K⁺ and Ca²⁺) were determined according to Chapman and Pratt (1978) using a flame photometer. Cl⁻ was determined according to the titration method described by Jackson (1973).

**Statistical Analysis**

All data were subjected to analysis of variance (ANOVA) for a randomized complete block design, after testing for homogeneity of error variances according to the procedure outlined by Gomez and Gomez (1984). Statistically significant differences between means were compared at $p \leq 0.05$ using Duncan’s multiple range test and presented with the standard errors.

**RESULTS**

**Growth Parameters**

All the measured growth parameters (plant height, leaves number/plant, fresh and dry weights of plant) as well as relative water content (RWC) decreased as a result of application of the two salinity levels ($S_1$, $S_2$) relative to control plants ($S_0$) (Table 1). These decreases were significant at $S_1$ treatment, whereas, decreases due to $S_2$ treatment were significant only in plant dry weight and RWC relative to control plant. Regarding melatonin effect on plants irrigated with tap water, it was noted that both melatonin concentrations ($ME_1$ and $ME_2$ as 100 µM and 500 µM) increased all examined growth parameters. These increases were significant at 500 µM melatonin ($ME_2$), meanwhile, melatonin at 100 µM ($ME_1$) caused non-significant increases in all parameters except RWC that showed significant increase relative to control plants ($S_0ME_0$). Under salinity stress ($S_1$ and $S_2$), 100 µM and 500 µM melatonin ($ME_1$ and $ME_2$) caused increases in all the examined growth parameters relative to corresponding controls, where the increases in plant dry weight and RWC were significant. It is worthy to mention that the beneficial effects of melatonin treatments in alleviating the harmful effect of salinity stress on the growth parameters was more pronounced in the plants grown under the higher salinity level ($S_2 = 7.69$ dS/m) than those grown under lower salinity level ($S_1 = 3.85$ dS/m) relative to corresponding controls.

**Photosynthetic Pigments**

The two applied salinity levels ($S_0ME_0$ and $S_1ME_0$) caused a decrease in all components of photosynthetic pigments (chlorophylls a, b and carotenoids) and consequently chlorophyll a+b as well as total photosynthetic pigments relative to the control plants ($S_0ME_0$) (Table 2). On the other hand, chlorophyll a, a+b, and total photosynthetic pigments were significantly increased in faba bean plants irrigated with tap water under the effect of 100 µM melatonin, meanwhile 500 µM melatonin caused significant increases in all the components of photosynthetic pigments except the carotenoid content relative to the control ($S_0ME_0$). Interaction between the two salinity levels (3.85 dS/m and 7.69 dS/m) and melatonin (100 µM and 500 µM) showed that both melatonin concentrations caused significant increases in chlorophyll a and non-significant changes in chlorophyll b and carotenoids relative to corresponding

**Table 1.** Effect of melatonin at 100 µM ($ME_1$) and 500 µM ($ME_2$) on growth parameters of faba bean plants irrigated with tap water ($S_0$) or diluted seawater at 3.85 dS/m ($S_1$) and 7.69 dS/m ($S_2$). The presented results are means of the measurements taken at two successive seasons (2011/2012 and 2012/2013) for six replicates at each season.

