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IMPROVING AGENTS FOR PHYTOREMEDIATION OF SOIL CONTAMINATED WITH PYMETROZINE INSECTICIDE

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ABSTRACT: The present work was designed to investigate the potential of using plantain (*Plantago major* L.) for the phytoremediation of pymetrozine contaminated soil. The use of soluble silicon dioxide (SiO₂), Tween 80, hydroxypropyl- β -cyclodextrin (HP β CD) and liquid humic acid (HA) for enhancing the availability and uptake of pymetrozine contaminated soil by *P. major* were evaluated. Results revealed that pymetrozine concentrations in soil with *P. major* reduced by 30.00 – 83.25% throughout 1 to 12 days exposure, compared with 12.50–61.90% in the soil of control. Pymetrozine uptake in the roots and translocated in the leaves of *P. major* to reach the maximum levels, 53.41 mg/kg and 58.08 mg/kg, respectively, after 4 days. The phytoremediation efficiency of *P. major* amended with SiO₂ was greater than that of other solubility-enhancing agents with respect to the removal of pymetrozine uptake in roots and translocation in leaves by about 170.84% and 322.83% compared with *P. major* roots and leaves alone, respectively within 4 days. The most-effective to least-effective supplements to use in combination with *P. major* for the purpose of pymetrozine phytoremediation in roots and leaves were found to be as follows: SiO₂> HA > HP β CD > Tween 80. This study indicates that SiO₂ can improve the efficiency of phytoremediation of pymetrozine.

Key words: Phytoremediation, Plantago major, pymetrozine, improving agents, soil.

INTRODUCTION

Pymetrozine was the compound of the pyridine azomethine family, representing a newly developed chemical class of insecticides (Shen et al., 2009). Pymetrozine is efficient toward aphids, whiteflies and plant hoppers in pest control programmes (Lashkari et al., 2007). Pymetrozine acts by interfering in the regulation of the nervous system for feeding behavior, which results in death of the insect due to starvation a few days after application (Guoging et al., 2009). EFSA (2014) showed that pymetrozine exhibits low to moderate persistence in soil under laboratory aerobic conditions. It degraded by hydroxylation of the methylene group of the triazine ring and by oxidation. Degradation under field conditions from 8 sites in Europe (Switzerland, France and Germany) resulted in pymetrozine dissipating with single first order DT_{50} of 19.6 – 183 d (5sites) and biphasic DT_{50} 3.81-10.3 d with associated DT_{90} 167 \geq 1000 d. Pymetrozine degraded from water mainly by distribution to the sediment. Single first order DT_{50} of pymetrozine in the whole systems ranged between 289 and 495 d.

Phytoremediation is an environmentally sound technology for pollution prevention, control and remediation. One medicinal herb is plantain (*Plantago major* L., Plantaginacea family). *Plantago major* has a wide geographic distribution through the temperate grasslands of the world and grows in a wide area of Europe, temperate regions of Asia and South Australia, North Africa and North America (Velasco-

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Lezama et al., 2006). It is a familiar perennial weed and may be found at roadsides, meadowlands, cultivated fields, waste areas, and even cracks in sidewalks and canal banks. The seeds and husks contain high levels of fiber; they expand and become highly gelatinous when soaked in water (Samuelsen, 2000; Sharifa et al., 2008). P. major can accumulate a variety of inorganic metals or metabolize a variety of organic compounds including imidacloprid, chlorpyrifos, diethyl and dioctyl phthalates, azoxystrobin and cyanophos (Romeh, 2010; Romeh, 2013; Romeh and Hendawi, 2013 : Romeh, 2014; Romeh, 2015a&b). Therefore, the goal of this work was to assess the capability of using P. major for the phytoremediation of pymetrozine -contaminated soil. The utilization of soluble silicon dioxide (SiO₂) and improving agents such as the surfactants, Tween 80, hydroxypropyl-\beta-cyclodextrin (HPBCD) and liquid humic acid (HA) for enhancing the bioavailability and uptake of pymetrozine contaminated soil by P. major were evaluated.

MATERIALS AND METHODS

Pesticide and Plant

Pymetrozine (Technical grade 95.00%), 6methyl-4-[(E)- pyridin-3-ylmethylideneamino]-2, 5-dihydro-1, 2, 4-triazin-3-one was acquired from Central Agricultural Pesticides Laboratory, Agricultural Research Center, Dokki - Giza, Egypt. The normal broadleaf plantain (P. major) was acquired as seedlings in phytoremediation experiment. Seeds of P. major grow naturally on meadow land in Zagazig University, Zagazig, Sharkia Governorate. After the seeds germinated, the seedlings (the age of the seedlings is 30 days and 10-12 cm height with 4 -6 leaves) were collected for the experiment.

