Interaction between sulfur and lead in toxicity, iron plaque formation and lead accumulation in rice plant

Junxing Yang^a, Zhiyan Liu^b, Xiaoming Wan^a, Guodi Zheng^a, Jun Yang^a, Hanzhi Zhang^a, Lin Guo^c, Xuedong Wang^d, Xiaoyong Zhou^a, Qingjun Guo^{a,*}, Ruixiang Xu^a, Guangdong Zhou^a, Marc Peters^a, Guangxu Zhu^a, Rongfei Wei^a, Liyan Tian^a, Xiaokun Han^a

^a Center for Environmental Remediation, Institute of Geographic Sciences and Natural Resources Research, Chinese Academy of Sciences, Beijing 100101, PR China

^b School of Life Sciences, South China Normal University, Guangzhou 510631, PR China

^c Department of Biological and Environmental Sciences, Texas A&M, Commerce, TX 75428, United States

^d College of Resource Environment and Tourism, Capital Normal University, Beijing 100048, PR China

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ABSTRACT

Human activities have resulted in lead and sulfur accumulation in paddy soils in parts of southern China. A combined soil–sand pot experiment was conducted to investigate the influence of S supply on iron plaque formation and Pb accumulation in rice (*Oryza sativa* L.) under two Pb levels (0 and 600 mg kg^{−1}), combined with four S concentrations (0, 30, 60, and 120 mg kg^{−1}). Results showed that S supply significantly decreased Pb accumulation in straw and grains of rice. This result may be attributed to the enhancement of Fe plaque formation, decrease of Pb availability in soil, and increase of reduced glutathione (GSH) in rice leaves. Moderate S supply (30 mg kg^{−1}) significantly increased Fe plaque formation on the root surface and in the rhizosphere, whereas excessive S supply (60 and 120 mg kg^{−1}) significantly decreased the amounts of iron plaque on the root surface. Sulfur supply significantly enhanced the GSH contents in leaves of rice plants under Pb treatment. With excessive S application, the rice root acted as a more effective barrier to Pb accumulation compared with iron plaque. Excessive S supply may result in a higher monosulfide toxicity and decreased iron plaque formation on the root surface during flooded conditions. However, excessive S supply could effectively decrease Pb availability in soils and reduce Pb accumulation in rice plants.

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1. Introduction

Lead levels in paddy soils of China have been significantly enhanced recently by anthropogenic activities, such as mining, atmospheric deposition or irrigation with contaminated ground-water (Ye et al., 2004; Luo et al., 2011). Rice grown on Pb-contaminated paddy soil can accumulate high levels of Pb in shoots and grains (Zhuang et al., 2009; Liu et al., 2013). The Pb accumulated in rice plants can enter the food chain, leading to serious health risks to the human body, particularly the central nervous system of children (Wasserman et al., 1997). Thus, preventing Pb uptake and translocation in rice grown in Pb-contaminated soils is important (McLaughlin et al., 1999).

The increasing sulfur accumulation in paddy soils in southern China was confirmed by agricultural activities, such as anthropogenic SO₂ emission, super-phosphate fertilization, and wastewater irrigation

(Hu and Xu, 2002; Wang et al., 2004). Sulfur plays an important role in regulating plant growth and development (Anjum et al., 2008). However, more S accumulation in the paddy soils may disturb the uptake of other elements (e.g., As and Cd) by rice, as S can be easily affected by the change of redox potential (Eh) of paddy soils because of the transformation of the redox state of inorganic species from −2 to +6 (Hu and Xu, 2002; Hu et al., 2007; Fan et al., 2010).

The oxidizing capacity, which is characterized as radial oxygen loss (ROL), of paddy rice roots can lead to the formation of iron oxyhydroxide plaque on the root surface (Mei et al., 2012; Cheng et al., 2014; Yang et al., 2016). The effect of Fe plaque on the uptake of metals (e.g., Zn) depends on the amount of Fe plaque on the root surfaces (Otte et al., 1989; Zhang et al., 1998; Yang et al., 2014). The enhancement of Fe plaque formation was demonstrated to reduce the accumulation of As, Cd, and Pb in grains of rice (Hu et al., 2007; Cheng et al., 2014). Fe plaque can act as a barrier or buffer to the uptake of heavy metals probably because of the adsorption and immobilization of metals on plaque (Taylor and Crowder 1983; Liu et al., 2004; Mei et al., 2012; Yang et al., 2016).

* Corresponding author.

E-mail address: guojq@igsnrr.ac.cn (Q. Guo).

