Enhancement of Lead Phytoremediation by Perennial Ryegrass (Lolium perenne L.) Using Agent of Streptomyces pactum Act12

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Abstract

It has been well recognized that microbes can assist phytoremediation of lead (Pb) by promoting plant growth and metal uptake. However, little is known about the potential of soil inoculation with actinomycetes to enhance Pb uptake and translocation in plants, particularly in semi-arid water-deficient areas. This study was focused on exploring the resistance of a soil actinomycete strain (Streptomyces pactum Act12) to Pb and its effects on plant growth, antioxidant responses, Pb uptake and translocation in perennial ryegrass (Lolium perenne L.). Special attention was given to experimental conditions in semi-arid area in Northwest China. A fermentation culture of Act12 was applied in powder form to soil with or without Pb treatment (0–1,000 mg kg⁻¹), and ryegrass plants were immediately grown in this soil, in pots, for 60 days under 20% relative soil moisture. Act12 showed tolerance to up to 1,200 mg Pb L⁻¹ in plate culture and well colonized the soil containing less than 500 mg Pb kg⁻¹. Under Pb stress, inoculated plants had higher biomass with greater plant height and root tiller number than the uninoculated controls. Additionally, higher catalase, superoxide dismutase and peroxidase activities were detected in leaves of inoculated plants under Pb stress. Inoculating soil with Act12 significantly increased Pb concentrations and uptake in plants grown in soil containing 200 to 1,000 mg Pb kg⁻¹. The translocation and bioconcentration factors of inoculated plants were 10.5–36.2% and 37.3–133.1% higher than uninoculated plants, respectively. Streptomyces pactum Act12 can enhance Pb phytoremediation by perennial ryegrass and its powder form facilitates the use in semi-arid areas.

Keywords: Lead contamination; Phytoremediation; Microbial agent; Streptomyces pactum; Lolium perenne

Introduction

Lead (Pb) is a major pollutant in contaminated soil under long-term sewage irrigation, abuse of fertilizer, and industrial activities [1,2]. Excessive Pb results in reduced soil fertility and health, affecting plant growth and leading to reduced crop production [3]. Moreover, Pb is the most toxic pollutants to living organisms and Pb exposure is related to the etiology of cardiovascular, renal, nervous, and blood diseases in human [4]. It is therefore crucial to develop effective measures for remediating Pb-contaminated soils.

Phytoremediation is a low-cost, efficient, environmentally friendly method for eliminating toxic metals from soil. The process involves using metal-tolerant plants to extract contaminants from soil and to accumulate these contaminants in harvestable tissues [5]. Plants with great potential for metal phytoextraction include black nightshade (Solannum nigrum L.) [5], rape (Brassica napus L.) [6], and perennial ryegrass (Lolium perenne L.) [7]. However, slow plant growth, low biomass, and heavy-metal toxicity restrict the efficiency of phytoremediation [8,9].

Microbes can assist in the process of phytoremediation by promoting plant growth and metal uptake from soil. The microbial mechanism of promoting plant growth is to facilitate phytohormone (e.g., indole acetic acid) and siderophore production, nitrogen fixation, minerals solubilization, and/or nutrient transformation in plants [10]. Additionally, microbes can increase the activity of antioxidant enzymes and the production of protective substances in plants, thereby protecting them from phytotoxic substances under metal stress and ultimately improving the efficiency of phytoremediation [11,12].

Plant–microbe interactions utilized in phytoremediation include mutually beneficial symbiotic associations of plants with growth-promoting bacteria and mycorrhizal fungi [13]. The available bacteria have been studied extensively under optimal laboratory conditions with 60%–80% relative soil moisture [6,14]. It is unclear whether inoculated plants can maintain an efficient phytoremediation of heavy metals in soil under water deficiency, e.g., 10%–30% relative soil moisture in extensive mainland of semi-arid areas in China [15,16]. Moreover, laboratory-cultivated bacteria or spore suspension has been frequently used for microbe-assisted phytoremediation. The liquid form of microbial agent was time-consuming to prepare and difficult to apply in large areas.