| Treatments | Salinity (dS/m) | Melatonin (µM) | Plant height (cm) | Change % | Leaves number / Plant | Change % | Plant fresh weight (g) | Change % | Plant dry weight (g) | Change % | RWC % | Change % |
|------------|----------------|----------------|-------------------|----------|----------------------|----------|---------------------|----------|--------------------|----------|-------|----------|
| $S_0$      |                |                |                   |          |                      |          |                     |          |                    |          |       |          |
| $ME_0$     |                | 59.06±0.78a    | -                 | 12.77±0.70b  | -              | 21.85±0.98bc | -        | 3.60±0.18c        | -        | 80.35±0.49f | -        |       |          |
| $ME_1$     |                | 62.56±0.67b   | 9.92             | 13.50±0.29ab  | 5.71         | 27.28±0.45ab  | 24.85    | 4.04±0.03bc       | 12.22    | 82.75±0.44d | 2.98    |       |          |
| $ME_2$     |                | 74.76±0.57a   | 26.58             | 14.95±0.37ab  | 17.07        | 27.95±0.83b   | 27.91    | 4.70±0.06a        | 30.55    | 84.76±0.48a | 5.49    |       |          |
| $S_1$      |                |                |                   |          |                      |          |                     |          |                    |          |       |          |
| $ME_0$     |                | 54.50±0.38ad  | -                 | 11.65±0.39bcd | -              | 16.75±0.66ab | -        | 2.48±0.23c       | -        | 58.77±0.45e | -        |       |          |
| $ME_1$     |                | 56.23±0.38bcd | 3.17             | 12.24±0.15c   | 5.06         | 20.48±0.64bc  | 22.26    | 3.33±0.06c        | 34.27    | 62.60±0.12d | 6.52    |       |          |
| $ME_2$     |                | 57.83±0.71bc  | 6.11             | 12.88±0.13ab  | 10.55        | 22.08±0.52bc  | 31.82    | 3.58±0.04bc      | 44.35    | 64.01±0.02e | 8.91    |       |          |
| $S_2$      |                |                |                   |          |                      |          |                     |          |                    |          |       |          |
| $ME_0$     |                | 44.50±1.15c   | -                 | 9.13±0.22c   | -             | 13.06±0.89bc | -        | 1.48±0.05c       | -        | 43.82±0.39f | -        |       |          |
| $ME_1$     |                | 50.03±0.99de  | 12.42             | 9.99±0.18ab   | 9.42         | 17.97±0.71cd  | 37.59    | 2.09±0.06d       | 41.21    | 54.46±0.31f | 24.28   |       |          |
| $ME_2$     |                | 53.10±0.5d    | 19.32             | 10.82±0.10abcd | 18.51       | 19.87±0.16c   | 52.14    | 2.40±0.05d       | 62.16    | 55.32±0.19f | 26.24   |       |          |

RWC (Relative Water Content).

Means followed by the same letter for each tested parameter are not significantly different by Duncan’s test ($p \leq 0.05$) and presented by± SE.
controls. It is imperative to mention that the enhancement effect of 500 μM melatonin (ME) on photosynthetic pigments was more pronounced than that by 100 μM melatonin (ME), either in the plants irrigated with tap water (S0) or diluted seawater (S1 and S2). These increases in total photosynthetic pigments were 22.31%, 12.87% and 15.85% in the plants treated with 500 μM melatonin and irrigated with tap water (S0), diluted seawater at lower (S1) and higher (S2) concentrations, respectively, as compared with corresponding controls.

**Biochemical Constituents and Compatible Solutes**

The two applied salinity levels (S0,ME0 and S1,ME1) caused significant and gradual decreases in total carbohydrate and indole acetic acid contents relative to the control plant (S0,ME0) (Table 3). However, soluble carbohydrate, phenolic content, free amino acid and proline were significantly and gradually increased by increasing salinity levels relative to control plant (S0,ME0). Melatonin concentrations (ME and ME) in absence of salinity caused gradual increases in total carbohydrate, phenolic content and indole acetic acid accompanied by gradual decreases in soluble carbohydrate, free amino acid and proline content relative to control plant (S0,ME0). Under salinity stress, the effect of melatonin on some biochemical constituents and compatible solutes of faba bean plants was more or less similar to its effect on the plants irrigated with tap water. It is worthy to mention that, 500 μM melatonin was more effective than 100 μM melatonin either in enhancement of some parameters (total carbohydrate, phenolic content and indole acetic acid) or inhibition of the others (soluble carbohydrate, free amino acid and proline).

### Table 2. Effect of melatonin at 100 μM (ME) and 500 μM (ME) on photosynthetic pigments (mg/g fresh weight) of faba bean plants irrigated with tap water (S0) or diluted seawater at 3.85 dS/m (S1) and 7.69 dS/m (S2). The presented results are means of the measurements taken at two successive seasons (2011/2012 and 2012/2013) for six replicates at each season.