Sulfur may alleviate the influence of heavy metal on plant growth by S metabolism. Reduced glutathione (GSH), which is a low molecular weight tri-peptide, detoxifies heavy metals through the formation of phytochelatin (PCs), which can synthesize metal-PC complex and transport the complex to the vacuole (Sun et al., 2005; Khan et al., 2008; Gupta et al., 2013). In addition, S metabolism is closely linked to the biosynthesis of PCs in plants and metal sequestration (Thangavel et al., 2007). Recent experimental studies showed that S application could protect plants from heavy metal (e.g., As and Cd) toxicity because of the increase of GSH contents in the leaves of the plants (Hu et al., 2007; Fan et al., 2010, 2013; Zhang et al., 2013). It is necessary to clarify the interaction between S and Pb in toxicity, iron plaque formation and Pb accumulation in rice plant. In order to evaluate variation of S supply and Fe plaque on Pb accumulation and the distribution in rice plants, the major aim of the present study was to investigate the variations and correlations in GSH contents, degrees of Fe plaque formation, and the uptake and distribution of Pb in shoot and root tissues and Fe plaque on root surfaces and in the rhizospheres under moderate and excessive S conditions.

2. Materials and methods

2.1. Soil used and experimental design

The soil used in the pot trial was collected from a hydric paddy field (0–20 cm) located at Soil Fertility and Fertilizer Efficiency Long Term Monitoring Base of Qiyang City, Hunan Province, China. The soil was thoroughly mixed, air-dried, and ground to < 2 mm. The physical and chemical properties of the soil were analyzed and presented as follows: pH: 5.31; organic matter: 22.6 g kg⁻¹; total N: 1.65 g kg⁻¹; total S: 191 mg kg⁻¹; available S: 13.45 mg kg⁻¹; total Pb: 41.99 mg kg⁻¹; available Pb: 7.32 mg kg⁻¹; available Fe: 128.7 mg kg⁻¹; and available Mn: 25.4 mg kg⁻¹.

A rhizobag system with a soil-sand combination was used to collect rhizosphere and non-rhizosphere soils separately and study the effect of S application on iron plaque formation, rice leaf GSH, and Pb uptake in rice plants. Six treatments were employed with two levels of Pb (as PbCl₂) [without Pb (Pb0), 600 mg Pb kg⁻¹ (Pb600)], combined with four levels of S (as Na₂SO₄) [0 (S0), 30 (S30), 60 (S60), 120 (S120) mg S kg⁻¹]. In total, 1.5 kg of dried soil was placed in each pot. The soil used was firstly spiked with Pb (0 and 600 mg kg⁻¹ supplied as PbCl₂), then mixed thoroughly and allowed to equilibrate for 2 months. After 2 months, the bulk soil was air-dried and passed through a 2-mm sieve, and then the air-dried Pb polluted-soil was spiked with S (0, 30, 60 and 120 mg kg⁻¹ supplied as Na₂SO₄) and equilibrated for 1 month. After 1 month, the bulk soil was also air-dried, passed through a 2-mm sieve and then received (per kg) a basal application of 100 mg P as KH₂PO₄; 125 mg K as KH₂PO₄; and 110 mg N as urea. During the equilibration, the soil water was maintained at 70% of maximum water holding capacity by weight. Rhizobags, which were made of nylon netting with a mesh size of 40 µm, were 4 cm in diameter and 10 cm height and filled with 0.3 kg of quartz sand. The quartz sand was collected from Jinwuxing market, Beijing, PR China, and was not contaminated with heavy metals. Before use, the quartz sand was deionized water-washed, air-dried, and sieved (< 2 mm). The sand-filled rhizobags were placed in the center of each soil pot (15 cm diameter × 17 cm height). The sand-filled rhizobags were placed in the center of each soil pot. This rhizobag design could successfully prevent the roots and root hairs from entering the adjacent non-rhizosphere soil zone while allowing the transfer of microfauna and root exudates between the two compartments. This outcome meant that although the soil was used to enclose the outside portion of the rhizobag, the rhizosphere was confined to the sand compartment and effectively separated from the non-rhizosphere soil compartment (Hu et al., 2007; Yang et al., 2014, 2016). The plants were

transplanted into sand at the beginning of the pot experiment, and the sand is here referred to as “rhizosphere” material and at the end of the study and the soil outside of the rhizobag is referred to as “non-rhizosphere” (the comparable soil zone). The rest of the pot outside the rhizobag was filled with 1.5 kg of air-dried soil. Each treatment was repeated four times. The soils were balanced at 100% of the water holding capacity for two weeks before planting.

The rice seeds of cultivar no. 12 Zhe-You were sterilized in 10% H₂O₂ (v/v) solution for 30 min and thoroughly washed with deionized water. The seeds were germinated in the moist filter paper at a temperature of 26 °C. In each pot, two uniform germinated seeds were transplanted into each rhizobag and grown for 120 days. Soil moisture content was increased to 100% of the water holding capacity before seedling emergence and kept submerged in deionized water for the whole growth period. All the pots were arranged randomly in the greenhouse with a relative humidity of 85% and light/dark cycle of 14 h day/10 h night.

2.2. Harvest and sampling

Before harvesting, the plant height was measured, and two pieces of fresh penultimate leaves of each plant were sampled, wrapped round with aluminum foils, and stored in liquid nitrogen for GSH analysis. The rice plants were harvested by carefully removing the rhizobags from the pots. The roots were separated from the quartz sand. The plants were divided into three parts as roots, straw, and grains. All the plants were rinsed with deionized water. The straw and grains were dried to a constant weight at 60 °C for 48 h, and the dry weights were recorded. The fresh roots were used for the extraction of iron plaque. The quartz sand inside the rhizobags was referred to as the rhizosphere material. The soils in the pots were homogenized thoroughly and further referred to as the non-rhizosphere soils (Hu et al., 2007).

