Research Article

*Trichoderma* and spermidine improve cadmium tolerance and phytoremediation potential in purslane (*Portulaca oleracea* L.) plant

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Abstract

In order to understand, the physiological and biochemical mechanisms of *Trichoderma longibrachiatum* (TL) and spermidine (SPD) polyamine treatment on cadmium (Cd) tolerance phytoremediation in purslane (*Portulaca oleracea*) plant and the activity of anti-oxidants enzyme (CAT, APX, POX, SOD), hydrogen peroxide and proline content as well as determination of cadmium accumulation in shoots, roots, soil and their ratio to each other, a factorial experiment was performed in a completely randomized design with three replications and three treatments. In the current study, mitigative roles of SPD and TL were assessed in Cd stressed *Portulaca oleracea* plants. SPD (1, 0.5, and 1 mM) was applied after 20 days of sowing on the branches and leaves of plants inoculated or without TL inoculating in the presence of Cd (0, 30, 60 and 90 mg.kg⁻¹). Cd stress and coexistence with TL increased the activity of antioxidant enzymes and leaf soluble protein in purslane plants. Also, the application of SPD, especially at 0.5 mM, resulted in a higher increase in leaf protein under cadmium stress in inoculated plants. Proline parameter responds differently to TL. SPD application reduced the severity of these changes. The amount of H₂O₂ was significantly reduced in plants when treated by both TL and SPD. Significant differences were observed between 0.5 and 1 mM of SPD in terms of the Cd uptake in the TL inoculated purslane shoots. Inoculated purslane plants treated by either 0.5 or 1 mM of SPD had lower Cd uptake and greater BF. In general, the results showed a synergetic effect between TL fungi and SPD application on improving the Cd phytoremediation in the purslane plant.

Keywords: Antioxidant enzymes, Cadmium toxicity, Phytoremediation, Purslane, Spermidine, *Trichoderma*

Introduction

The natural weathering of minerals, along with recent human activities, bring large quantities of heavy metals into the environment (Zhang et al., 2018). Heavy metals have become a serious threat to human health due to their direct and indirect toxic effects through accumulation in the food chain (Ghaderian and Ghotbi Ravandi, 2012). Cd is listed as the sixth most dangerous element in the US Agency for the study of toxic substances (Kumar Rai, 2008). Since some plants such as corn, canola, and purslane (Hechmi et al., 2013; Yaghoubian et al., 2016; Belouchrani et al., 2016) can absorb and remove metal contaminants such as cadmium from the contaminated environment, phytoremediation could introduce as an environmentally friendly, low-cost and new technology that uses plants to reduce the concentration of organic and metal pollutants (Luo et al., 2016). One of the aspects of phytoremediation is risk control through the stabilization of pollutants by the roots of plants in the soil.

Cd leads to increased production ROS (Shahid et al., 2014), and in return ROS rapidly damages existing macromolecules including membrane lipids, proteins, nucleic acids, and pigments, resulting in irreversible metabolic defects (Maldonado-Magna et al., 2011). To minimize the deleterious effects of ROS, plant cells have developed a complex network of enzymatic and non-enzymatic antioxidant systems to protect plant cells from oxidative disadvantage caused by different types of ROS (Fahr et al., 2013; Afzal et al., 2014). SOD, CAT, APX and GPX are ROS-scavenging enzymes that maintain ROS content within a specified range (Gupta et al., 2009). The key role of CAT, SOD, and APX as three enzymes involved in controlling Cd-induced H₂O₂ content has recently been described (Guo et al., 2019). It has been reported that annually a lot of cadmium from, industrial and municipal wastewater, along with unnecessary consumption of insecticides and high levels