| Treatments | Salinity (dS/m) | Melatonin (μM) | Cholorophyll a | Change | Cholorophyll b | Change | Carotenoid | Change | Cholorophyll (a+b) | Change | Total pigments | Change |
|------------|----------------|---------------|----------------|--------|----------------|--------|------------|--------|-------------------|--------|---------------|--------|
| S0         | ME0            | 1.70±0.014a   | -              | 0.45±0.014ab | -      | 0.26±0.017abc | -      | 2.16±0.028b     | -      | 2.42±0.046b     | -      |
| S1         | ME0            | 1.91±0.046a   | 12.35          | 0.51±0.006b  | 13.33  | 0.27±0.014abc | 3.85   | 2.42±0.051b     | 12.04  | 2.69±0.066b     | 11.16  |
| S2         | ME0            | 2.01±0.020c   | 18.23          | 0.60±0.006c  | 33.33  | 0.35±0.023a   | 34.61  | 2.61±0.014a     | 20.83  | 2.96±0.008c     | 22.31  |
| S0         | ME0            | 1.40±0.009d   | -              | 0.36±0.020d  | -      | 0.25±0.031abc | -      | 1.77±0.028d     | -      | 2.02±0.060d     | -      |
| S1         | ME0            | 1.52±0.014c   | 8.57           | 0.35±0.029c  | -2.78  | 0.24±0.025abc | -4.00  | 1.87±0.043b     | 5.65   | 2.12±0.017b     | 4.95   |
| S2         | ME0            | 1.59±0.003e   | 13.57          | 0.37±0.023d  | 2.78   | 0.32±0.040abc | 28.00  | 1.96±0.025c     | 10.73  | 2.28±0.066c     | 12.87  |
| S0         | ME0            | 1.20±0.017a   | -              | 0.26±0.009a  | -      | 0.17±0.011a   | -      | 1.46±0.025a     | -      | 1.64±0.014a     | -      |
| S1         | ME0            | 1.32±0.017a   | 10.00          | 0.28±0.011a  | 7.69   | 0.18±0.008bc  | 5.88   | 1.60±0.028b     | 9.59   | 1.78±0.020b     | 8.53   |
| S2         | ME0            | 1.39±0.002d   | 15.83          | 0.29±0.025d  | 11.54  | 0.21±0.014abc | 23.53  | 1.69±0.023c     | 15.75  | 1.90±0.037c     | 15.85  |

Means followed by the same letter for each tested parameter are not significantly different by Duncan’s test (p ≤ 0.05) and presented by± SE.

### Table 3. Effect of melatonin at 100 μM (ME) and 500 μM (ME) on some biochemical constituents and compatible solutes of faba bean plants irrigated with tap water (S0) or diluted seawater at 3.85 dS/m (S1) and 7.69 dS/m (S2). The presented results are means of the measurements taken at two successive seasons (2011/2012 and 2012/2013) for six replicates at each season.

| Treatments | Salinity (dS/m) | Melatonin (μM) | Total carbohydrate (% dry weight) | Soluble carbohydrate (% dry weight) | Total phenolic content (mg/g dry weight) | Free amino acid (mg/g dry weight) | Proline (mg/g dry weight) | Indole acetic acid (μg/100 g fresh weight) |
|------------|----------------|---------------|----------------------------------|-------------------------------------|--------------------------------------|---------------------------------|--------------------------|----------------------------------------|
| S0         | ME0            | 16.27±0.62a   | 3.41±0.26d                      | 26.89±0.15a                        | 9.80±0.17a                          | 0.47±0.008c                    | 53.04±0.03a               | 16.90±0.75a                            |
| S1         | ME0            | 18.54±0.37a   | 3.18±0.18a                      | 27.86±0.88a                        | 7.87±0.41a                          | 0.40±0.002d                    | 61.73±1.00a               | 20.07±0.53a                            |
| S2         | ME0            | 19.22±0.40a   | 2.61±0.02a                      | 29.83±0.24a                        | 6.36±0.11a                          | 0.34±0.023c                    | 78.09±0.34a               | 25.07±0.47a                            |
| S0         | ME0            | 14.14±0.32a   | 5.12±0.03a                      | 27.52±0.29a                        | 16.90±0.75a                        | 0.69±0.020bc                   | 41.43±0.40a               | 50.58±0.74a                            |
| S1         | ME0            | 16.84±0.35a   | 5.07±0.04a                      | 28.72±0.56a                        | 13.77±0.19a                        | 0.61±0.014c                    | 50.58±0.74a               | 51.25±0.43a                            |
| S2         | ME0            | 17.15±0.34b   | 4.82±0.1a                       | 31.92±1.06a                        | 9.10±0.06a                          | 0.57±0.14d                     | 49.04±0.52a               | 51.25±0.43a                            |