2.3. Extraction of iron plaque

During harvest, one of fresh root in the rhizobag was used for Pb, Fe, and Mn on the root surface by a dithionitecitrate-bicarbonate (DCB) method (Taylor and Crowder, 1983), and the other fresh root was used for the S on the root surface by 1 M HCl solution (Hu et al., 2007). The quartz sand that was collected from the rhizosphere was extracted by 0.1 M HCl solution for Pb, Fe, Mn, and S analysis (Hu et al., 2007). The rice roots or quartz sand were immersed in 45 mL DCB or HCl solution, respectively, and shaken at 280 rpm for 3 h at 25 °C. The solution was filtered with quantitative filter papers to 100 mL volumetric flasks, rinsed four times, and diluted to the volume with deionized water. After the extraction with DCB or HCl solution, the roots were oven dried to constant weight at 60 °C for 48 h, and the dry weights were recorded. The dry weights of roots, straw, and grains for each plant were referred to as plant biomass.

2.4. Plant digestions

The oven-dried root, straw, and grain samples were homogenized using a Retsch grinder (Type: 2 mm, Retsch Company, Germany) and digested by HNO₃ (guaranteed reagent) (Alexander et al., 2006). The sub-samples of plant tissues (0.5 g) were digested in flasks on an electric heating plate at 60 °C and then increased to 110 °C and kept stable until the sample solution became clear. The sample volume was adjusted to 50 mL with ultrapure water. Blank and reference material (BGW-07603) (China Standard Materials Research Center, Beijing, PR China) were utilized for quality control. The Pb recovery rates were 90 ± 10%.

2.5. Chemical analysis

The Pb, Fe, and Mn concentrations in the DCB extracts and HCl

solution were determined by inductively coupled plasma optical emission spectrometry (ICP-MS, Elan 5000, Perkin Elmer, USA). The sulfur concentrations in the HCl extracts were identified by turbidimetric spectroscopy (Verma et al., 1977). Available Pb, Fe, and Mn in soils were extracted by diethylenetriamine pentaacetic acid (DTPA: 0.005 M DTPA, 0.1 M triethanolamine and 0.01 M CaCl₂ at pH 7.3) in a soil:solution ration of 1:2 (v/v) (Lindsay and Norvell, 1978) and determined by ICP-MS. To determine the total Pb contents, the soil samples were digested in 4 mL of “aqua-regia” (HNO₃/HCl=1/3, V/V) (Allen, 1989) and determined by ICP-MS. Soil organic C was analyzed by potassium dichromate (K₂CrO₇) and total N by the Kjeldahl method (Kirk, 1950). Soil pH was measured in the soil:water ratio of 1:2.5 (v/v). Soil total S was digested by Mg(NO₃)₂ (Butters and Chenery, 1959). Available S was extracted by 0.01 M Ca(H₂PO₄)₂ (Warman and Sampson, 1992) and determined by a turbidimetric method. Available P was identified according to a Olsen method (Olsen and Sommers, 1982). The GSH content in the penultimate leaves of the rice plants was determined according to a DTNB [5, 5'-dithiobis-(2-nitrobenzoic acid)] method (Griffith, 1980).

2.6. Data analysis

DCB extractable Fe of the root surface ($T_{\text{DCB-Pb}}$) was calculated by multiplying rice tissue weight and Pb concentration in iron plaque, whereas the total Pb in the rice tissue (root, straw, and grain) ($T_{\text{Rice tissue-Pb}}$) was obtained by multiplying rice tissue weight and Pb levels in each tissue. The total Pb amount (T_{Pb}) was calculated as the sum of $T_{\text{DCB-Pb}}$ and $\sum T_{\text{Rice tissue-Pb}}$. Thus, the Pb distribution in iron plaque or rice tissue was calculated by dividing $T_{\text{DCB-Pb}}$ or $T_{\text{Rice tissue-Pb}}$ by T_{Pb} .

$$T_{\text{Pb}}(\%) = \frac{T_{\text{DCB-Pb}} \text{ or } T_{\text{Rice tissue-Pb}}}{T_{\text{DCB-Pb}} + T_{\text{Rice tissue-Pb}}} \times 100$$

where $T_{\text{DCB-Pb}}$ (g) and $T_{\text{Rice tissue-Pb}}$ (g) represent the Pb amount on root surface and Pb amount of tissue (root, straw and grain), respectively.

Data on plant performances were tested for their normality and variance prior to a one-way or two-way analysis of variance (ANOVA), as no data transformation was needed. If the differences between Pb treatments for rice plant, or among different S treatments for rice plant, were significant at 5% level, the least significant difference (LSD) or *t*-test was calculated to determine where differences lay. Pearson correlation analysis was performed to establish the relationship among Cd concentrations in roots, straw, grain and DCB-extractable Fe, Mn and Cd concentrations on

root surface and in the rhizosphere. All statistical analyses were performed using the SPSS 11.0 statistical package.