Experimental Design

To assess the elimination of pymetrozine from the soil, two treatments were performed in this experiment, and each treatment consisted of five replicates: pymetrozine contaminated soil without plants, pymetrozine contaminated soil with *P. major* only (each pot contained one seedling of *P. major*). Experimental uptake was performed on soil in a pot experiment for 21-day exposure. Air-dried sieved clay loam soil (organic matter, 1.79%, pH 7.8, electric

conductivity 2.36) was obtained from Kamrona Village, Menia EL-Kamh district, Sharkia Governorate, Egypt, and then placed in plastic pots. The pots were provided with 0.5 kg of air dried soil. After planting, pymetrozine dissolved in water was spiked into the 150 ml of distilled water used for irrigation to obtain the original concentration of 20 mg/kg. The irrigation water containing pymetrozine was dropped into the pots with a caution to avoid the direct contact of plant leaves. Samples from exposed and control plants were collected through 1, 3, 7, 10, 14 and 21 days. Roots of plant from soil were rinsed in running tap water for 2 min and were blotted dry. The plants were dissected into individual roots and leaves then, 10 g of each leaves and roots and 20 g of soil were analyzed for the pesticide. All pots were watered with 50 ml tap water every 4 days or additionally watered when essential.

Enhancing Agents for Phytoremediation of Soil Contaminated by Pymetrozine

To assess the removal of pymetrozine from the soil, eight treatments each consisting of five replicates were performed, as follows: Autoclaved soil contaminated with pymetrozine without plants. Soil contaminated bv pymetrozine without plants. Contaminated soil with pymetrozine plus P. major only. Contaminated autoclaved soil with Pymetrozine containing P. major only. Contaminated soil with pymetrozine plus P. major and amended with soluble silicon dioxide (SiO_2) , these called silica, at 750 mg/l for a total concentration of 187.5 mg/kg. Contaminated soil with pymetrozine plus P. maior and amended with 2-Hydroxypropyl-beta-cyclodextrin (HPBCD) at 1.0% (Chen et al., 2010). Contaminated soil with pymetrozine plus P. major and amended with humic acid solution (HA) at 10 mg/l (humus WSG 90, produced by organist-Hungary). The latter concentration is reported to be the critical micelle concentration of HA (Guetzloff and Rice, 1994). Contaminated soil with pymetrozine plus P. major and amended with polyoxyethylene sorbitan monooleate (Tween 80) at 9.2 mg/l, corresponding to 0.5 critical micelle concentration (CMC), where the CMC of Tween 80 was determined as 13-45 mg/l (Edwards et al., 1991; Mitton et al., 2012).

In treatments (3)-(8), each pot contained one seedling of *P. major*. Each whole plant uptake experiment was performed in potted soil for 12 days as described above.

Residues Analysis

The system followed in this work depended on **OuEChERS** strategy depicted by Anastassiades et al. (2003) and Lehotay et al. (2005). Soil tests were homogenized sieved (2 mm mesh) and air-dried at room temperature. A 10 g of homogenised soil was weighed into a 50 ml polypropylene tube, then 5 ml of HPLC water was included and the mixture was shaken for 1 min with a vortex apparatus. After that 10 ml of acetonitrile was included (acidified with acetic acid 1%) and the mixture was shaken for 1 min by hand and for 1 min with a vortex apparatus. Four grams of anhydrous magnesium sulfate, and 1 g sodium chloride were added, and the mixture was instantly hand-shaken for 30 sec., and centrifuged for 5 min at 4000 rpm in a Sigma 2-5 rotator (Sigma, Steinheim, Germany). At that point, a clean-up dispersive solid phase extraction step was implemented by including the supernatant (7.5 ml, i.e. 1.33 g of soil for every ml), to a 15 ml polypropylene tube that contained 1.125 g of MgSO₄ (150 mg MgSO₄ per ml of concentrate) and 0.225 g of C_{18} (30 mg C_{18} per ml of concentrate), handshaken for 30 sec. and centrifuged for 5 min at 4000 rpm (Asensio-Ramos et al., 2010; Padilla-Sanchez et al., 2010). For the determination of pymetrozine, 1.0 ml of the final concentrate was determined by HPLC.