Means followed by the same letter for each tested parameter are not significantly different by Duncan’s test (p ≤ 0.05) and presented by± SE.
Mineral Content

The two applied salinity levels \((S_0, ME_0)\) and \((S_1, ME_1)\) caused significant and gradual increases in \(Na^+\) and \(Cl^-\) percentages accompanied by significant and gradual decreases in \(Ca^{2+}\) and \(K^+\) percentages as well as ratios of \(K^+/Na^+\) and \(Ca^{2+}/Na^+\) relative to the control plants \((S_0, ME_0)\) (Table 4). Concerning melatonin effect on mineral content of faba bean plant irrigated with tap water, it was found that \(ME_1\) and \(ME_2\) caused non-significant decreases in \(Na^+\) and \(Cl^-\) percentages accompanied by increases in the percentages of \(K^+\) and \(Ca^{2+}\) as well as ratios of \(K^+/Na^+\) and \(Ca^{2+}/Na^+\) relative to control plants \((S_0, ME_0)\). Furthermore, melatonin effect on mineral content of faba bean plants irrigated with two levels of diluted seawater was more or less similar to those obtained in the plants irrigated with tap water. Both melatonin concentrations caused significant decreases in \(Na^+\) and \(Cl^-\) percentages accompanied by significant increases in \(K^+\) and \(K^+/Na^+\) ratio, as well as non-significant increases in \(Ca^{2+}\) relative to corresponding controls. It was noted also that 500 \(\mu M\) melatonin was more effective than 100 \(\mu M\) in alleviating the harmful effect of the two salinity levels \((3.85 \text{ dS/m and 7.69 dS/m})\) on the mineral content of faba bean plants.

### Table 4. Effect of melatonin at 100 \(\mu M\) \((ME_1)\) and 500 \(\mu M\) \((ME_2)\) on some mineral content (%) of faba bean plants irrigated with tap water \((S_0)\) or diluted seawater at 3.85 \(\text{dS/m} (S_1)\) and 7.69 \(\text{dS/m} (S_2)\). The presented results are means of the measurements taken at two successive seasons \((2011/2012 \text{ and 2012/2013})\) for six replicates at each season.

| Treatments | Salinity (dS/m) | Melatonin \((\mu M)\) | Na\(^+\) | K\(^+\) | Ca\(^{2+}\) | Cl\(^-\) | K\(^+\)/Na\(^+\) | Ca\(^{2+}/Na^+\) |
|------------|---------------|-------------------------|--------|-------|--------|--------|----------------|--------------|
|            | S\(_0\)       | ME\(_0\) 1.23±0.011\(^e\) | 2.08±0.005\(^d\) | 0.94±0.008\(^bc\) | 0.30±0.002\(^ef\) | 1.69±0.011\(^e\) | 0.77±0.002\(^e\) |
|            |               | ME\(_1\) 1.22±0.11\(^a\) | 2.62±0.028\(^a\) | 1.10±0.057\(^ab\) | 0.24±0.005\(^b\) | 2.14±0.002\(^c\) | 0.90±0.040\(^b\) |
|            |               | ME\(_2\) 1.20±0.005\(^c\) | 2.36±0.023\(^c\) | 1.33±0.069\(^b\) | 0.22±0.005\(^b\) | 1.96±0.025\(^c\) | 1.11±0.063\(^c\) |
|            | S\(_1\)       | ME\(_0\) 1.81±0.008\(^a\) | 1.66±0.040\(^d\) | 0.69±0.005\(^bc\) | 0.48±0.017\(^bc\) | 0.91±0.025\(^b\) | 0.37±0.002\(^d\) |
|            |               | ME\(_1\) 1.47±0.005\(^a\) | 1.87±0.014\(^d\) | 0.81±0.008\(^cd\) | 0.40±0.005\(^d\) | 1.27±0.014\(^a\) | 0.55±0.002\(^d\) |
|            |               | ME\(_2\) 1.35±0.014\(^a\) | 1.88±0.054\(^a\) | 0.82±0.014\(^a\) | 0.33±0.011\(^d\) | 1.39±0.057\(^a\) | 0.60±0.008\(^c\) |
|            | S\(_2\)       | ME\(_0\) 2.45±0.002\(^a\) | 1.33±0.020\(^b\) | 0.58±0.005\(^a\) | 0.67±0.005\(^a\) | 0.54±0.057\(^a\) | 0.23±0.002\(^d\) |
|            |               | ME\(_1\) 2.20±0.063\(^b\) | 1.69±0.005\(^b\) | 0.68±0.011\(^bc\) | 0.52±0.002\(^c\) | 0.76±0.020\(^d\) | 0.30±0.002\(^d\) |
|            |               | ME\(_2\) 1.99±0.002\(^c\) | 1.90±0.051\(^d\) | 0.72±0.005\(^b\) | 0.46±0.037\(^bc\) | 0.95±0.023\(^e\) | 0.36±0.005\(^d\) |