3. Results

3.1. Plant height, biomass, and root color

For rice grown in 600 mg kg⁻¹ Pb, S30, S60, and S120 treatments significantly ($P < 0.05$) increased plant height and biomass (roots, straws, and grains) compared with the S0 treatment (Table 1). According to the two-way analysis of variance test analysis, the biomasses of the rice were significantly influenced by S doses ($P < 0.01$) and Pb exposures ($P < 0.01$).

The rice roots appeared reddish under S0 and S30 treatments, whereas the roots turned black with the application of S60 and S120. However, the black color was lighter on roots exposed to the Pb600 treatment than the color on roots treated with Pb0 (Fig. S1). Evidently, excessive S (S60 and S120) resulted in H₂S poison, and Pb addition could alleviate the poison (Fig. S2).

3.2. Pb uptake by rice plants

The concentration of grain Pb in soils without Pb treatment (Pb0) was below the detection limit (< 0.01 mg kg⁻¹) (Table 2). When rice was grown in soil added with Pb600, the Pb levels in straw and grain of rice treated with S were significantly ($P < 0.05$) lower than that in rice treated with S0. However, the root Pb concentrations were significantly ($P < 0.05$) elevated under the treatment of S60 and S120 (Table 2).

3.3. Glutathione contents

Under Pb600 treatment, S supply significantly ($P < 0.05$) increased GSH content, and the peak value was observed at S30 treatment (Fig. 1). Under Pb0 treatment, S supply tended to increase GSH in rice leaves, but these differences were less obvious compared with that of the Pb600 treatment (Fig. 1). A higher GSH concentration resulted in the Pb600 treatment than in Pb0 treatment when rice was supplied with 30 and 60 mg S kg⁻¹, but no difference was observed in the other two S treatments (Fig. 1).

3.4. Concentrations of Fe, Mn, Pb, and S in iron plaque on the root surface and in rhizosphere quartz sand

Applying S (S60 and S120) significantly ($P < 0.05$) decreased

Table 1
The effect of sulfur application on plant height (cm) and biomass (g plant⁻¹) of rice exposed to paddy soils with (+Pb) and without (−Pb) addition of 600 mg Pb kg⁻¹ (mean ± S.E., $n=4$).

S levels (mg/kg)	Plant height		Root weight		Straw weight		Grain weight	
	−Pb	+Pb	−Pb	+Pb	−Pb	+Pb	−Pb	+Pb
0	52.55 ± 2.33 bB	60.05 ± 1.30 aA	2.34 ± 0.09 aA	1.32 ± 0.05 cB	5.74 ± 0.29 aA	4.73 ± 0.12 bB	3.19 ± 0.42 aA	2.21 ± 0.04 cA
30	56.83 ± 1.10 aB	61.23 ± 0.78 aA	2.47 ± 0.04 aB	2.73 ± 0.07 aA	5.97 ± 0.10 aA	6.27 ± 0.05 aA	3.31 ± 0.06 aB	4.94 ± 0.08 aA
60	57.95 ± 0.75 aA	60.43 ± 2.12 aA	1.25 ± 0.03 bB	2.59 ± 0.13 aA	4.41 ± 0.19 bB	6.03 ± 0.28 aA	1.79 ± 0.44 bB	4.65 ± 0.17 aA
120	56.30 ± 1.01 abA	58.83 ± 1.19 aA	0.89 ± 0.06 cB	2.00 ± 0.21 bA	2.29 ± 0.16 cB	4.71 ± 0.13 bA	0.84 ± 0.07 cB	3.78 ± 0.09 bA
Analysis of variance								
Pb	18.308**		33.811**		40.19**		97.468**	
S levels	1.894		43.337**		69.874**		23.17**	
Pb × S levels	1.428		53.854**		33.439**		31.327**	

Different lowercase letters within plant height, root weight, straw weight, and grain weight under the same Pb treatment indicates a significant difference among the different S levels at $P < 0.05$.

Different capital letters within plant height, root weight, straw weight, and grain weight under the same S treatment indicates a significant difference between Pb and non-Pb treatments at $P < 0.05$.

The effects of Pb treatments and S levels on rice tested were analyzed by the two-way analysis of variance test.

** $P < 0.01$.

Table 2

The effect of sulfur application on Pb concentrations (mg kg^{-1} DW) in rice exposed to paddy soils with (+Pb) and without (–Pb) addition of $600 \text{ mg Pb kg}^{-1}$ (mean $\pm$ S.E., $n=4$).