A 10 g of fine macerated plant tissue (roots or leaves) was weighed into a 50 ml polypropylene tube. Then, 10 ml of acetonitrile was added (acidified with acetic acid 1%) and the mixture was hand-shaken for 2 min. Four grams anhydrous magnesium sulfate, and 1 g sodium chloride, was added, and the mixture was immediately hand-shaken for 30 sec. and centrifuged for 5 min at 4000 rpm. The clean-up step was implemented by adding the supernatant (7.5 ml, *i.e.* 1.33 g of plant tissue per ml), to a 15 ml polypropylene tube that included 1.125 g of MgSO₄ (150 mg MgSO₄ per ml of extract), and 0.188 g of PSA (25 mg PSA per ml of extract), hand-shaken for 30 sec. and centrifuged for 5 min at 4000 rpm (Anastassiades et al.,

2003; Lehotay *et al.*, 2005). For the determination of pymetrozine residues, 1.0 ml of the final extract was analyzed by HPLC.

HPLC analysis

Soil, root, and leaf samples were analyzed for pymetrozine using high-performance liquid chromatography (HPLC) according to (Shen *et al.*, 2009). A 10 μ l aliquot of final extract was injected into HPLC system and determined using a C₁₈ reversed-phase column [250 mm x 4.6 mm (i.d.)] and eluted isocratically with a mobile phase of water and acetonitrile (15:85, V/V) at the rate of 1 ml/min. The UV detection was adjusted at 299 nm. The retention time (RT's) was 2.63 minutes under these conditions. The performance of HPLC method was tested by evaluating quality parameters, such as recovery values.

Recovery Samples

The extraction effectiveness of the analytical procedure was assessed by recovery experiments prepared in triplicate using the fortified blank leaves and roots of *P. major*, and soil samples at 0.5 mg/kg. The percent recoveries were respectively, 92.33, 90.12, and 88.5% in roots, leaves, and soils.

Data Analysis

The rate of degradation (K_r) and half-life $(t_{1/2})$ was obtained according to Gomaa and Belal (1975) and Ashour (1976). The rate of degradation (K_r) = 2.303 x slope. Half-life $(t_{1/2})$ = 0.693/K_r.

RESULTS AND DISCUSSION

Phytoremediation of Soil Contaminated with Pymetrozine

Results obtained during the present investigation revealed that all experimental sets containing plantain removed a high amount of pymetrozine. As shown in Table 1, pymetrozine concentrations in soil containing *P. major* declined by 30.00-83.25% during 1 to 12 days of the experimental period, compared with 12.50-61.90% in un-planted soil. The half-life value ($t_{1/2}$) of pymetrozine, estimated by first-order reaction, for soil plus *P. major* was found to be 5.65 days, compared with 8.58 days for

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Treatment	Days after application								
	1	2	4	6	8	12	T _{1/2} (days)	K _r (days)	AUC _s mg/kg (days)
In soil									
mg/kg	17.50	15.30	13.52	11.40	9.05	7.62	8.58	0.08	142.19
Loss (%)	12.50	23.50	32.40	43.00	54.75	61.90			
In soil with <i>P. major</i>									
mg/kg	14.00	11.52	9.71	7.87	5.66	3.35	5.65	0.12	97.45
Loss (%)	30.00	42.40	51.45	60.65	71.70	83.25			
In <i>P. major</i> roots									
mg/kg	38.74	45.56	53.41	42.91	37.69	22.58			
In <i>P. major</i> leaves									
mg/kg	13.58	44.95	58.08	37.72	28.13	16.06			
Total uptake	52.32	90.51	111.49	80.63	65.82	38.64			

Table 1. Dissipation of pymetrozine in soil planted with Plantago major L

 $T_{1/2}$, half-life; k_r, disappearance rate constant; AUCs, areas under the curve represent compound concentration during the period of study

soil alone (Table 1). Results with the disappearance rate constant (kr) values showed that pymetrozine had the highest k_r value and lowest $t_{1/2}$ in soil with *P. major*, while pymetrozine had the shortest k_r and longest $t_{1/2}$ in unplanted soil.