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of chemical fertilizers, especially phosphate, are released into the environment (Chiapello et al., 2015). Therefore as sedentary plants, they need to have defense mechanisms for acclimating to change environmental conditions. Numerous studies indicate that Cd signals can be transmitted by polyamines and causes increased production of ROS (Arasimowicz-Jelonek and Floryszak-Wieczorek, 2012). Polyamines include Put, SPD and spermidine are known as a group of natural compounds with aliphatic nitrogen structure. Polyamines play an important role in the physiological mechanisms of plants and improve tolerance to abiotic stress (Tiburcio et al., 2014; Cai et al., 2015). Polyamines should not only be considered as protective molecules but should also be regarded as compounds involved in a complex signaling system that plays in regulating stress tolerance (Kasukabe et al., 2004). For example, Yang et al. (2013) reported enhanced salt and heavy metal enhanced tolerance in spermidine-treated Hydrocharis dubia, and Salvinia natans (Xu et al., 2008) and sunflower (Mutlu and Bozuk, 2005).

Phytoremediation is an emerging, low-cost technology that uses plants to clean and remove pollutants from the environment, meanwhile only some plants (the metallophytes) can survive in adverse environmental conditions (soil contamination with heavy metals), so they can help clean up the contaminated environment (Yadav, 2010; Ghaderian and Ghotbi Ravandi, 2012). Some plants (the metallophytes) can survive in adverse environmental conditions (soil contamination with heavy metals), so they can help clean up the contaminated environment.

Fungal strains grouped in the genus Trichoderma species are known to alter the response of plants to abiotic stresses and a widespread free-living with different species that can benefit plants by promoting root extension and plant growth, succor the remediate soil and water pollution (Akladious and Abbas, 2012; Ghorbanpouro et al., 2018; Yaghoubian et al., 2019). Recently, phytoremediation as a diverse eco-friendly remediation option, has been explored to restore the contaminated environments. This process combined two methods of phytoremediation and bioremediation simultaneously. Since purslane is a well-known plant for its antioxidant properties (Cai et al., 2004) along with its well-documented and valuable potential to phytoremediation the heavy metals, the purpose of this study was to investigate the effect of fungi symbiosis and polyamine on the improvement of phytoremediation by purslane.

Materials and methods

Plant materials and growth conditions: The experiment was designed as a factorial with completely randomized design with three replications. Treatments consisted of Trichoderma longibrachiatum (TL) fungi, three spermidine (SPD) levels: 0, 0.5 and 1 and four cadmium (Cd) levels: 0, 30, 60 and 90 mg.kg⁻¹ Cd in the soil. The level of cadmium studied in this experiment was selected based on the initial method for measuring the growth and tolerance of purslane (Portulaca oleracea L.) in Cd-contaminated soils at 11 levels of cadmium concentration. The soil was collected at 0-30 cm depth at the site of the research farm of Sari Agricultural Sciences and Natural Resources University (SANRU), Mazandaran province, Iran (N 36°39′40.1″ E 53°04′16.6″). The soil texture was sandy loam mixed with the river sand (2/1: v/v). Soil Cd initial concentration was measured prior to the experiment. To reduce microorganism’s side effects, we autoclaved the soil in autoclavable bags at 120 °C for 30 min. For each treatment, the appropriate amount of Cd (from the source CdCl₂) was dissolved in 1 liter of double-distilled water and sprayed on the soil with thin layers. To ensure the uniformity of the Cd distribution, this procedure was repeated several times until the entire solution was applied. Contaminated soils were kept in the greenhouse for four weeks in order to perform the interactions between Cd and soil at a relative humidity of 53% to ensure the necessary balance between Cd and soil (Huang et al., 2009).