S levels (mg/kg)	Root		Straw		Grain	
	–Pb	+Pb	–Pb	+Pb	–Pb	+Pb
0	6.78 ± 0.19 aB	896.3 ± 9.33 cA	3.36 ± 0.09 bB	99.23 ± 4.9 aA	BDL	8.96 ± 0.64 a
30	6.72 ± 0.03 aB	816.2 ± 14.4 cA	3.21 ± 0.07 bB	43.64 ± 2.39 cA	BDL	2.24 ± 0.05 c
60	6.66 ± 0.81 abB	1424.6 ± 51.36 bA	2.51 ± 0.08 cB	71.23 ± 3.54 bA	BDL	5.02 ± 0.19 b
120	6.31 ± 0.09 bB	1915.1 ± 65.2 aA	3.65 ± 0.05 aB	40.69 ± 3.07 cA	BDL	2.27 ± 0.09 c
Analysis of variance						
Pb						
S levels	3502.263**		1132.449**		88.114**	
Pb $\times$ S levels	145.103**		57.506**			
	145.324**		58.543**			

Different lowercase letters within root, straw and grain under the same Pb treatment indicates a significant difference among the different S levels at $P < 0.05$.

Different capital letters within root, straw and grain under the same S treatment indicates a significant difference between Pb and non-Pb treatments at $P < 0.05$.

** $P < 0.01$.

Fig. 1. The effect of S application on GSH content in the penultimate leaves of rice (mg kg^{-1} , FW) exposed to paddy soils with (■) and without (□) addition of $600 \text{ mg Pb kg}^{-1}$ [different upper- and lower-case letters indicate significant difference at $P < 0.05$ (LSD test) for S levels with and without addition of $600 \text{ mg Pb kg}^{-1}$, respectively; * indicates a significant difference between Pb and non-Pb treatments under the same S level at $P < 0.05$ (t-test)].

Table 3

The effect of sulfur application on DCB-extractable Fe (g kg^{-1}), Mn (mg kg^{-1}), Pb (mg kg^{-1}) and HCl extractable S (g kg^{-1}) concentrations in iron plaque on root surface and in the rhizosphere of rice exposed to paddy soils with (+Pb) and without (–Pb) addition of $600 \text{ mg Pb kg}^{-1}$ (mean $\pm$ S.E., $n=4$).

S levels (mg kg^{-1})	DCB-Fe		DCB-Mn		DCB-Pb		HCl-S	
	–Pb	+Pb	–Pb	+Pb	–Pb	+Pb	–Pb	+Pb
Root surface								
0	26.32 ± 1.81 bB	38.96 ± 1.27 bA	94.32 ± 3.55 bB	131.25 ± 3.63 bA	25.9 ± 2.23 aB	211.3 ± 14.87 bA	0.55 ± 0.03 dA	0.67 ± 0.04 cA
30	39.41 ± 0.88 aB	67.23 ± 3.26 aA	138.46 ± 3.08 aB	197.94 ± 4.39 aA	25.3 ± 0.41 aB	289.3 ± 4.52 aA	1.14 ± 0.02 cB	1.57 ± 0.04 bA
60	16.63 ± 0.54 cB	21.58 ± 1.18 cA	64.21 ± 2.72 cA	71.69 ± 3.39 bA	24.3 ± 1.11 aB	155.9 ± 5.32 cA	1.39 ± 0.05 bB	1.61 ± 0.07 bA
120	12.45 ± 0.82 dB	18.94 ± 0.62 cA	50.28 ± 1.00 dB	77.38 ± 2.96 bA	23.8 ± 1.19 aB	99.8 ± 5.46 dA	2.45 ± 0.07 aB	3.44 ± 0.13 aA
Analysis of variance								
Pb	141.336**		206.027**		1403.683**		93.254**	
S levels	244.579**		455.284**		86.601**		459.067**	
Pb $\times$ S levels	22.873**		22.453**		83.472**		18.2**	
Rhizosphere								
0	14.29 ± 0.77 cB	25.28 ± 1.27 bA	1.35 ± 0.15 abA	1.68 ± 0.19 bA	0.085 ± 0.006 abB	0.35 ± 0.03 aA	0.16 ± 0.02 dA	0.21 ± 0.03 dA
30	33.72 ± 1.60 bB	48.14 ± 1.76 aA	1.92 ± 0.05 aB	2.57 ± 0.03 aA	0.114 ± 0.011 aB	0.31 ± 0.02 abA	2.41 ± 0.03 cA	2.33 ± 0.03 cA
60	23.34 ± 1.19 bB	47.56 ± 6.16 aA	1.13 ± 0.14 bA	1.54 ± 0.23 bA	0.105 ± 0.016 abB	0.26 ± 0.04 abA	5.73 ± 0.12 bA	5.41 ± 0.09 bA
120	19.34 ± 0.99 aB	26.69 ± 2.39 bA	1.72 ± 0.32 abA	1.68 ± 0.32 bA	0.074 ± 0.004 bB	0.23 ± 0.02 bA	10.21 ± 0.48 aA	9.87 ± 0.14 aA
Analysis of variance								
Pb	60.216**		5.268*		144.861**		1.707	
S levels	29.881**		7.185**		3.636*		1051.166**	
Pb $\times$ S levels	3.9*		0.93		2.579		0.516	

Different lowercase letters within DCB-Fe, DCB-Mn, DCB-Pb and HCl-S under the same Pb treatment indicates a significant difference among the different S levels at $P < 0.05$. Different capital letters within DCB-Fe, DCB-Mn, DCB-Pb and HCl-S under the same S treatment indicates a significant difference between Pb and non-Pb treatments at $P < 0.05$.