Pymetrozine concentrations (mg/kg) in various parts of P. major are found in Fig. 1. Concentrations of pymetrozine in the root were always higher than those in the leaves of the plant, with the exception of 4 days period; the translocation ratio was about 1.08 times higher than for pymetrozine contaminated soil with P. major roots (Fig. 1). In the roots of P. major, pymetrozine accumulated to reach the maximum levels through 4 days (53.41 mg/kg). Afterwards, concentration decreased gradually during the experiment (Table 1). Pymetrozine translocated into the leaves of P. major and reached the maximum through 4 days of treatment (58.08 mg/kg) then decreased until the end of exposure. Pymetrozine is highly moved in plants. It can be taken up from the soil as well as through the leaves (Fliickige et al., 1992). Pymetrozine is both systemic and translaminar, making it highly mobile within plants (Wyss and Bolsinger, 1997). From bioassays and autoradiographic techniques of pymetrozine, it has been shown that this systemic behavior originates not only from xylem but also from phloem mobility. After foliar application, the growing points of plants are protected by pymetrozine imports mainly from leaves. This indicates a high importance of phloem mobility for the systemic activity of pymetrozine for plant-sucking insects (Wyss and Bolsinger, 1997).

Combination of *P. major* and Agents that Increase the Pymetrozine Availability in Soil

Changes in the levels of pymetrozine in soils subject to various treatments were measured following 1 to 12 days (Table 2) to estimate (1) plant's capacity to remove pymetrozine, (2) the role of various agents in increasing pymetrozine accessibility. microorganisms, plants and combination of plants with microorganisms, which lead to the dissipation of pymetrozine in the soil. Different agents contributed in increasing pymetrozine availability and reached the maximum enhancing through 4 - 8 days; therefore, the removal percentage of pymetrozine in a control group (C) was compared with removal percentages in experimental treatments after 4 days.



Fig. 1. Uptake and translocation of pymetrozine by Plantago major L. in soil

In C, 24.70% of the pymetrozine was degraded through natural biotic losses, whereas degradation percentages in treatments T_1 , T_2 , and T₃ were 32,40%, 49.50%, and 51.45%, respectively. We examined the contribution of three factors in the degradation of pymetrozine: biotic dissipation or natural hydrolyzation, as in the control group (C), microbial dissipation (T_1) , remediation by *P. major* (T_2) , and a combination of microbial degradation and P. major remediation (T_3) (Table 2). Abiotic degradation was not considered in the treatments. The degradation of pymetrozine in the soil due to contribution by microorganisms (T_1) was 7.70%, while contribution by plant alone (T_2) and a combination of microorganisms and plants (T_3) resulted in dissipation amounts of 24.80 % and 26.75%, respectively. The results showed that a combination of microorganisms plus plants (T_3) is the most effective treatment for the dissipation of pymetrozine in soil, followed by plant alone (T_2) and then microorganisms (T_1) , compared with natural dissipation processes.

It could be concluded that enhancement in pymetrozine dissipation in phytoremediation system could be achieved possibly due to the degradation induced by effects of plant with microorganisms in rhizosphere (Cheng *et al.*, 2007). The microbial-enhanced phytoremediation offers much potential for the remediation of organic pollutants in the soil (Chen *et al.*, 2010).

Pollutant-degrading bacteria may accelerate plants adaptation to contaminants by detoxifying contaminated soils during direct mineralization of these organic contaminants (Escalante-Espinosa *et al.*, 2005). In addition to, plant exudates increase the density and activity of potential pollutant-degrading bacteria in the area surrounding the roots (Siciliano and Germida, 1998). Romeh (2010) found that short-rod gramnegative bacteria that isolated from the water solution containing *P. major* was able to induce 93.34% loss of imidacloprid as a source of both nitrogen and carbon through 48 hr.