Actinomycetes have been reported less frequently than bacteria and mycorrhizal fungi to enhance Pb uptake and translocation in plants. For example, sorghum plants inoculated with Streptomyces mirabilis P16B-1 exhibit increased growth and dry weight, suggesting that this strain may enhance phytoremediation by sorghum (Sorghum bicolor) [17]. A soil actinomycete strain, designated Streptomyces pactum Act12, was recently isolated from the Qinghai-Tibet Plateau [18], and S. pactum Act12 demonstrated beneficial effects on crop growth and resistance to plant diseases in pot experiments and field trials [19,20]. Given its

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origin from an extreme environment (drought, low-temperature, high-latitude), it is hypothesized that S. pactum Act12 would increase the efficiency of Pb phytoremediation by plants under water deficiency in semi-arid regions.

In this study, pot experiments were conducted using perennial ryegrass (Lolium perenne L.), a cultivated perennial herbaceous plant capable of Pb uptake and transport from the soil [7]. The resistance of S. pactum Act12 to Pb was examined and its effects on perennial ryegrass under Pb stress in a semi-arid area of natural water deficiency were then assessed. The study establishes an approach for actinomycete-assisted phytoremediation of Pb-contaminated soil using a powder form of microbial agent, which may facilitate the use in water-deficient fields of arid or semi-arid areas.

Materials and Methods

Soil, strain and plant

Top soils of 0-30 cm in depth were sampled in a local farmland in Xi’an, Shaanxi Province, China (34°09′24″N, 108°56′36″E). The soil represents a typical cinnamon soil in semi-arid area, Northwest China. Heavy metal contamination has occurred in the surface soil layer due to long-term irrational use of chemical fertilizers and pesticides [21], while the individual potential risk index of Pb content ranks the strong ecological hazard level [22]. After collection, the soil samples were air-dried, sieved using a 5-mm nylon sieve before physico-chemical characterization. The basic soil properties were organic matter of 14.57 g kg\(^{-1}\), total nitrogen of 1.27 g kg\(^{-1}\), nitrate nitrogen of 14.43 mg kg\(^{-1}\), total phosphorous of 0.49 g kg\(^{-1}\), Olsen-phosphorous of 15.82 mg kg\(^{-1}\), total potassium of 240 mg kg\(^{-1}\), available potassium of 392.80 mg kg\(^{-1}\), Pb of 85.62 mg kg\(^{-1}\), zinc of 78.23 mg kg\(^{-1}\), bulk density of 1.21 g cm\(^{-3}\), pH of 6.77 (1:1 w/v water), and soil moisture of 20.56% (fresh soil).

Streptomyces pactum Act12 was provided by the Laboratory of Microbial Resources, College of Resources and Environment, Northwest A&F University (Yangling, China) [18]. The agent Act12 was prepared by solid-state fermentation and applied in powder form (5.3 × 10\(^{10}\) CFU g\(^{-1}\)) [23]. The viable Act12 count was performed for colony forming units by the dilution-plate method on Gauze’s No. 1 agar supplemented with antibiotics which were specific for the identification of actinomycete [24].

Perennial ryegrass (Lolium perenne L.) seeds were purchased from the Xin Nong Feng Agricultural Technology Research Institute (Beijing, China). Full-grain seeds of uniform size (1,000-grain-weight: 2.2 g) were surface-disinfected with 0.5% NaClO solution (10 min, 25°C) and thoroughly rinsed with distilled water before sowing.

Pot experiments

Dry soil sample was treated with Pb at five levels: 0, 200, 300, 500 and 1,000 mg Pb kg\(^{-1}\). A Pb solution was prepared by dissolving analytical-grade Pb(NO\(_3\))\(_2\) in distilled water. Soil moisture content was adjusted to approximate 20%, close to the soil moisture content in natural farmland of semi-arid regions [25]. Prior to the experiment, Pb-treated soil subsamples were kept in darkness at 20°C for 3 months to allow equilibrium to be reached between metal and soil.

Pot experiments were performed in two treatments: Act12 inoculation and non-inoculation. Each treatment with the five Pb concentrations had three replications respectively. Thirty plastic pots (diameter 23 cm, depth 21 cm) were filled with Pb-treated soil using 3 kg per pot. For inoculation treatment, Act12 was pre-mixed with soil at 1.5 g kg\(^{-1}\) on a dry soil basis and placed into a pot. Non-inoculation treatment was prepared with Pb-treated soil without Act12 and provided as the control.