* $P < 0.05$.

** $P < 0.01$.

Table 4
The effect of sulfur application on distribution (%) of Pb in iron plaque on root surface and rice plants exposed to paddy soils with (+Pb) and without (–Pb) addition of 600 mg Pb kg^{–1} (mean ± S.E., n=4).

S levels (mg kg ^{–1})	DCB-Fe on root surface		Root		Straw		Grain	
	– Pb	+ Pb	– Pb	+ Pb	– Pb	+ Pb	– Pb	+ Pb
0	62.95 ± 2.03 aA	14.27 ± 0.78 bB	16.75 ± 1.17 aB	61.96 ± 1.85 dA	20.30 ± 1.44 abA	24.01 ± 0.78 aA	BDL	1.02 ± 0.06 a
30	63.59 ± 0.51 aA	23.94 ± 0.66 aB	16.90 ± 0.24 aB	67.46 ± 0.26 cA	19.50 ± 0.39 bA	8.27 ± 0.51 bB	BDL	0.34 ± 0.01 c
60	60.95 ± 1.44 aA	8.93 ± 0.59 cB	16.76 ± 0.41 aB	81.04 ± 0.75 bA	22.30 ± 1.20 abA	9.52 ± 0.83 bB	BDL	0.51 ± 0.03 b
120	60.10 ± 1.76 aA	4.73 ± 0.33 dB	15.99 ± 0.57 aB	90.37 ± 0.70 aA	23.92 ± 1.63 aA	4.69 ± 0.64 cB	BDL	0.21 ± 0.03 d
Analysis of variance								
Pb	3465.97**		8502.859**		188.429**			
S levels	35.187**		97.968**		28.296**		87.007**	
Pb × S levels	16.521**		108.28**		45.378**			

Different lowercase letters within DCB-Fe on root surface, root, straw and grain under the same Pb treatment indicates a significant difference among the different S levels at $P < 0.05$.

Different capital letters within DCB-Fe on root surface, root, straw and grain under the same S treatment indicates a significant difference between Pb and non-Pb treatments at $P < 0.05$.

** $P < 0.01$.

Table 5
The effect of sulfur application on Ca(H₂PO₄)₂-extractable SO₄^{2–} (mg kg^{–1}), DTPA-extractable Pb (mg kg^{–1}) concentrations and pH in the non-rhizosphere of rice exposed to paddy soils with (+Pb) and without (–Pb) addition of 600 mg Pb kg^{–1} (mean ± S.E., n=4).

S levels (mg kg ^{–1})	Ca(H ₂ PO ₄) ₂ -SO ₄ ^{2–}		DTPA-Pb		pH	
	– Pb	+ Pb	– Pb	+ Pb	– Pb	+ Pb
0	14.03 ± 0.26 dA	11.4 ± 0.22 dB	7.44 ± 0.08 bB	327.78 ± 5.16 aA	5.56 ± 0.09 bA	5.67 ± 0.08 aA
30	87.24 ± 1.67 cA	61.24 ± 1.05 cB	7.89 ± 0.08 abB	308.21 ± 6.75 abA	5.61 ± 0.01 abB	5.68 ± 0.01 aA
60	115.88 ± 1.36 bA	96.25 ± 2.76 bB	8.17 ± 0.63 abB	283.99 ± 2.96 bA	5.74 ± 0.19 aA	5.76 ± 0.03 aA
120	142.70 ± 1.53 aA	129.63 ± 1.99 aB	11.89 ± 0.68 aB	206.54 ± 21.6 cA	5.73 ± 0.13 aA	5.71 ± 0.04 aA
Analysis of variance						
Pb	190.14**		2194.652**		2.167	
S levels	2257.819**		20.398**		3.195*	
Pb × S levels	20.143**		21.034**		0.535	

Different lowercase letters within Ca(H₂PO₄)₂-SO₄^{2–}, DCB-Mn, DTPA-Pb and pH under the same Pb treatment indicates a significant difference among the different S levels at $P < 0.05$.

Different capital letters within Ca(H₂PO₄)₂-SO₄^{2–}, DCB-Mn, DTPA-Pb and pH under the same S treatment indicates a significant difference between Pb and non-Pb treatments at $P < 0.05$.

* $P < 0.05$.

** $P < 0.01$.

3.5. Lead distribution in iron plaque and rice plants

In the Pb0 combined S0 treatment, Pb was predominately distributed in the following order: iron plaque > roots > straw. No significant difference was observed for Pb distribution in iron plaque and roots except in straw (Table 4). However, in Pb600-combined S0 treatment, Pb was predominately distributed in the rice roots and iron plaque, followed by the straws and grains. Under Pb600 treatment, applying S (S60 and S120) significantly ($P < 0.05$) decreased DCB-extractable Pb distribution in iron plaque, but significantly ($P < 0.05$) increased Pb levels in the roots. The addition of S significantly ($P < 0.05$) decreased the distribution of straw Pb and grain Pb under the Pb600 treatment (Table 4).