TL specie was prepared from the laboratory of mycology, Department of Plant Pathology at SANRU cultured in medium contained PDA (Potato-Dextrose Agar, Merck) and incubated for 7 days at 25 °C (Lopez Errasquin and Vazquez, 2003). After the incubation, 15 ml of double-distilled water was added to each medium. The resulting solution combined with Tween 20 (biological detergent) was used to inoculate germinated seeds. Therefore, a fungus suspension with 10⁵ conidia.ml⁻¹ was used for the inoculum in the trial (Hjeljord and Tronsmo, 2003). Purslane seeds obtained from Karaj Seed Pakan Center, Iran then were surface-sterilized for 15 min in 1% (W/V) sodium hypochlorite and thereafter thoroughly washed with distilled water. After 24 h of germination, the seedlings were inoculated with TL spores as described below. Then inoculated seedlings were planted into 20 cm wide plastic pots filled with 5 kg of air-dried soil, then placed in a greenhouse at 26/17 °C (day/night) temperature with a 16 h photoperiod. SPD was purchased from Sigma company. After the acclimation period of at least two months, seedlings purslane was sprayed at the two-week interval (in the tenth leaf stage) with 0, 0.5 and 1 millimolar (mM) of SPD, until both sides of all the leaves were completely wet. At each level, spermidine was dissolved in water. Nonyl soap was added as a surfactant to both SPD solutions and also to the control treatment. The Cd and SPD concentrations were optimized and selected in a preliminary study. After 60 days of the experiment (before the emergence of flowers stage), various analyses were performed.

Enzyme extraction and assays: Antioxidant enzymes were extracted at 4 °C using 0.5 g tissue from the fresh samples of purslane seedling leaves after 14 days of SPD treatment. The samples were homogenized with liquid nitrogen, followed by adding 5 ml extraction buffer consisting of 0.1 M phosphate buffer with pH
7.5, including 0.5 mM EDTA in case of CAT and POX, or 0.1 M phosphate buffer, pH 7.5, 0.5 mM EDTA, 2 mM ASA and 5% PVP in case of APX at 5 °C (Nakano and Asada, 1981). CAT activity was measured according to the method of Aebi (1984) in 240 nm. APX activity was measured by the ascorbate oxidation rate at 290 nm (E = 2.8 mM/cm) according to Yoshimura et al. (2000). POX activity was measured according to Tang and Newton's method (2005) in 470 nm. Superoxide dismutase (SOD) activity was determined by measuring the inhibition in photo reduction of nitro blue tetrazolium (NBT) by SOD enzyme (Kumar et al., 2012). The reaction mixture contained 50 mM sodium phosphate buffer (pH 7.6), 0.1 mM EDTA, 50 mM sodium carbonate, 12 mM L-1 methionine, 50 μM NBT, 10 μM riboflavin and 100 μL of crude extract in a final volume of 3.0 mL. A control reaction was performed without crude extract. The SOD reaction was carried out by exposing the reaction mixture to white light for 15 min at room temperature. After 15 min incubation, absorbance was recorded at 560 nm using a spectrophotometer. One unit (U) of SOD activity was defined as the amount of enzyme causing 50% inhibition of photochemical reduction of NBT.

**Determination of hydrogen peroxide:** First, 0.5 g of leaf sample was homogenized in 5 ml of 0.1% TCA solution (w/v) and then centrifuged at 12000 rpm for 15 min. Then, the reaction complex containing 0.5 ml of the supernatant, 0.5 ml potassium phosphate buffer 10 Mm (pH = 7), and 1 ml KI 1 M. was prepared, and their adsorption rate was measured at 390 nm (Sergiev et al., 1997).

**Proline determination:** Proline measurement was performed using the method of Bates et al. (1973). In this method, 0.5 g of fresh leaves of *Portulaca oleracea* was homogenized and after transferring to the tube and adding 10 ml of sulfosalicylic acid 3% (w/v) for 10 to 15 minutes at 4 °C, it was centrifuged in 15000 rpm. Then, we added 2 ml acid ninhydrin and 2 ml glacial acetic acid to 2 ml of filtered extract and incubated in a boiling water bath for 1 h. The reaction was finished in an ice bath, finally, 4 ml of toluene was added to each test tube and Vertex was performed for 20 seconds. The concentration of soluble proline in toluene was determined using a spectrophotometer at a wavelength of 520 nm, and finally, the amount of proline in the plant was determined according to the standard proline curve.