Seeds were sown in 1-cm deep holes (50 seeds per pot) in late March. Three-week-old seedlings were thinned to 15 plants per pot and the pots were then randomly arranged in a plastic greenhouse. The plants were watered once per day to maintain approximately 20% soil moisture content. The growth conditions were recorded and plants were carefully harvested after 60 d of growth.

Rhizosphere soil was removed from plant roots by gently shaking and collected for bacteria determination. Plant samples were washed with deionized water before measuring growth parameters including plant height, tiller number, and fresh weight. Ten gram of fresh leaf samples were excised and frozen in liquid nitrogen for physiochemical analysis. Plant roots and shoots were deactivated at 105°C for 30 min, followed by heating at 80°C for 24 h for Pb analysis. Pot soils were dried and passed through a 0.075-mm sieve and used to determine the bioavailable and residual Pb contents in the soil.

Pb tolerance analysis of actinomycetes strain: Pb resistance of S. pactum Act12 was analyzed using phosphate-poor morpholine propane sulfonic acid (MOPS) medium with 0.1% glucose as carbon source [26]. Act12 was streaked on MOPS agar plates containing gradient concentrations of Pb\(^{2+}\) from 0 to 1500 mg L\(^{-1}\) (100 mg L\(^{-1}\) interval). The culture was incubated at 28°C for 7 d. The minimum inhibitory concentration (MIC) of Pb was defined as the metal ion concentration that completely inhibited the growth of Act12 [6].

Sample analysis: Dry pot soil samples were ground in an agate mortar and digested in boiling aqua regia for 2 h. The digested sample was adjusted to 50 mL with distilled water, before Pb analysis using the PerkinElmer Optima 5300DV ICP-OES system [27]. Bioavailable Pb in the soil was extracted by EDTA [27]. Dry plant samples were ground into a powder and digested in a mixture of HNO\(_3\) and HClO\(_4\) (4:1, v/v). Pb concentration in the digested sample was determined by ICP analysis [28].

Fresh leaves were cleaned and the leaf veins were removed. Leaf superoxide dismutase (SOD) activity was assayed using the nitroblue tetrazolium reduction method [29]. Catalase (CAT) activity was assayed by measuring the decomposition of hydrogen peroxide as reflected by change of absorbance at 240 nm [30]. Peroxidase (POD) activity was analyzed in accordance with Polle et al. [31].

The number of viable Act12 in the rhizosphere soil was determined by serial dilution on MOPS plates containing 500 mg L\(^{-1}\) Pb. The inoculated plates were incubated at 30°C for 7 days. With account taken of Pb-resistant microorganisms and the appearance of characteristic colonies from the inoculated samples, only gray compact spiral colonies morphologically similar to colonies of Act12 were considered as reisolates of Act12 [18].

Data analysis: For the parameters tested, rate of increase (ΔCtrl%) was obtained as the percentage of the difference between Act12 inoculation and non-inoculation treatments to the non-inoculation control at the same Pb level. Translocation factor (TF) was calculated as the ratio of Pb concentration in shoots to that in roots. Bioconcentration factor (BCF) was calculated as the ratio of Pb concentration in plant (roots and shoots) to that in soil.

Average values of mean ± SD (n = 3) are presented. Significance of difference between treatments was determined by DUNCAN test at α = 0.05. Statistical analysis was performed using SPSS 20.0 (IBM...
SPSS, Somers, NY, USA). An asterisk (*) denotes significant difference between Act12 inoculation and uninoculated control within the same soil Pb treatments (P < 0.05). Different lower-case letters indicate significant difference of Act12 inoculation or uninoculated control between different soil Pb treatments (P < 0.05).

**Results**

**Pb tolerance of Act12**

*S. pactum* Act12 demonstrated high Pb resistance in agar medium and the MIC of Pb to Act12 was up to 1,200 mg Pb L⁻¹. Plate culture results showed that the strain had good growth in the presence of 0–800 mg Pb L⁻¹, while the growth was slower in the presence of 800–1,200 mg Pb L⁻¹.