3.6. Concentrations of Ca(H₂PO₄)₂-extractable SO₄^{2–}, DTPA-extractable Pb, and pH in the non-rhizosphere soil

Irrespective of Pb addition, the concentrations of Ca(H₂PO₄)₂-extractable SO₄^{2–} were significantly higher ($P < 0.01$) in the non-rhizosphere soils with S addition than that in the non-rhizosphere soils without S addition (Table 5). In S60 and S120 treatments, Pb addition (Pb600) significantly ($P < 0.05$) decreased the concentrations of Ca(H₂PO₄)₂-extractable SO₄^{2–} in the non-rhizosphere soils.

Lead addition significantly increased ($P < 0.05$) DTPA-extractable Pb concentrations in the non-rhizosphere soils compared with that of the Pb0 treatment, irrespective of S addition (Table 5). No significant difference was observed in soil pH with respect to S treatments and Pb addition (Table 5).

3.7. Correlation analysis

Significant negative correlations were found between HCl-extractable S concentrations and Fe and Mn concentrations on the root surfaces and in the rhizosphere (Table 6). Significant negative correlations were also found between grain Pb concentrations and HCl-extractable S concentrations on the root surfaces and in the rhizosphere.

Significant positive correlations were determined among DCB-Fe, DCB-Mn, and DCB-Pb concentrations on the root surfaces and sand surfaces under both Pb treatments (Table 6).

4. Discussion

4.1. S and Pb toxicity

The results indicated that S could alleviate Pb stress on rice, whereas Pb also could mitigate excessive S-induced stress on rice. Excessive S resulted in H₂S poison, but some heavy metals (e.g. Cd, and As) addition could alleviate the poison (Hu et al., 2007; Fan et al., 2010, 2013). The differences of root surface color and biomass between Pb0 and Pb600 treatments also strongly demonstrated that Pb could mitigate excessive S-induced stress on rice (Table 1, Fig. S1).

Sulfur may alleviate the influence of Pb on rice growth in two mechanisms. First, S induced the increase of GSH for the synthesis PCs, thus increasing Pb tolerance of plants. In this study, S30 and S60 treatments caused an evident increase in GSH content in rice leaves with Pb stress (Fig. 1). Previous research also indicated that

Table 6Correlation between concentrations of Pb in root, straw, and grain and concentrations of Fe, Mn, Pb, and S on root surfaces and in the rhizospheres of rice plants tested ($n=32$).

	Fe root surface	Fe rhizosphere	Mn root surface	Mn rhizosphere	Pb root surface	Pb rhizosphere	Pb root	Pb straw	Pb grain	S root surface	S rhizosphere
Fe root surface	1	0.529**	0.972**	0.601**	0.727**	0.519**	0.043	0.285	−0.133	−0.343	−0.609**
Fe rhizosphere		1	0.457**	0.356*	0.635**	0.497**	0.424*	0.399*	−0.331	0.063	−0.08
Mn root surface			1	0.61**	0.675**	0.504**	0.029	0.26	−0.093	−0.363*	−0.635**
Mn rhizosphere				1	0.471**	0.387*	0.128	0.165	−0.362	0.041	−0.078
Pb root surface					1	0.849**	0.543**	0.763**	0.072	−0.066	−0.311
Pb rhizosphere						1	0.664**	0.854**	0.348	−0.037	−0.251
Pb root							1	0.684**	−0.399	0.529**	0.276
Pb straw								1	0.95**	−0.085	−0.236
Pb grain									1	−0.728**	−0.631**
S root surface										1	0.897**
S rhizosphere											1

** $P < 0.01$.* $P < 0.05$.

exposure to Pb stress initially resulted in an increased consumption of GSH for PC production, whereas GSH content will recover to a higher level after prolonged Pb exposure (Verma and Dubey, 2003; Islam et al., 2008; Gupta et al., 2013). Phytochelatin and Pb could form Pb-PCs chelate complex and be taken up into the vacuole (Sun et al., 2005; Gupta et al., 2013). This condition further decreased Pb transportation from roots to shoots in plant and alleviated Pb stress (Verma and Dubey, 2003). However, the GSH decreased in rice leaves treated with Pb600 and S120 (Fig. 1). This outcome indicated that S-rich compounds induced by Pb tolerance could be related to S levels. Second, the application of excessive S decreased Pb uptake into rice by decreasing Pb availability in soil because of the formation of PbS precipitates (Table 6).