Treatment	Days after treatments							
Amount added to the soil (20 mg/kg)	mg/kg	Removal (%)	Contribution (%)					
1 day		_ · ·						
C:In autoclaved soil	18.55	7.25	0.0					
T1: In soil	17.50	12.50	5.25 microorganisms (T_1-C)					
T2: Autoclaved soil with plantain	14.30	28.50	21.25 Plant (T ₂ -C)					
T3: Soil with plantain	14.00	30.00	22.75 Combination (T ₃ -C)					
T4: Soil with plantain+ SiO2	11.10	44.50	14.50 $SiO_2(T_4-T_3)$					
T5: Soil with plantain+ HPβCD	11.20	44.00	14.00 HPβCD (T ₅ - T ₃)					
T6: Soil with plantain+ HA	11.16	44.20	14.20 Humic acid $(T_6 - T_3)$					
T7: Soil with plantain +Tw 80	11.60	42.00	12.00 Tw 80 (T_7 - T_3)					
2 days								
C:In autoclaved soil	17.39	13.05	0.0					
T1: In soil	15.30	23.50	10.45 microorganisms (T1-C)					
T2: Autoclaved soil with plantain	12.00	40.00	26.95 Plant (T2-C)					
T3: Soil with plantain	11.52	42.40	29.35 Combination (T3-C)					
T4: Soil with plantain+ SiO2	7.42	62.90	20.50 SiO2 (T4- T3)					
T5: Soil with plantain+ HPβCD	8.24	58.80	16.40 HPβCD (T5- T3)					
T6: Soil with plantain+ HA	7.71	61.45	19.05 Humic acid (T6- T3)					
T7: Soil with plantain+ Tw 80	8.80	56.00	13.60 Tw 80 (T7-T3)					
4 days								
C:In autoclaved soil	15.06	24.70	0.0					
T1: In soil	13.52	32.40	7.70 microorganisms (T1-C)					
T2: Autoclaved soil with plantain	10.10	49.50	24.80 Plant (T2-C)					
T3: Soil with plantain	9.71	51.45	26.75 Combination (T3-C)					
T4: Soil with plantain+ SiO2	5.30	73.50	22.05 SiO2 (T4-T3)					
T5: Soil with plantain+ HPβCD	6.45	67.75	16.30 ΗΡβCD (T5- T3)					
T6: Soil with plantain+ HA	6.01	69.95	18.50 Humic acid (T6-T3)					
T7: Soil with plantain+ Tw 80	6.90	65.50	14.05 Tw 80 (T7-T3)					
6 days	13.07	34.65	0.0					
C:In autoclaved soil T1: In soil	13.07	43.00	8.35 microorganisms (T1-C)					
T2: Autoclaved soil with plantain	8.29	43.00 58.55	23.90 Plant (T2-C)					
T3: Soil with plantain	7.87	60.65	26.00 Combination (T3-C)					
T4: Soil with plantain+SiO2	3.47	82.65	22.00 SiO2 (T4- T3)					
T5: Soil with plantain+ HPβCD	4.20	79.00	18.35 HPβCD (T5- T3)					
T6: Soil with plantain+ HA	3.90	80.50	19.85 Humic acid (T6- T3)					
T7: Soil with plantain+ Tw 80	4.70	76.50	15.85 Tw 80 (T7-T3)					
8 days								
C:In autoclaved soil	11.50	42.50	0.0					
T1: In soil	9.05	54.75	12.25 microorganisms (T1-C)					
T2: Autoclaved soil with plantain	6.00	70.00	27.50 Plant (T2-C)					
T3: Soil with plantain	5.66	71.70	29.20 Combination (T3-C)					
T4: Soil with plantain+SiO2	1.50	92.50	20.80 SiO2 (T4-T3)					
T5: Soil with plantain+ HPβCD	1.72	91.40	19.70 HPβCD (T5- T3)					
T6: Soil with plantain+ HA	1.60	92.50	20.80 Humic acid (T6- T3)					
T7: Soil with plantain+ Tw 80	2.30	88.50	16.80 Tw 80 (T7-T3)					
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Table 2. Contribution of agents that increase the pymetrozine availability in soil

The potential of Р. major for phytoremediation of pymetrozine in a the solution amended with SiO_2 (T₄) was greater than those amended by other solubility enhancing agents, as measured by the removal of pymetrozine from contaminated soil at all experimental periods (Table 2). The percentage removal of pymetrozine through 4 days was ~73.50% in T₄. In addition to, the removal percentages of pymetrozine in T_6 , T_5 and T_7 were 69.95%, 67.75%, and 65.50%. The contribution of T₄ to the release of pymetrozine from soil was 22.05% while the contributions of T_6 , T_5 and T_7 were 18.50%, 16.30%, and 14.05% (Table 2). Results in Table 2 indicat that, the maximum contribution of T_1 , T_2 , T_3 , T_5 , T₆, and, T₇ (12.25%, 27.50%, 29.20%, 19.70%, 20.80%, and 16.80%) to the release of pymetrozine from soil during 8 days, while, T₄ during 4 days cleared (22.05%) (Table 2).

The above results showed that, most of the pymetrozine disappearance in $T_2 - T_8$, may be attributed to bioavailability of pymetrozine by enhancing agents. Romeh (2015b) found that the phytoremediation potential of *P. major* plus liquid silicon dioxide, SiO₂ was more potent than other solubility enhancing agents in removing cyanophos from the contaminated soil, improving removal percentage to 74.05% from 45.90% in soil with *P. major* only.