**Determination of Cd accumulation:** An atomic absorption device (AAS-6300, Shimadzu, Kyoto, Japan) was used to measure the concentration of cadmium in soil and aerial and terrestrial portions of purslane (Helrich, 1990). Also, the BCF and the TF, assessed to determine the relative translocation of Cd from belowground parts to the aboveground shoot of the plant. BCF and TF calculated as follows (Ghosh and Singh, 2005; Gupta et al., 2008):

\[
BCF = \frac{\text{concentration of a metal in roots}}{\text{concentration of metal in the medium}}
\]

\[
TF = \frac{\text{concentration of a metal in the shoot}}{\text{concentration of metal in the roots}}
\]

The experiment was conducted in a completely randomized design with three replicates. Mean comparisons were made by LSD (alpha = 0.05) using SAS version 9.1. The normality of data was tested using the one-sample Kolmogorov–Smirnov test (Massey, 1952).

**Results and discussion**

In the present study, the activities of four antioxidant enzymes SOD, APX, GPX, CAT were evaluated to assess the function of TL and SPD in regulating plant antioxidant against cadmium stress (Fig. 1 to 4). Our results indicated that SOD (Fig. 1A) showed a positive response to Cd concentration gradients. SOD activity increased by about 5, 14, and 14% at 30, 60, and 90 mg.kg⁻¹ of Cd concentrations, respectively as compared to the control. This result was consistent with the results of Guo et al. (2019) who found that Cd concentration increased the activities of SOD in different wheat varieties in different growth stages. Application of 0.5 mM SPD increased SOD activity at low levels of cadmium toxicity (0 and 30 mg.kg⁻¹ Cd) by more than 17.84 and 13.52% compared to the control, respectively. On the other hand, the use of spermidine in higher concentrations of cadmium had no significant effect on the SOD activity. These results are similar to some studies showing SPD was a collector of ROS (e.g. Satish et al., 2018). Most cases, during abiotic stress in plants, a close relationship has been reported between plant protection at the time of SPD application and increased antioxidant levels (Satish et al., 2018). Also, we demonstrated here that TL induced SOD expression in chloroplasts and significantly improved SOD activity (12.19%) in plant as compared to the uninoculated control (Fig. 1B). Decreased SOD activity in Cd toxicity had been previously reported in both wheat and bean plants (Cardinals et al., 1984; Milone et al., 2003), although it increased in sunflower, rice, and soybean (Kuo and Cao, 2004; Sobkowiak et al., 2004; Laspina et al., 2005).

The results of the APX are shown in Figure 2A. APX activity negatively responds to Cd concentrations (not always). However, a marked increase in APX activity was monitored after spraying 1 mM of SPD (Fig. 2C) which was more pronounced than the Cd treatment alone

SPD is a messenger regulator in stress signaling pathways leading to changes in the expression of different genes associated with stress in plants (Kasukabe et al., 2004) and may alter the antioxidant content in the plant (Radhakrishnan and Lee, 2014). APX activity was significantly increased in the presence of Cd, particularly in the inoculated plants as compared to the control. CAT, along with POX and APX, converts H₂O₂ to water and oxygen (Ahmad et al., 2010).

Significant results were obtained in CAT activity in...
purslane under Cd stress at different doses of spermidine. Most of the changes occurred at lower levels of cadmium toxicity, with the difference that a spray of 0.5 mM of this polyamine solution increased the activity of this enzyme in plants inoculated with fungi (Fig. 3B). While the highest activity of this enzyme was observed during foliar application of 1 mM spermidine in plants not inoculated with fungi (Fig. 3C). In the first place, this effect was due to the exogenous Spd contribution to the improvement of the antioxidant Halliwell-Asada path way enzyme functioning (Kubis, 2001), and secondly, resulted from the fact that PAs are able to affect the other hydrogen peroxide scavenging enzyme activities – catalase and guaiacol peroxidase (Kubis, 2003). Shen et al. (2000) reported that SPD can counteract the chill-induced activation of NADPH oxidase, and by this way, diminishes ROS generation in cucumber cultivars.