After 60 days of cultivation in pots, Act12 was recovered from the rhizosphere of ryegrass plants with a high probability. Distinctive gray compact spiral colonies that morphologically resembled those of the original strain were considered as Act12 (Figure 1a). In the non-inoculated soil samples, no colonies resembling Act12 were found. The maximum Act12 count of the soil after 60 days of ryegrass growth was 5.4±5×10⁶ CFU g⁻¹ at 500 mg Pb kg⁻¹. The viable count of Act12 was reduced to a minimum of 0.90±10⁴ CFU g⁻¹ at 1,000 mg Pb kg⁻¹ (Figure 1d).

**Plant growth and biomass**

The growth of ryegrass plants was affected by Pb stress (Figure 2). As soil Pb level increased, plant height decreased in both inoculated and uninoculated treatments (Figure 1c); however, shoot fresh weight, root fresh weight, and tiller number initially increased before decreasing, with the maximum value at 300 mg Pb kg⁻¹.

Act12 inoculation had positive effects on the growth of ryegrass plants under Pb stress (Figure 2). Plant height, root fresh weight, and shoot fresh weight were significantly higher in inoculated plants than in the uninoculated controls, regardless of soil Pb level (P < 0.05). The greatest increases in plant height and root fresh weight were 12.3% and 94.6%, at 300 mg Pb kg⁻¹, respectively. The maximum increase in shoot fresh weight was 29.6% at 1,000 mg Pb kg⁻¹. Root tiller number significantly increased at all Pb levels except 500 mg Pb kg⁻¹; the greatest increase was 44.2% at 200 mg Pb kg⁻¹.

**Leaf antioxidant enzyme activity**

The leaf antioxidant enzyme activity in ryegrass plants was altered by the presence of Pb in soil (Table 1). As soil Pb level increased, leaf CAT and POD activities both decreased; however, SOD activity initially increased and peaked at 500 mg Pb kg⁻¹, followed by decreases.

Act12 inoculation positively affected leaf antioxidant enzyme activity in ryegrass plants (Table 1). Leaf CAT, SOD, and POD activities were all higher in inoculated plants than in the uninoculated controls, regardless of Pb level, especially at high soil Pb levels (500 to 1,000 mg Pb kg⁻¹), where inoculated plants showed significantly increased enzyme activities (P < 0.05). The greatest increase in CAT and SOD activities were 70.3% and 58.4% at 1,000 mg Pb kg⁻¹, respectively, while a 94.2% increase in POD activity was observed at 500 mg Pb kg⁻¹.

**Lead bioavailability in soil and Pb accumulation and uptake in plants**

The bioavailable soil Pb content in pot soil without Pb treatment was 32.1±3.51 mg kg⁻¹. Inoculation with Act12 significantly increased bioavailable soil content (8.4-37.2%) in a dose-dependent manner compared with the non-inoculation controls (P < 0.05) (Figure 1b). The greatest increase in response to Act12 inoculation was observed at 500 mg Pb kg⁻¹ (Table 2).

Pb uptake and accumulation by ryegrass plants varied in a dose-dependent manner. As soil Pb levels increased, there was an increase in root and shoot Pb concentrations and uptake in both inoculated and uninoculated plants. Act12 inoculation positively affected Pb uptake and accumulation under Pb stress. Both shoot and root Pb concentrations and uptakes were significantly higher in inoculated plants than in the uninoculated controls at 200 to 1,000 mg Pb kg⁻¹ (P < 0.05). For the inoculated plants, there were 61.7–130.4% (shoots) and 34.8–107.2% (roots) increases in Pb concentration, and 33.6–146.3% (shoots) and 36.4–173.4% (roots) increases in Pb uptake, compared with the uninoculated controls. The greatest increase in Pb concentrations and uptakes in response to Act12 inoculation was observed at 300 mg Pb kg⁻¹ (Table 2).