Means followed by the same letter for each tested parameter are not significantly different by Duncan’s test \((p \leq 0.05)\) and presented by SE.

### Table 5. Effect of melatonin at 100 \(\mu M\) \((ME_1)\) and 500 \(\mu M\) \((ME_2)\) on SDS-PAGE electrophoretic protein patterns of faba bean plants irrigated with tap water \((S_0)\) or diluted seawater at 3.85 \(\text{dS/m} (S_1)\) and 7.69 \(\text{dS/m} (S_2)\).

| Mr (kDa) | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 |
|---------|---|---|---|---|---|---|---|---|---|
| 125     | + | + | + | + | + | + | + | + | + |
| 103     | + | + | + | + | + | + | + | + | + |
| 96      | - | - | - | - | - | + | + | + | + |
| 92      | - | - | + | + | + | + | + | + | + |
| 82      | - | - | - | + | + | + | + | + | + |
| 72      | + | + | + | + | + | + | + | + | + |
| 69      | - | - | - | - | + | + | + | - | - |
| 60      | + | + | + | + | + | + | + | + | + |
| 56      | - | - | - | + | + | + | + | - | - |
| 44      | + | + | + | + | + | + | + | + | + |
| 35      | + | + | + | + | + | + | + | + | + |
| 31      | + | + | + | + | + | + | + | + | + |
| Total number | 7 | 8 | 9 | 9 | 10 | 10 | 8 | 9 | 9 |

Number of new bands: 1 = \((S_0, ME_0)\); 2 = \((S_1, ME_1)\); 3 = \((S_1, ME_2)\); 4 = \((S_2, ME_0)\); 5 = \((S_2, ME_1)\); 6 = \((S_2, ME_2)\); 7 = \((S_0, ME_0)\); 8 = \((S_0, ME_1)\); 9 = \((S_0, ME_2)\).
bands (Mr: 125, 103, 72, 60, 44, 35, 31 kDa, respectively). This number (7 bands) represented the least number of bands that were common among the different treatments. The high concentration of salinity (S₂) alone or in combination with melatonin at the relatively low (ME₁) or high (ME₂) concentration induced a new protein band having a molecular rate 96 kDa. A unique protein band (Mr: 92 kDa) was induced by 500 μM melatonin (S₀ME₂). Furthermore, a protein band having a Mr 82 kDa seemed to be a marker for all melatonin treatments either alone (S₀ME₁) or (S₀ME₂) or in combination with salinity at lower and higher levels (S₁ME₁, S₁ME₂, S₂ME₁, S₂ME₂). The low concentration of salinity (S₁) alone or in combination with melatonin at the relatively low (ME₁) or high (ME₂) concentration induced two new protein band having a molecular rate 69 and 56 kDa.

**DISCUSSION**

**Growth Parameters**

It is well known that the responses of plant growth to salinity stress vary to different extents according to the degree and duration of stress, plant variety or species and developmental stage. In the present work, the vegetative growth parameters of faba bean plants were generally adversely affected by salinity stress (Table 1). Such a reduction in growth could
be attributed to the influence of high osmotic stress and ion toxicity (Hasanuzzaman et al., 2013) or due to altered cell wall structure induced by stress (Sweet et al., 1990). Further, salinity stress might inhibit cell division, cell enlargement and expansion as mentioned by Radi et al. (2013).