By contrast, GPX activity was greater at higher concentrations of Cd, especially in inoculated plants (Fig. 3D). There was a synergistic effect between TL fungi and SPD. However, spraying of SPD at the rate of 0.5 Mm had a significant and positive effect on GPX activity in the uninoculated plants (Fig. 3E).
Trichoderma and spermidine improve cadmium tolerance...

Fig. 3 - Effects of exogenous SPD (0, 0.5, 1 mM) and Trichoderma on CAT (A, B, C) and GPX (D, E, F) in purslane seedlings under four concentrations (0, 30, 60, 90 mg kg⁻¹) CdCl₂ stress. Different letters indicate significant differences (P ≤ 0.05).

Differences observed in antioxidant activities in different plant species under Cd toxicity probably depended on the amount of metal concentration and the duration of exposure to stress and the type of plant. (Gallego and Tomaro, 2005; Jawad Hassan et al., 2020). Our results could be indicative of cell adaptation to metal after prolonged exposure to the plant. Abiotic stresses such as heavy metals by ROS production lead to molecular damage to plant cells (Zhang et al., 2005). Although Cd does not generate ROS directly, it generates oxidative stress by interrupting the antioxidant defense system (Toppi and Gabrielli, 1999). Antioxidant enzymes such as GPX, APX, and CAT balance the production and destruction of ROS. Meanwhile Cd inhibits the coenzyme by inhibiting the Calvin cycle enzymes, preventing the electron from receiving PSI (Vassilev et al., 2004).

Protein: As shown in Figure 4A, TL pretreatment caused a great induction of soluble protein content in the leaves of purslane plant (P < 0.05), particularly in the 0.5 mM of SPD. Alternatively, SPD spraying (particularly in the 0.5 mM concentration), showed a significant increase in soluble protein content as compared to the control. SPD showed an ameliorating effect on the soluble protein content in purslane under Cd contamination (Fig. 4B). Cd toxicity caused a slight decrease in soluble protein content while SPD spraying increased soluble protein content in purslane leaves. On the other hand, Cd, at higher concentrations, exhibited a strong suppression on insoluble protein synthesis. The soluble protein content reached its maximum level (about 2.4 times over the control level) when purslane
plants sprayed with 0.5 mM of SPD spraying. High levels of Cd accumulation in different parts of the plant may interfere with plant growth by reducing the activity of antioxidant enzymes, closing the stomata, and inhibiting nutrient uptake (Parrotta et al., 2015; Paunov et al., 2018; Guo et al., 2019). The amount of leaf protein was greatly affected by Cd treatments. When purslane plants were grown at 30, 60, and 90 mg.kg$^{-1}$ of Cd, root protein content was declined by 14.5, 27.3, and 34.5%, respectively (Fig. 4B).

The decrease in protein content may be due to increased protein degradation or reduced synthesis (Balestrassi et al., 2003). The results indicated that the toxic effects of high levels of Cd on the mechanisms of protein synthesis resulted from decreased protease activity (Lee et al., 1976).

**Proline activity:** The results of the present experiment indicated an increase in proline content in response to an increase in cadmium concentration (Fig. 5A). Also, SPD increased the proline content, particularly in inoculated plants with fungi (Fig. 5B). Proline acts as an osmolyte in stress conditions and may increase antioxidant enzyme activity and thereby minimize the side effects of oxidative stress (Malhotra et al., 2017; Khamsuk et al., 2018).

In another finding, Islam et al. (2009) reported that proline acted as a growth regulator and maintains osmotic in plants, also the ability to protect cells from the accumulation of ROS. Proline is thought to be able to reduce the negative effects of cadmium toxicity on plant growth. Proline accumulates in plant tissues in response to exposure to heavy metals stress (Teklic et al., 2008; Radic et al., 2010). Despite the very low percentage of proline from the total free amino acid in the plant, it has been regarded as one of the important osmotic and common metabolites that are found in the cellular system exposed to stress (Matysik et al., 2002; Liang et al., 2013). Pal et al. (2015) and Szalai et al. (2017) have reported an increase in proline content by Put treatment. Sharma et al. (1998) supposed that proline may act as a metal Celator.