**Phytoremediation efficiency**

As soil Pb levels increased, TF decreased, while root BCF increased in both inoculated and uninoculated plants. Act12 inoculation promoted the translocation of Pb from roots to shoots, which resulted in a 10.5–36.2% increase in TF compared with the uninoculated controls (P < 0.05). Meanwhile, inoculation with Act12 enhanced Pb absorption by roots and shoots from the soil, resulting in 37.3–109.4% increase in root BCF and 62.7–133.1% increase in shoot BCF (P < 0.05). The greatest increase in TF was observed at 500 mg Pb kg⁻¹, while the greatest increases in root BCF and shoot BCF were observed at 300 mg Pb kg⁻¹ (Table 3).

**Discussion**

The role of *S. pactum* Act12

Microbes isolated from extreme environments provide a potential source for phytoremediation in contaminated soils under adverse conditions. Strain Act12 was isolated from a unique environment (drought, low-temperature, and high-latitude) on the Qinghai-Tibet Plateau [18]. Owing to the long-term adaption, Act12 may have evolved versatile mechanisms to cope with adverse stress. In the present study, *Streptomyces pactum* Act12 demonstrated high resistance to Pb (MIC = 1,200 mg Pb L⁻¹) in plate culture. As has been reported [32], processes of biosorption, reduction, biomineralization, extracellular binding by chelators, efflux by transport systems, and/or intracellular binding of metals are all involving as the role of *Streptomycetes*.

The rhizosphere, as an interface of soil and plant, supplies a niche for metal-tolerant bacteria that are essential for promoting plant-growth activities, alleviating plant stress and mobilizing or immobilizing metals [33]. Pot experiments showed that Act12 well colonized the soil containing less than 500 mg Pb kg⁻¹ and suffered a certain degree of inhibition at 1,000 mg Pb kg⁻¹ under 20% relative soil moisture, in agreement with the results of plate culture analysis for Pb tolerance. Taken together, the above results indicate that *S. pactum* Act12 is resistant to Pb and can thrive in soil with relatively low water content. This strain exhibits great potential for agricultural use in semi-arid regions.

It has been noted that low plant biomass could be a bottleneck in phytoremediation of heavy metal-contaminated soils [8]. Thus, plant-growth-promoting and heavy-metal-tolerant microbes are preferentially used to improve the efficiency of phytoextraction [12,14]. In the present study, *S. pactum* Act12 demonstrated plant-growth-promoting effects on ryegrass plants under Pb stress, as indicated by the greater root tiller number plant height and biomass (Figure 2). This is consistent with the results observed in cotton and ginseng [20,34]. The effect of Act12 may be related to the production of indole-3-acetic acid,
The plant growth-promoting effect of Act12 observed in the current study was more significant in soil at higher Pb levels. Similar results have been observed in *Streptomyces mirabilis* P16B-1, which promotes the growth of *S. bicolor* in a dose-dependent manner [17]. A combination of mycorrhiza and *Streptomycetes* also achieves a slight, albeit statistically significant increase in the biomass productivity of the same plant [36]. Previous studies have added the microbes as cell suspensions after laboratory cultivation for days and used them in soil under optimal water conditions (60%–80%). In the present study, Act12 was applied in powder form, which is convenient for preservation, maintenance of high activity, and application in large fields. With respect of soil condition, Act12 was tested in soil with 20% fields. With respect of soil condition, Act12 was tested in soil with 20%

### Table 1: Leaf antioxidant enzyme activity (catalase activity, CAT; superoxide dismutase, SOD; peroxidase activity, POD) of perennial ryegrass in Pb-treated soil inoculated with or without *Streptomyces pactum* Act12 (means ± SD, n = 3).

| Pb treatment (mg kg⁻¹) | CAT U g⁻¹ min⁻¹ | SOD U g⁻¹ FW | POD U mg⁻¹ min⁻¹ |
|------------------------|------------------|--------------|------------------|
| Control                | 2.63 ± 0.13      | 80.70 ± 5.72 | 113.25 ± 7.11    |
| Act12 200              | 3.14 ± 0.14      | 85.70 ± 4.41 | 146.02 ± 5.38    |
| μCtrl (%)              | 19.3%            | 6.2          | 16.8%            |
| Control                | 2.48 ± 0.23      | 95.69 ± 4.17 | 107.35 ± 3.72    |
| Act12 300              | 2.57 ± 0.22      | 104.22 ± 2.92| 113.25 ± 7.05    |
| μCtrl (%)              | 3.6              | 8.9%         | 5.5              |
| Control                | 2.33 ± 0.19      | 107.35 ± 5.72| 94.22 ± 1.86     |
| Act12 500              | 2.77 ± 0.12      | 113.82 ± 4.64| 115.69 ± 6.65    |
| μCtrl (%)              | 19.0%            | 6.0          | 22.8%            |
| Control                | 1.55 ± 0.24      | 135.05 ± 6.62| 67.70 ± 3.52     |
| Act12 1,000            | 2.52 ± 0.10      | 176.02 ± 6.63| 131.49 ± 7.70    |
| μCtrl (%)              | 62.4%            | 30.3%        | 94.2%            |