The obtained data show an enhancement effect of melatonin on vegetative growth parameters of faba bean plants irrigated either with tap water or diluted seawater (Table 1). In this connection, Janas and Posmyk (2013) mentioned that exogenously applied melatonin improved developmental processes during both vegetative and reproductive growth under stress conditions. Moreover, due to its antioxidant properties, melatonin protected the roots of barley from the damaging effects of NaCl, ZnSO{	extsubscript{4}} and H{	extsubscript{2}}O{	extsubscript{2}} (Tan et al., 2010). Li et al. (2012) also reported that melatonin mediated many physiological processes in plants i.e. growth regulation and ion homeostasis and partially alleviated the salt-induced inhibition on plant growth. Melatonin has similar chemical structure as auxin-IAA so, it may play a similar role in plants as this hormone (Sarropoulou et al., 2012). Further, Hernández-Ruiz and Arnao (2008) mentioned that melatonin stimulated the vegetative growth in etiolated lupine hypocotyls in a manner similar to that of indole acetic acid. Tan et al. (2007) investigated the potential relationships between melatonin supplementation and environmental tolerance of plants. Plants having a higher content of melatonin coped better with stress under adverse environmental conditions, compared to those with lower levels of this compound (Zhang et al., 2013). Recently, there are several reports demonstrating the function of melatonin in alleviating the adverse effects of abiotic stresses (Arnão and Hernández-Ruiz, 2009, Li et al., 2012, Wang et al., 2013).

**Photosynthetic Pigments**

The depressive effect of salinity stress on chlorophyll content (Table 2) may be due to the formation of proteolytic enzymes such as chlorophyllase, which is responsible for the chlorophyll degradation and/or damaging the photosynthetic apparatus (Radi et al., 2013). In addition, Santos (2004) pointed out that the decrease in chlorophyll content in severely NaCl stressed leaves was mainly due to a decrease of ALA (5-aminolinolic acid) synthesis. This acid is a precursor of protochlorophyllide, which converts to chlorophyll when exposed to light. Sabra et al. (2012) concluded that salt concentration (100 mM NaCl) reduced Chl a, Chl b and carotenoid contents in *Echinacea purpurea* and *E. angustifolia*, where that reduction was correlated with shoot Na{	extsuperscript{+}} content rather than Cl{	extsuperscript{-}}, suggesting that Na{	extsuperscript{+}} was the major ion causing pigment reduction. Nevertheless, in other plant species like *Vicia faba*, the decline in the leaf chlorophyll was strictly attributed to Cl{	extsuperscript{-}} accumulation in the leaves (Tavakkoli et al., 2010). In the present work, melatonin at two applied concentrations (100 μM and 500 μM) positively affected the photosynthetic pigments of faba bean plants either on irrigation with tap water or diluted seawater. XD et al. (2010) mentioned that treatment with melatonin played an important role in preservation of chlorophyll and promotion of photosynthesis due to raising the antioxidant enzyme activities and antioxidant contents and thus inhibiting the production of reactive oxygen species. In addition, Arnão and Hernández-Ruiz (2009) cited that melatonin treatments lowered chlorophyll degradation and slowed down the senescence process in barley plants, where 1 mM melatonin was optimal. Li et al. (2012) illustrated that pretreatment of *Malus hupehensis* Rehd with melatonin under high salinity conditions significantly improved plant growth and photosynthetic capacity. Zhang et al. (2013) stated that treatment with 100 μM melatonin significantly reduced chlorophyll degradation in cucumber seedlings and alleviated the effect of water stress. Furthermore, the ultrastructure of chloroplasts in water-stressed cucumber leaves was maintained after melatonin treatment. Melatonin molecule significantly reduced chlorophyll degradation and suppressed the up-regulation of senescence-associated gene and increased the photosynthetic efficiency of many plants (Tan et al., 2012, Wang et al., 2013).

**Biochemical Constituents and Compatible Solutes**

* a. Total carbohydrates

Carbohydrates are supplied mainly through the process of photosynthesis and photosynthetic rates are usually lower in plants exposed to salinity and especially to NaCl (Ashraf and Harris, 2004). Hence, the reduction in photosynthetic pigments in faba bean leaves under the effect of salinity stress (Table 2) might have led to decreased levels of photo-assimilates in the leaves, mainly total carbohydrates (Table 3). Most of the plants are sensitive to salt stress, and high level of salinity caused reduction in carbohydrates (Hassanein et al., 2009). On the other hand, the melatonin enhancement effects on photosynthetic pigments that have been recorded in the present work might interpret the overproduction of total carbohydrates, and thus the concomitant enhancement of plant growth.