**Hydrogen peroxide (H$_2$O$_2$):** The results showed that the amount of H$_2$O$_2$ production in plants inoculated with TL was affected at all levels of Cd (Fig. 6A). However, this change was less obvious in seedlings treated with SPD. On the other hand, H$_2$O$_2$ production was higher at all cadmium levels in non-inoculated plants (Fig. 6B, C). Also, H$_2$O$_2$ production decreased significantly with increasing SPD concentration in all the studied plants. Also, it has been previously reported that *Trichoderma* enhances antioxidant enzyme activities and reduces reactive oxygen species

![Fig. 4](image1.png)

Fig. 4- Effects of exogenous SPD (0, 0.5, 1 mM) with *Trichoderma* (A), on proteine content free in purslane seedlings under four concentrations (0, 30, 60, 90 mg.kg$^{-1}$) CdCl$_2$ stress (B). Different letters indicate significant differences (P≤0.05).

![Fig. 5](image2.png)

Fig. 5- Effects of four concentrations (0, 30, 60, 90 mg.kg$^{-1}$) CdCl$_2$ stress (A) and exogenous spermidine (0, 0.5, 1 mM) with *Trichoderma* (B) on prolin content in purslane seedlings. Different letters indicate significant differences (P≤0.05).
Trichoderma and spermidine improve cadmium tolerance

Application of Cd alone produces no change in the H$_2$O$_2$ content, contrasting with the results observed by Tang et al. (2005) in maize. Polyamines are likely to be effective in protecting plant tissues from oxidative damage by increasing the activity of antioxidant enzymes (Groppa et al., 2007, 2008; Liu et al., 2015).

Bioaccumulation of heavy metals: The interaction effects between TL and SPD on uptake and translocation of Cd in purslane are shown in Figure 7. Cd was accumulated in both plant shoots and the roots at all Cd concentrations. This result clearly shows the Cd ability to accumulate and transport in all parts of the purslane (Fig. 7). On the other hand, the highest levels of Cd accumulation were obtained in purslane shoots contrary to the pattern of accumulation heavy metal by other plants. Similarly, Linger et al. (2005) has reported Cd accumulation in Cannabis sativa and Hibiscus cannabinus plants. This abundance amount of Cd accumulation in the aerial part cannot be explained solely by the adsorption of active ions. Immobilization, by attaching to cell walls, may play a minor role (Chakravarty and Banerjee, 2012). Although, plants inoculated with Trichoderma had no significant difference in Cd values in the aerial and terrestrial part of the plant, spraying of 0.5 and 1 mM of SPD in leaf tissues caused a significant difference in the amount of Cd in those organs. In fact, under the application of spermidine, the amount of cadmium accumulation in the aerial and terrestrial portions of Portulaca oleracea decreased in both non-co-existent and coexisting plants with TL.

The highest Cd accumulation in the shoot (318.5 mg.kg$^{-1}$ of DW) was measured when purslane control plants were grown at 90 mg.kg$^{-1}$ of Cd with no SPD spraying. The effects of TL fungi and SPD on Cd concentration in shoot, root and soil were shown in Figure 7. Inoculation of plants with TL fungi showed varied changes in Cd concentration in the soil. The Cd content in shoot increased at 30 mg.kg$^{-1}$ of Cd while decreased at 90 mg.kg$^{-1}$ of Cd as compared to the control. The concentration of Cd in shoot and root significantly increased with increasing of Cd toxicity, however, SPD spraying particularly in the Low-Cd concentration significantly decreased Cd accumulation. These results were similar to those obtained by Hsu and Kao (2007), which observed a decrease in Cd uptake by rice leaves treated with SPD. Also, these results were similar to those obtained by Hsu and Kao (2007), which observed a decrease in Cd uptake by rice leaves treated with SPD. It seemed that SPD was able to protect Cd-induced oxidative damage and this protection was related to the reduction of H$_2$O$_2$ generation and Cd uptake.