4.2. Fe plaque formation

In this study, various S doses exhibited different effects on Fe plaque formation under Pb0 and Pb600 exposures. The present results strongly suggested that moderate S supply caused an evident increase in Fe formation on root surface and in the rhizosphere, whereas excessive S supply could decrease the formation of iron–manganese oxides plaque on the root surface (Table 3). Excessive S that induced the decrease of plaque formation is probably attributed to an increase in the formation of abundant S^{2-} in bulk soil (non-rhizosphere) and the root surface. In this study, the 0.01 M $Ca(H_2PO_4)_2$ -extractable SO_4^{2-} concentrations in S-supplied non-rhizosphere soil ranged from 61 mg kg^{−1} (S30 combined with Pb600 treatment) to 142 mg kg^{−1} (S120 with Pb0 treatment), which are greater than the S deficiency critical value (12 mg kg^{−1}) (Hu and Xu, 2002). Under flooded conditions, large amounts of Fe^{2+} , Mn^{2+} , and S^{2-} can be produced by SO_4^{2-} reducing bacteria or the metabolic activities of microorganisms (Murase and Kimura, 1997; Canfield et al., 2005). Thus, the S^{2-} in bulk soil (non-rhizosphere) and on the root surfaces could result in FeS and MnS precipitation, thus ultimately inducing the decrease in iron plaque formation and/or lower stability of iron plaque. Previous studies also suggested that S application can increase or decrease the formation of iron plaque of rice (Hu et al., 2007; Fan et al., 2010, 2013). The outcome depends on the concentrations and species of S fertilization applied, capacities of root oxidation, rice growth stage, available S concentrations in tested soil, or species of metal exposure (Hu et al., 2007; Mei et al., 2012; Fan et al., 2010, 2013). Our results, together with previous reports, suggest that the concentrations of S applied may play an important role in iron plaque formation on the root surface of rice.

4.3. Pb uptake and distribution

In this study, the concentrations and distribution of Pb in straw

and grains of rice decreased with the increasing rates of S application (Tables 2 and 4). This result strongly suggested that S supply could significantly decrease Pb uptake by rice. Our studies found that different rates of S supply undergo different mechanisms to decrease Pb uptake and accumulation in rice.

Under the moderate S (S30) treatment, Fe plaque formation on the root surface and in the rhizosphere was significantly enhanced. Iron plaque was reported to be likely a barrier to the adsorption, uptake, and accumulation of heavy metals (e.g., As, Cd, and Zn) on rice surfaces (Liu et al., 2011; Mei et al., 2012). Significant positive correlations were also found between DCB-Fe, DCB-Mn, and DCB-Pb concentrations on root surfaces and sand surfaces under both Pb treatments (Table 6). This result suggested that Fe plaque induced by moderate S supply could adsorb more Pb on the root surface and in the rhizosphere. However, the effects of Fe plaque on metal uptake and translocation by wetland plants (rice) are still unclear (Ye et al., 1998; Liu et al., 2004; Cheng et al., 2014). Fe plaque formation was reported to reduce Pb, Cd, and As uptake and translocation in rice with higher rates of ROL (Mei et al., 2009; Cheng et al., 2014). On the contrary, other reports showed that metal uptake by plants was facilitated or unaffected by Fe plaque formation (Ye et al., 1998). The functions of Fe plaque on metal uptake may depend on the amount of Fe/Mn deposition on the root surfaces and metal bioavailability in the rhizosphere.

The excessive S supply (S60 and S120) significantly decreased the iron plaque on the root surface. Under Pb600 treatment, Pb was predominately distributed in rice roots, irrespective of S supply, followed by iron plaque, straws, and grain (Table 4). These results indicated that the rice root was more effective in preventing Pb from entering the plants than iron plaque under excessive S treatments. Similarly, most of the absorbed Pb was reported to accumulate in the outer cortex of roots of wetland plants (Ye et al., 1998; Vesik et al., 1999; Yang et al., 2010). By using energy dispersive X-ray microanalysis, Vesik et al. (1999) found that Pb was not localized at the root surface and increased toward the root center of water hyacinth (*Eichhornia crassipes*). Liu et al. (2011) indicated that iron plaque on rice roots can reduce Pb uptake only in relatively mild or moderate Pb-contaminated soil. The capacity of iron plaque to immobilize Pb may also be related to other factors, such as the extent of plaque (Otte et al., 1989; Zhang et al., 1998), concentrations of metal and pH in the culture solution or rhizosphere, and capacity of oxidation and the age of roots (Ye et al., 1997; Batty et al., 2000; Liu et al., 2011; Mei et al., 2012).

Compared with that of the Pb0 treatment, the addition of Pb increased both DCB-extracted Fe, Mn, and Pb concentrations in iron plaque and HCl-extractable Fe, Mn, and Pb concentrations in the rhizosphere (Table 3), indicating that Pb addition stimulated the formation of iron plaque. Ksp of PbS (3.4×10^{-28}) is lower than Ksp of FeS (3.7×10^{-19}) and MnS (1.4×10^{-18}). Thus, S^{2-} can mainly react with

Pb to form PbS rather than reacting with Fe and Mn (Kashem and Singh, 2001; Beeston et al., 2010). Thus, Pb supply relatively induced the formation of iron plaque on the root surface of rice.

In summary, S supply affected Pb uptake and distribution in rice tissues in different ways: 1) moderate S supply increased iron plaque on the root surface and in the rhizosphere; 2) excessive S supply decreased iron plaque on the root surface; 3) S supply decreased Pb availability in soils; 4) S supply induced GSH formation in the leaves. Further studies are needed to investigate the dynamics of iron plaque formation on rice roots of different growth stages under various S levels.

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Appendix A. Supplementary Information

Supplementary data associated with this article can be found in the online version at <http://dx.doi.org/10.1016/j.ecoenv.2016.02.021>.

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