* b. Indole acetic acid

The decreases in indole acetic acid (IAA) under the effect of salinity stress (Table 3) were concurrent with the decrease in vegetative growth parameters (Table 1). The reduced IAA levels under salinity stress might be attributed to the inhibition in biosynthesis of indole acetic acid and/or increases in their degradation or transformation into inactive form. However, recent work indicated that the decreased auxin contents under salinity might be rather complicated as being underplayed by stress-responsive transcription factors.
and microRNAs that modulate auxin- and environment-mediated root development (Kazan, 2013). On the other hand, the promotive effect of melatonin on IAA was confirmed by Chen et al. (2009) who reported that melatonin doses increased indole acetic acid. In addition, Posmyk et al. (2009) mentioned that hydro-priming of cucumber seeds with melatonin increased IAA content.

Phenolic compounds
In response to various environmental stresses such as salinity stress, plants have developed different physiological and biochemical mechanisms to adapt or to tolerate stress. Table 3 shows that salinity stress and/or melatonin treatments enhanced the phenolic content. Salinity induced disturbances in the metabolic process leading to the increase in the synthesis of phenolic compounds (Keutgen and Pawelzik, 2009). Actually, the accretion of reactive oxygen species (ROS) under salt stress is generally coupled with changes in net carbon gain which may strongly affect the biosynthesis of carbon-based secondary compounds, particularly leaf polyphenols (Radi et al., 2013). Phenolic compounds play an important role as antioxidants in scavenging free radicals arising from their high reactivity as hydrogen or electron donors, to stabilize and delocalize the unpaired electron (chain-breaking function), and from their ability to chelate transition metal ions (Huang et al., 2005). Beneficial effects of melatonin may also result from its signaling function, through the induction of different metabolic pathways and stimulate the production of various substances, preferably operating under stress (Tan et al., 2012). Szafranska et al. (2012) showed that melatonin added to Vigna radiata L. seeds protected the roots of chilled seedlings after re-warming and increased the synthesis of phenolic compounds, particularly derivatives of p-coumaric acid.

Compatible solutes
In the present work, salinity stress caused increase of compatible solutes, whereas melatonin treatments decreased them (Table 3). The osmotic adjustment in plants subjected to salt stress occurs by the accumulation of high concentrations of osmotically active compounds known as compatible solutes such as proline, glycinebetaine, soluble sugars, free amino acids and polyamines (Jagesh et al., 2010). These authors revealed that such substances play an important role in the adaptation of cells to various adverse environmental conditions through raising osmotic pressure in the cytoplasm, stabilizing proteins and membranes, and maintaining the relatively high water content obligatory for plant growth and cellular functions.

Proline accumulation is considered as an indicator in several plant species under salt stress conditions, acting as an osmotic protectant and contributing to the turgor maintenance of cells (Jagesh et al., 2010). Proline was involved in the synthesis of key proteins that are necessary for stress responses (Iyer and Caplan, 1998). Further, the increase in proline content could be attributed to a decrease in proline oxidase activity under saline conditions (Roodbari et al., 2013). The free amino acid accumulation associated with stress may actually be a part of an adaptive process contributing to osmotic adjustment (Dubey, 1994).

Elevated sugar levels in salt stressed plants may contribute to the turgor maintenance and stabilization of cellular membranes (Jouve et al., 2004). The accumulation of soluble carbohydrates in plants has been widely reported as a response to salinity despite a significant decrease in net CO₂ assimilation rate (Murakeozy et al., 2003). According to Bohnert and Jensen (1996) carbohydrates may act as ROS scavengers and contribute to increase in membrane stabilization.

On the other hand, melatonin is a free radical scavenger and broad-spectrum antioxidant that might directly eliminate ROS when produced under stressful conditions. The accumulation of ROS under stress was inhibited by melatonin applications due to direct scavenging and/or enhanced activities of antioxidant enzymes (Tan et al., 2000, Tan et al., 2007). The results of the present work show that melatonin treatments decreased the harmful effect of salinity on faba bean plants and increased its salinity tolerance.