The highest Cd accumulation was obtained in the shoot of the plants grown on soil contaminated with 60 and 90 mg/kg Cd in the absence of spermidine foliar application (Fig. 7A).
Fig. 7 - Effects of exogenous SPD (0, 0.5, 1 mM) and *Trichoderma* on distribution of absorbed cadmium in the shoot (A-C), root (D-F), soil (G-I) and bioaccumulation factor (J-L) in purslane seedlings under four concentrations (0, 30, 60, 90 mg.kg⁻¹) CdCl₂ stress. Similar letters indicate the absence of significant differences (P≤0.05).

(shoot) and terrestrial (root) parts, and the shoot/root ratio are shown in Table 1. Plants grown under both levels of SPD had lower Cd in the shoot than the control.

A significant interaction effect was obtained (P≤0.05) between Cd and SPD in terms of Cd concentration in the shoot and root parts. The decrease of the shoot Cd concentrations triggered by elevated levels of SPD was more substantial in 30 mg.kg⁻¹ of Cd concentration with 1 mM of SPD (Fig. 7). Our results
confirmed that purslane possesses a high ability to accumulate Cd in its tissues, as in previous studies reported by Coakley et al. (2019) on plants such as *Impatiens glandulifera* in the environment contaminated with Cd.

On the other hand, *Trichoderma* inoculation improved purslane phytoremediation efficiency. Cd concentration ratios of shoots to roots are shown in Table 1. This value was greater than one. Our results showed that purslane especially when inoculated with TL and were treated with spermidine could be recommended as a suitable species for phytoremediation of Cd contaminated soils.

**Conclusions**

In the present study, we examined the effects of *Trichoderma* fungi coexistence and the application of polyamine spermidine on soil purification from cadmium contamination in the purslane plant. Our results showed that the Purslane plant could be recommended as a suitable species for soil purification from heavy metal such as cadmium due to the proper accumulation of Cd in its branches, especially when inoculated with TL. In other words, it increased the *Portulaca oleracea* phytoremediation in cadmium-contaminated soil. Also, SPD suppressed H$_2$O$_2$ production and improved proline content in plants under Cd stress. In general, the synergistic effect between the TL fungi and SPD could be an appropriate strategy to improve Cd phytoremediation efficiency in purslane plants.

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**References**


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**Table 1 - The shoot/root cadmium concentration ratio±standard error (MS/MR) in the treatment Trichoderma and levels of the SPD**

<table>
<thead>
<tr>
<th>Spermidin</th>
<th>Pretreatment</th>
<th>Cd0</th>
<th>Cd30</th>
<th>Cd60</th>
<th>Cd90</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Control</td>
<td>0.63±0.04*</td>
<td>1.13±0.05*</td>
<td>2.03±0.06*</td>
<td>1.59±0.04*</td>
</tr>
<tr>
<td></td>
<td>Trichoderma</td>
<td>0.10±0.02*</td>
<td>2.79±0.26*</td>
<td>2.04±0.08*</td>
<td>1.25±0.09*</td>
</tr>
<tr>
<td>SPD 0.5</td>
<td>Control</td>
<td>0.45±0.04*</td>
<td>1.54±0.07*</td>
<td>2.29±0.10*</td>
<td>1.23±0.04*</td>
</tr>
<tr>
<td></td>
<td>Trichoderma</td>
<td>0.31±0.04*</td>
<td>0.62±0.02*</td>
<td>1.12±0.08*</td>
<td>1.31±0.04*</td>
</tr>
<tr>
<td>SPD 1</td>
<td>Control</td>
<td>0.52±0.11*</td>
<td>1.39±0.04*</td>
<td>2.41±0.12*</td>
<td>1.32±0.03*</td>
</tr>
<tr>
<td></td>
<td>Trichoderma</td>
<td>0.73±0.12*</td>
<td>1.19±0.11*</td>
<td>2.39±0.09*</td>
<td>1.25±0.04*</td>
</tr>
</tbody>
</table>

Means with common letters in the LSD test are not significantly different from each other (P<0.05)