Improvement the Phytoremediation of Soil Contaminated with Pymetrozine Using Soluble-Enhancing Agents

Pymetrozine amounts (mg/kg) in the roots and leaves of P. major are shown in Table 3. Pymetrozine accumulated in the roots of P. major to reach the maximum levels after 4 days (53.41 mg/kg). Afterwards, the concentration decreased gradually throughout the test (Table 3). In the leaves, pymetrozine translocated into the P. major leaves and reached the maximum after 4 days of exposure (58.08 mg/kg), then decreased until the end of testing. In autoclaved soil with P. major, pymetrozine accumulated in the roots of P. major to reach the maximum level after 4 days (52.67 mg/kg). Afterwards, concentration decreased gradually throughout the test (Table 3). Pymetrozine translocated into the leaves of P. major and reached the maximum through 4 days of exposure (57.72 mg/kg) then declined until the end of testing. Liquid silicon dioxide (SiO₂) produced a synergistic effect on pymetrozine uptake and translocation. The phytoremediation potential of *P. major* plus SiO_2 was greater than that of other solubility-enhancing agents with respect to the removal of pymetrozine from contaminated soil within 1-12 days exposure (Table 3). Amending the soil with P. major containing SiO₂ resulted in a decrease in pymetrozine, half-life $T_{1/2}$; k_r, disappearance rate constant; AUCs, areas under the curve represent compound concentration during the period of study in the soil, and increased pymetrozine concentrations in plant leaves and roots (Table 3); it was detected in roots after one day then increased gradually and reached the maximum after 4 days (91.25 mg/kg) then decreased to the end of experiment. The pymetrozine concentration in P. major roots amended with SiO₂ increased by about 170.84% compared to those treated with pymetrozine in P. major roots alone through 4 days. The pymetrozine concentrations in P. major roots amended with SiO₂ HA, HPBCD and Tween 80 achieved 91.25 mg/kg, 87.50 mg/kg, 86.12 mg/kg, and 58.04 mg/kg within 4 days, respectively. Pymetrozine concentrations in P. major roots alone reached 53.41 mg/kg (Fig. 2). Combination of P. major with enhancing agents for the purpose of pymetrozine phytoremediation in roots and leaves followed the order: $SiO_2 > HA > HP\beta CD > Tween. SiO_2$ caused increased pymetrozine addition translocation to leaves by about 322.83% compared with P. major leaves alone through 4 days. The concentrations of pymetrozine in P. major enhanced with SiO2. HA, HPBCD and Tween 80 reached 187.50 mg/kg, 176.87 mg/kg, 105.12 mg/kg, and 94.25 mg/kg through 4 days, respectively. Pymetrozine concentrations in P. major leaves alone reached 58.08 mg/kg.

The total pymetrozine accumulation in entire plant amended with SiO₂, HA, HP β CD and Tween 80 arrived 278.75 mg/kg, 264.37 mg/kg, 191.24 mg/kg, and 152.29 mg/kg during 4 days compared with 111.49 mg/kg in *P. major* alone. Soil containing *P. major* and amended with SiO₂ caused a decrease in pymetrozine in soil and increases in plant leaves and roots, which is explained by the silicic acid Si (OH)₄ that enhance availability of compound from the soil to the roots and leaves of plant (Ma and Yamaji,

Table 3. Efficiency	of	solubility-enhancing	agents	in	phytoremediation	of	pymetrozine
contaminated soil							