Mineral Content
The decreases in K⁺ and Ca²⁺ were accompanied by increases in Na⁺ and Cl⁻ in faba bean leaves under the effect of salinity levels (Table 4). This conclusion agrees with those reported by Cramer (1997) and Taibi et al. (2012) who revealed that plants exposed to NaCl take up high amounts of Na⁺, whereas the uptake of K⁺ and Ca²⁺ is significantly reduced. K⁺/Na⁺ ratio in plants under saline conditions is considered as one of the important selection criteria for salt tolerance (Ashraf and Harris, 2004). Tester and Davenport (2003) pointed out that low K⁺/Na⁺ ratios could disrupt protein synthesis in the cell, given that Na⁺ competes with K⁺ for binding sites essential for cellular function, but cannot substitute for K⁺ to activate functional enzymes. Furthermore, the maintenance of Ca²⁺ acquisition and transport under salinity constitutes an important determinant of salinity tolerance (Unno et al., 2002), making plants to be less susceptible to osmotic and specific ion injury. Taibi et al. (2012) mentioned that Na⁺ and Ca²⁺ can enter cells through ion channels. These channels may control the transport of cations to the xylem and therefore, may control cation transport to the shoot. In some cases, these channels are more selective for Na⁺ than K⁺ and the increase in Ca²⁺ concentration reduces Na⁺ conductance through these channels. K⁺ also moves through the Ca²⁺ channels and can interfere with Ca²⁺ transport (Piñeros and Tester, 1997). Chlorine is widespread in the nature,
and plants easily absorb chloride ions. Chloride ions play an important role in PSII, membrane potential stabilization and turgor and pH regulation. External Cl− concentrations higher than 20 mM can be toxic in susceptible plant species, whereas in tolerant species external concentrations four to five times higher may show no effect on plant growth (Broadley et al., 2012).

Melatonin treatments partially alleviated the harmful effect of salinity stress on mineral content of faba bean leaves (Table 4). Li et al. (2012) mentioned that melatonin might control the expression of ion-channel genes under salinity, which may possibly contribute to the maintenance of ion homeostasis and thus, improves salinity resistance in plants. The ability to limit Na+ transport into the shoots and to reduce Na+ accumulation in the rapidly growing shoot tissues is critically important for maintenance of high growth rates and protection of metabolic processes in elongating cells from the toxic effects of Na+ (Roodbari et al., 2013). Enhanced uptake of K+ and/or Ca2+ at the cost of reduced uptake Na+ in the cells of salt stressed plants is considered vital for maintaining high cellular K+/Na+ and Ca2+/Na+ ratios. Significant ameliorative effects of Ca2+ on Na+ toxicity have been reported to improve water transport (Knight et al., 1997), and growth of plants (White and Broadley, 2003).

Protein Pattern
The results of the present work showed appearance of new protein bands in faba bean plants under the effect of salinity stress and/or melatonin treatments (Table 5 and Fig. 1). The most important mechanism involved in cell protection against salt stress is the induction of de novo synthetic protein groups (Kermode, 1997). These stress proteins may provide a storage form of nitrogen that may be re-utilized when stress is over (Badr et al., 1998). It is suggested that these new proteins may play an important role in triggering a special system that help the whole plant against salinity stress. These proteins may have an osmoprotection function (Dure, 1993) or protecting cellular structures (Close and Lammers, 1993). Shukry and El Bassiouny (2002) reported that salinization induced de novo synthesis of some salt responsive proteins of Vicia faba seeds during germination and the salt responsive proteins might be osmotin, dehydrin and ubiquitin.

Moreover, the melatonin-induced new protein bands, shown in the present work, might be attributed to the synthesis of new polypeptides or might represent degradative product(s) of proteins due to the effect of hydrolytic enzymes on high molecular weight proteins. This assumption might be reinforced by that of Posmyk et al. (2009) who reported that although melatonin protected membrane structure against peroxidation during chilling, excessive melatonin levels (~4 μg/g fresh weight) in cucumber seeds provoked oxidative changes in proteins.

CONCLUSION
It could be concluded that melatonin treatments (100 μM and 500 μM) improved growth parameters, relative water content, photosynthetic pigments, total carbohydrate, total phenolic content, indole acetic acid, K+/Ca2+ and reduced the levels of compatible solutes, Na+ and Cl− contents in leaf tissues of faba bean plants irrigated with diluted seawater (3.85 dS/m and 7.69 dS/m). Melatonin at 500 μM had a more pronounced effect in alleviating the adverse effects of the two salinity levels on the performance of faba bean plants than 100 μM melatonin.

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