### Table 2: Pb concentration and uptake in perennial ryegrass plants, and bioavailable soil Pb in Pb-treated soil inoculated with or without *Streptomyces pactum* Act12 (means ± SD, n = 3).

| Pb treatment (mg kg⁻¹) | Pb concentration (mg kg⁻¹) | Pb uptake (μg pot⁻¹) | Bioavailable soil Pb (mg kg⁻¹) |
|------------------------|-----------------------------|----------------------|-------------------------------|
| Control                | Shoot 1.53 ± 0.13            | Root 6.35 ± 0.48     | Shoot 31.71 ± 3.89            |
| Act12 200              | 6.51 ± 0.82                 | 35.44 ± 2.90         | Root 50.23 ± 3.69             |
| μCtrl (%)              | 5.0                          | 11.8                 | Act12 157.45 ± 15.93          |
| Control                | 19.70 ± 1.59                | 131.34 ± 16.95       | SOD U g⁻¹ FW 311.98 ± 35.27   |
| Act12 300              | 26.51 ± 2.16                | 174.45 ± 13.95       | Root 85.32 ± 2.93             |
| μCtrl (%)              | 61.7%                       | 34.8%                | Ctrl (%) 163*                 |
| Control                | 27.27 ± 2.57                | 152.65 ± 12.34       | Ctrl Act12 123.98 ± 8.95      |
| Act12 500              | 56.51 ± 5.89                | 376.02 ± 29.18       | SOD U g⁻¹ FW 837.42 ± 67.95   |
| μCtrl (%)              | 130.4%                      | 107.2%               | Root 165.86 ± 3.99            |
| Control                | 63.25 ± 5.26                | 281.56 ± 12.40       | Ctrl Act12 231.06 ± 22.33     |
| Act12 1,000            | 86.83 ± 6.93                | 423.68 ± 35.63       | SOD U g⁻¹ FW 371.05 ± 2.88    |
| μCtrl (%)              | 89.6%                       | 37.3%                | Root 13.8 13.8 8.4*           |
| Control                | 60.5 ± 2.25                 | 876.86 ± 106.84      | Ctrl 1682.01 ± 98.57          |
| Act12 200              | 107.35 ± 3.72               | 1195.69 ± 112.15     | SOD U g⁻¹ FW 370.87 ± 11.76   |
| μCtrl (%)              | 163%                        | 146.3%               | Root 2530.10 ± 231.38         |
| Control                | 2.52 ± 0.10                 | 135.05 ± 112.15      | Ctrl 418.71 ± 16.33           |
| Act12 300              | 62.7*                       | 113.25 ± 7.11        | SOD U g⁻¹ FW 418.71 ± 16.33   |
| μCtrl (%)              | 58.4%                       | 117.9%               | Root 13.8 13.8 8.4*           |

### Table 3: Average translocation factor (TF) and bioaccumulation coefficient (BCF) of perennial ryegrass plants in Pb-treated soil inoculated with or without *Streptomyces pactum* Act12 (means ± SD, n = 3).