Treatment	Days after application									
	1	2	. 4	6	8	12	t _{1/2} (days)	K _r (days)	AUC _s mg/kg (days)	
In soil										
mg/kg	17.50			11.40	9.05	7.62	8.58	0.08	1 42 .19	
Loss (%)	12.50	23.50	32.40	43.00	54.75	61.90				
In soil with P. major	14.00	11.53	0.71	7.07	F ()	2.25	E (E	0.10	07.45	
mg/kg	14.00 30.00	11.52 42.40	9.71 51.45	7.87 60.65	5.66 71.70	3.35	5.65	0.12	97.45	
Loss (%) In P. major roots	30.00	42.40	51.45	00.05	/1./0	83.23				
mg/kg	38.74	45.56	53.41	42.91	37.69	22.58				
In <i>P. major</i> leaves	2011				21105					
mg/kg	13.58	44.95	58.08	37.72	28.13	16.06				
Total uptake	52.32	90.51	111.49	80.63	65.82	38.64				
In soil with <i>P. major</i> + HPβCD						_	_		_	
mg/kg	11.20	8.24	6.45	4.20	1.72	0.16	3.10	0.22	57.84	
Loss (%)	44.00	58.80	67.75	79.00	91.40	99.20				
In P. major roots	56.22	72.63	86.12	70.19	60.05	35.09				
mg/kg In <i>P. major</i> leaves	30.22	12.05	80.1 <i>2</i>	/0.19	00.05	33.09				
mg/kg	43.15	80.12	105.12	65 43	47 66	26.85				
Total uptake	99.37	152.75								
In soil with P. major + Tween 80						••••				
mg/kg	11.60	8.80	6.90	4.70	2.30	0.60	3.41	0.20	63.28	
Loss (%)	42.00	56.00	65.50	76.50	88.50	97.00				
In P. major roots										
mg/kg	46.26	53.80	58.04	47.61	40.42	30.92				
In P. major leaves			04.95		26.00	ao 1a				
mg/kg	37.09 83.35		94.25 152.29			20.13 51.05				
Total uptake In soil with <i>p. major</i> + SiO ₂	83.33	110.54	152.29	105.25	//.24	51.05				
mg/kg	11.10	7.42	5.30	3.47	1.50	0.01	2.66	0.26	51.05	
Loss (%)	44.50	62.90	73.50		92.50		2.00	0.20	51.05	
In P. major roots										
mg/kg	45.41	65.62	91.25	56.25	35.00	0.00				
In <i>P. major</i> leaves										
mg/kg		156.21				6.25				
Total uptake	170.66	221.83	278.75	190.00	125.37	6.25				
In soil with <i>P. major</i> + HA	11.17		6.01	2 00	1.60	0.05	2.02			
mg/kg	11.16 44.20	7.71 61.45	6.01 69.95	3.90 80.50	1.60 92.00	0.05 99.75	2.92	0.23	54.65	
Loss (%)	44.20	01.43	09.93	80.30	92.00	99.73				
In <i>P. major</i> roots mg/kg	55.61	68 75	87.50	22.50	15 00	0.00				
In P. major leaves	00101	00110	01.00	22.00	10100	0.00				
mg/kg	36.25	126.25	176.87	131.25	56.25	12.50				
Total uptake	91.86		264.37							
In autoclaved soil with P. major										
mg/kg	14.30		10.10		6.00	3.90	6.01	0.11	102.44	
Loss (%)	28.50	40.00	49.50	58.55	70.00	80.50				
In <i>P. major</i> roots	27 21	11 06	57 67	41 70	26 46	31.00				
mg/kg In <i>P. major</i> leaves	37.31	44.80	52.67	41./0	30.40	21.99				
In P. major leaves mg/kg	17.34	43 46	57.72	36 88	27 26	15 68		·		
ng/kg Total uptake	54.65		110.39							
In autoclaved soil	5 1.05	00.02		, 0,00	00.14	2.107				
mg/kg	18.55	17.39	15.06	13.07	11.50	9.20	10.48	0.06	163.52	
Loss (%)	7.25		24.70							

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Fig. 2. Efficiency of solubility enhancing agents in phytoremediation of pymetrozine contaminated soil through 1-12 days of exposure; HPβCD: 3- hydroxyl, B-cyclodextrin, Tw 80: tween 80, SiO₂: silicon dioxide, HA: humic acid; (A): 1 day, (B): 2 days, (C): 4 days, (D): 6 days, (E): 8 days, (F): 12 days

2006). In addition to, humic acid (HA) caused a synergistic effect on pymetrozine uptake and translocation. Humic acid (HA) could act as a surfactant enhancing natural for the bioavailability of pymetrozine -contaminated soil. The combined effects of plants plus HA resulted in increasing the removal amount of pymetrozine from the soil, enhancing percentage degradation of 69.95% from 51.45% in soil with P. major only through 4 days of treatment (Table 3). The enhanced degradation performance for pymetrozine observed might be due to an increase in microbial activities and bioavailable in soils caused by the combined effects of plants and HA. Also, results in Table 3 show that HA was a little bit better than Tween 80 in decreasing pymetrozine - polluted soil. The surfactant activity of HA was found to increase solubility of organic contaminants on soils, hence enabling desorption-remediation of polycyclic aromatic hydrocarbon (PAH) (Holman et al., 2002), In addition to, the fraction of humic substances remaining in the soil cause a favorable role in the growth of plant and microbes and thus useful in the full recuperation of treated soils (Nardi et al., 2002). Several reports discussed the use of surfactantenhanced phytoremediation (Mitton et al., 2012). This work also showed that enhancing agents such as the surfactants, HPBCD, natural HA and Tween 80 removes amounts convergent of pymetrozine from a contaminated soil (Fig. 2). HPBCD helped in recovering 86.12 mg/kg pymetrozine and 105.12 mg/kg from contaminated soil by P. major roots and leaves within 4 days of treatment (Fig. 2). This increase in the removal of pymetrozine in soils amended with HPBCD may be due to the formation of an inclusion complex with pymetrozine (Villaverde et al., 2006). The use of plants plus surfactants has been proposed for improving phytoremediation strategies. These methods are based on the ability of agents to enhance the water solubility of hydrophobic organic compounds (HOCs) and to promote desorption, bio-degradation and phytoremediation processes (Wang and Keller, 2009).