| Pb treatment (mg kg⁻¹) | TF (%) | Root BCF (%) | Shoot BCF (%) |
|------------------------|--------|--------------|---------------|
| Control                | 24.11 ± 1.45 | 24.76 ± 1.61 | Act12 24.64 ± 0.57 |
| μCtrl (%)              | 2.7    | 7.55 ± 0.20  | Act12 11.3%    |
| Control                | 27.91 ± 0.09 | 33.45 ± 0.51 | 9.9%          |
| Act12 200              | 6.97 ± 0.12 | 9.46 ± 0.03  | 35.8%         |
| μCtrl (%)              | 1.82%  | 2.3          | Act12 50.5%    |
| Control                | 22.06 ± 0.2 | 24.6 ± 0.57  | 11.3%         |
| Act12 300              | 7.16 ± 0.05 | 14.99 ± 0.09 | 109.4%        |
| μCtrl (%)              | 1.58%  | 109.4%       | Act12 3.68 ± 0.1 |
| Control                | 14.47 ± 0.17 | 19.72 ± 0.18 | 36.2%         |
| Act12 500              | 11.00 ± 0.10 | 15.10 ± 0.18 | 37.3%         |
| μCtrl (%)              | 1.59%  | 2.98%        | Act12 2.98 ± 0.04 |
| Control                | 16.4 ± 0.24 | 18.13 ± 0.07 | 10.5%         |
| Act12 1,000            | 12.43 ± 0.33 | 18.86 ± 0.15 | 51.7%         |
| μCtrl (%)              | 2.04%  | 2.04%        | Act12 3.42 ± 0.03 |

siderophores and 1-aminocyclopropane-1-carboxylic acid deaminase, and/or solubilization of inorganic phosphate [6,12]. Additionally, metabolites such as organic acids, amino acids, and vitamins, as well as extracellular hydrolytic enzymes produced by branched mycelia [35], can stimulate root uptake of nutrients and subsequent plant growth. Moreover, actinomycetes are beneficial to rhizosphere soil microlora and thus could indirectly contribute to plant growth [20].
Figure 1: *Streptomyces pactum* Act12 colonies (a), powder form agent (b), the growth of perennial ryegrass plants in Pb-treated soil with or without Act12 inoculation (c) and the count of *Streptomyces pactum* Act12 after 60-d survival in Pb-treated rhizosphere soil with or without inoculation. (means ± SD, n = 3).

Figure 2: Growth parameters of perennial ryegrass in Pb-treated soil inoculated with or without *Streptomyces pactum* Act12 (a), plant height (b), tiller number (c), root fresh weight and (d) shoot fresh weight of perennial ryegrass after 60-d survival. (means ± SD, n = 3).
in semi-arid water-deficient areas. Therefore, the powder form of Act12 will facilitate the use in heavy metal remediation across semi-arid regions.

In addition, lead can produce physiological stress and induce the production of superoxide anion (O2−) and hydrogen peroxide (H2O2) in perennial ryegrass [30], which are harmful to plant cells. Plants can produce antioxidant enzymes to reduce the oxidative stress and maintain the resistance against heavy metals [37]. The current results show that soil inoculation with strain Act12 increased leaf CAT, SOD and POD activities in ryegrass plants under Pb stress compared with the un-inoculated control. The activities of the three enzymes markedly increased in inoculated plants grown in soil at higher Pb levels (500–1,000 mg kg−1). SOD can convert active and toxic superoxide radical into H2O2; CAT and POD then eliminate H2O2 through the conversion to H2O and O2 [37]. Improving the antioxidant enzyme activity thus can increase the detoxification of heavy metal stress by plants [38].

At the cellular level, high levels of Pb in plants can pose phytotoxic effects by inducing excessive accumulation of reactive oxygen species. Act12 in the rhizospheric microbial communities might enhance the tolerance of ryegrass plants against Pb toxicity by producing plant growth-promoting factors such as indole acetic acid (IAA), siderophores, 1-aminoacyclopeptide-1-carboxylate (ACC) deaminase [39,40]. Additionally, Act12 might produce organic acids and amino acids that are high-affinity ligands for the chelation of Pb by cytosol. This mechanism could lower ionic Pb in cytoplasm, attenuate the phytotoxic effect, and then normalize the antioxidative response of plants, ultimately contributing to metal detoxification and tolerance [13,41].

Similarly, co-inoculation of lentil with Pb-resistant bacteria (A. tumefaciens, R. aquatilis, and Pseudomonas sp.) increases CAT, SOD, and POD activities in plants under Pb stress [42]. An endophytic fungus (EF0801) benefits rice growth under moderate Pb stress by enhancing CAT, SOD, and POD and reducing MDA levels [12]. A recent study has reported that the expression of genes encoding antioxidant enzymes is upregulated in plants to help alleviate metal toxicity [11]. These findings confirm that Act12 can enhance the tolerance of perennial ryegrass to Pb stress by enhancing antioxidant enzyme activities. This can be regarded as a defense mechanism against Pb toxicity, and thus beneficial to plant growth.

Furthermore, phytoextraction capacity is influenced by metal accumulation in plant tissues and biomass productivity [38]. The present study showed that Act12 inoculation increased Pb bioavailability in the contaminated soil and improved Pb concentrations and uptake in shoots and roots of perennial ryegrass plants under Pb stress (Table 2). Microbes can alter metal availability and mobility to the plant by acidification, chelation and oxidation-reduction reactions in the rhizosphere [43], ultimately contributing to metal accumulation in plants. Similarly, an endophytic Rahnella sp. JN6 was reported to increase the availability of Pb in soil and enhance Pb uptake by B. napus [6].

In the 300 mg kg−1 soil treatment, the Pb concentrations and uptake in Act12-inoculated plant showed greater than 100% increase relative to those in the uninoculated controls. The effect of Act12 is markedly stronger than that of the rhizobacterium Burkholderia sp. D54 (PGPR) in increasing Pb levels in ryegrass (11.3% increase in roots, no effects in shoots) [14]. Streptomyces can produce versatile secondary metabolites such as organic acids which were considered as the sources of protons for mobilization and metal-chelating anions to form complexes with metal cations [44,45].

The TF and BCF are two indices used to assess the efficiency of phytoextraction in Pb-contaminated soil [7]. The TF was less than 1.0 in ryegrass plants, as Pb is more likely to be absorbed and accumulate in roots than in shoots [30]. When we inoculated strain Act12 into Pb-treated soil, there were greater TF and BCF of Pb in both the roots and shoots of ryegrass plants than in the uninoculated control. This result suggested that soil inoculation with Act12 can promote the translocation of Pb from roots to shoots and boost Pb absorption by ryegrass plants from the soil. The current findings confirm that strain Act12 can improve the phytoremediation efficiency of perennial ryegrass.

**Prospects of lead phytoremediation in arid and semi-arid areas**

A number of studies claimed that lead phytoremediation can be enhanced by the assistance of beneficial bacteria [6,13]. These bacteria improve plant growth and increase plant biomass, in turn assisting phytoremediation by its growth-promoting traits. In addition, microbes can protect plants from phytotoxic substances under lead stress by increasing the activity of antioxidant enzymes and protecting the production of protective substances in plants. Additionally, lead uptake can be promoted by the transformation of lead into bioavailable and soluble forms through the action of bacteria. However, in arid and semi-arid area, the application of lead phytoremediation should pay special attention to the restriction of water deficiency as drought stress could result in low biomass production, low rates of metal removal and insufficiently metal uptake into plant tissue [46]. More significantly, the effective plant microbial remediation technologies suitable for Pb-contaminated soil in the arid and semi-arid area are not properly documented in the literature. This study has demonstrated that actinomycetes could assist Pb phytoremediation under water deficiency. It is reasonable to believe that the findings of this study will promote the further studies and potential application of Pb control in arid and semiarid areas, especially in China as it has 3.07×10 6 km2 in Northwest China while most of the areas are reported to be polluted during the last two decades of the industrialization/urbanization [1,2,21].

**Conclusions**

*Streptomyces pactum* Act12 exhibited high Pb tolerance and colonized the rhizosphere containing a wide range of Pb. Soil inoculation with Act12 in powder form promoted plant growth and enhanced Pb tolerance in ryegrass grown in Pb-treated soil in a semi-arid area of natural water deficiency by increasing antioxidant enzyme activities. Meanwhile, Act12 improved the bioavailability of Pb in the soil and improved Pb accumulation and uptake by ryegrass, achieving high bioconcentration and phytoremediation efficiency. *S. pactum* Act12 represents Pb-resistant microbial agent that can be used in phytoremediation in semi-arid water-deficient areas. The powder form of Act12 guarantees its convenience for transport and application. Field trials will be conducted to assess the effects of Act12 on phytoremediation and reveal its protective mechanisms under drought conditions.

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