Conclusion

The results showed that *P. major* removes efficiently of pymetrozine residues in soil and

has a potential activity for pesticides phytoremediation. Also, this study indicates that SiO_2 can enhance the phytoremediation effectiveness for pymetrozine.

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المواد المحسنة للعلاج النباتي للتربة الملوثة بالمبيد الحشري بيميتروزين

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تم تصميم هذا البحث لدراسة أهمية استخدام نبات لسان الحمل (.Plantago major L) في المعالجة النباتية للتربة الملوثة بالمبيد الحشري بيميتروزين، وتقييم تأثير إضافة (Plantago major L) - β - cyclodextrin (HPβCD) و Tween 80 - β - cyclodextrin (HPβCD) و Tween 80 - silicon dioxide (SiO₂) (SiO₂) - ۳۰,۰۰۰ و Tween 80 - silicon dioxide (SiO₂) و Iquid humic acid (HA) التحسين كفاءة التقاط المبيد من التربة الملوثة (SiO₂) باستخدام نبات لسان الحمل، وأوضحت النتائج أن تركيز المبيد قد انخفض في التربة بواسطة النبات إلى ٣٠,٠٠ - ٣٠,٠٠ من التربة الملوثة (Ar,٠٠ – ٣٠,٠٠ ليستخدام نبات لسان الحمل، وأوضحت النتائج أن تركيز المبيد قد انخفض في التربة بواسطة النبات إلى ٣٠,٠٠ - ٣٠,٠٠ مجاركة للعترة التجربة ١ – ٢٢ يوماً مقارنة بانخفاض قدره ١٢,٥٠ – ٢٦,٥٠ في تربة الكنترول (بدون النبات) ، كما أمتص المبيد بجذور وأوراق نبات لسان الحمل ليصل إلى ٣٠,٤٠ مجاركجم م ٢٠,٠٠ و مجاركة بانخفاض قدره ١٢,٥٠ مجاركجم م ٢٠,٠٠ مجاركجم بعد ٤ أيام، أدت إضافة أكسيد المبيد المبيد والتربة المرواق نبات الى ١٢,٠٠ مجاركجم م ٢٠,٠٠ مجاركجم بعد ٤ أيام، أدت إضافة أكسيد المبيد النبات) معارفية بانخفاض قدره ١٢,٠٠ مجاركجم م مجاركجم بعد ٤ أيام، أدت إضافة أكسيد السيليكون إلى زيادة التقاط الجذور للمبيد وانتقاله الى الأوراق بحوالى ١٢٠ معارك و ٣٢٢,٠٠ على المان الحمان إلى معارزة بالغاني الموراق بحوالى ١٢٠ محارك و ٣٢٢,٠٠ على المنافة أكسيد السيليكون إلى زيادة التقاط الجذور للمبيد وانتقاله الى الأوراق بحوالى ١٢٠ معارك و ٣٢٢,٠٠ على التوالى مقارنة بتجربة استخدام النبات فقط، وأمكن ترتيب تأثير إضافة المواد المحسنة مع النبات تنازليا كما يلى: التوالى مقارنة بتجربة استخدام النبات فقط، وأمكن ترتيب تأثير إضافة المواد المحسنة مع النبات تنازليا كما يلى: التوالى مقارئة بالمحسنة مع النبات فقط، وأمكن ترتيب تأثير إضافة المواد المحسنة مع النبات تنازليا كما يلى: التوالى مقارنة بتجربة الميادة معادة مالاراسة أنه يمكن إضافة معالم ترارك معلى التوربوب تأثير إضافة مالمواد المحسنة مع النبات تنازليا كما يلى: التوالى محال حمالي معاني مالي معانيمان معاني معاني معاني معاني معاني

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