Identification and quantification of phytochelatins in roots of rice to long-term exposure: evidence of individual role on arsenic accumulation and translocation

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Abstract

Rice has the predilection to take up arsenic in the form of methylated arsenic (o-As) and inorganic arsenic species (i-As). Plants defend themselves using i-As efflux systems and the production of phytochelatins (PCs) to complex i-As. Our study focused on the identification and quantification of phytochelatins by HPLC-ICP-MS/ESI-MS, relating them to the several variables linked to As exposure. GSH, 11 PCs, and As–PC complexes from the roots of six rice cultivars (Italica Carolina, Dom Sofid, 9524, Kitrana 508, YRL-1, and Lemont) exposed to low and high levels of i-As were compared with total, i-As, and o-As in roots, shoots, and grains. Only Dom Sofid, Kitrana 508, and 9524 were found to produce higher levels of PCs even when exposed to low levels of As. PCs were only correlated to i-As in the roots ($r$=0.884, $P<0.001$). However, significant negative correlations to As transfer factors (TF) roots–grains ($r$=−0.739, $P<0.05$) and shoots–grains ($r$=−0.541, $P<0.05$), suggested that these peptides help in trapping i-As but not o-As in the roots, reducing grains’ i-As. Italica Carolina reduced i-As in grains after high exposure, where some specific PCs had a special role in this reduction. In Lemont, exposure to elevated levels of i-As did not result in higher i-As levels in the grains and there were no significant increases in PCs or thiols. Finally, the high production of PCs in Kitrana 508 and Dom Sofid in response to high As treatment did not relate to a reduction of i-As in grains, suggesting that other mechanisms such as As–PC release and transport seems to be important in determining grain As in these cultivars.

Key words: Arsenic transfer factor, arsenic uptake, HPLC-ICP-MS/ESI-MS, phytochelatins, rice.

Introduction

Rice (Oryza sativa L.) is a global staple food which needs monitoring for contaminants. In a market basket survey in the United States, rice was found to have higher concentrations of inorganic arsenic (i-As) than other commodities (Schoof et al., 1999). Other studies demonstrated that this cereal accumulates more arsenic (As) in shoots and grains than wheat.
and barley (Williams et al., 2007). Rice consumption may therefore represent a significant risk, mainly when As is present in its inorganic forms, arsenite (As\textsuperscript{III}) and arsenate (As\textsuperscript{V}), which are class 1 non-threshold carcinogens (ATSDR, 2007; Meharg et al., 2009).

Arsenic uptake by rice plants depends on several factors. Cultivation under flooded conditions (Xu et al., 2008), irrigation with As contaminated water, as well as soil naturally contaminated with As all increase the levels of this element in rice grains (Saha and Ali, 2007; Lu et al., 2009; Meharg et al., 2009). Organic As (o-As) species (dimethylarsinic acid (DMA) and monomethylarsonic acid (MMA)) are absorbed to a lesser extent than i-As (Raab et al. 2007b). As\textsuperscript{III} and As\textsuperscript{V} are mostly carried via silicate and phosphate transporters, respectively, probably due to molecular similarities with silica and phosphate (Meharg et al. 1994; Ma et al., 2006; Zhao et al., 2010). After entering the root cells, As\textsuperscript{V} is readily converted to As\textsuperscript{III} and the Lsi1 efflux channels can expel As\textsuperscript{III} out of the root or into the xylem (via Lsi2), leading to transport into shoots and grains (Ma et al., 2006; Zhao et al., 2010).

Another defence system based on binding i-As in roots is the formation of As–phytochelatin complexes and their sequestration into cell vacuoles (Raab et al., 2005, 2007a; Song et al., 2010). The ability to synthesize phytochelatins (PCs) depends on genetic factors and the availability of sulphate (Cobbett and Goldsborough, 2002; Zhao et al., 2010; Duan et al., 2011).

PCs are derived from glutathione (GSH). Zhang et al. (2010) and Duan et al. (2011) observed that sulphur and GSH/PC deprivation increased As translocation in rice from roots to shoots and from shoots to grains, respectively. Duan et al. (2011) reported diverse patterns of As and PC accumulation in rice plants from different cultivars. They concluded that PCs have a specific role in As translocation and As accumulation in rice grains, without identification or quantification of the involved thiols.

The aim of this study was to compare PC production and As–PC complex formation in six different rice cultivars exposed to long-term high and low As levels in order to study their influence on the As translocation in plants. The use of high performance liquid chromatography (HPLC) hyphenated simultaneously to inductively coupled plasma mass spectrometry (ICP-MS) and electrospray mass spectrometry (ESI-MS) allowed simultaneous identification and quantification of As–peptide complexes and peptides without the need for authentic standards. Furthermore, the potential of these thiol compounds to serve as biomarkers for As exposure and their relationship to As translocation in rice were evaluated.

Materials and methods

Reagents

Ultra-pure water (Elga Ltd., High Wycombe, Bucks, UK) was used throughout the experiments. All reagents used were of analytical grade. Formic acid (98%), nitric acid (67%), and hydrogen peroxide (30%) were purchased from Fluka (UK). Potassium phosphate dibasic (99%), di-sodium arsenate (Na\textsubscript{2}H\textsubscript{2}AsO\textsubscript{4}.7H\textsubscript{2}O) and N-acetylcysteine (99%) were purchased from Sigma–Aldrich. MMA and DMA were purchased from Strem Chemicals (Newburyport, USA). Elemental standard solutions (Ge, Ga, and As, 1000 mg l\textsuperscript{-1}) were obtained from High Purity Standards (Charleston, USA) and methanol from Fisher (London, UK).

Instruments

The instruments used for speciation were an Accela HPLC system coupled (split 1:4) with an ICP-MS Element 2 and an ESI-MS LTQ Orbitrap Discovery (Thermo Fisher Scientific, Germany) for the determination of PCs and As–PC complexes. For As species determinations in grains, an 1100 HPLC system coupled with an 7500c ICP-MS (Agilent Technologies Stockport, Cheshire, UK) was used. Total As analysis was performed using an ICP-MS Agilent 7500c.

Rice cultivation conditions

Six different rice cultivars (Oryza sativa L.), Italica Carolina (IC), 9524 (9), Lemont (L), Kitrana 508 (K), Dom-Sofid (DS), and YRL-1 (Y) were selected from the Rice Diversity Panel 1 (Zhao et al., 2011). The rice lines were selected based on a study of grain arsenic in over 300 cultivars grown in the field in Bangladesh and China (Norton et al., 2012) where Lemont was low in both, Dom Sofid and Italica Carolina were low in Bangladesh but high in China, 9524 and Kitrana 508 were high in Bangladesh and low in China, and YRL-1 was high in both. In addition, in Bangladesh, 9524 and YRL-1 were identified as having a high shoot to grain As transfer factor while Dom Sofid and Italica Carolina were low.

After germination, plants were individually potted (eight plants each) and grown under flooded conditions (2.0 l-pots, soil immersion under 3–4 cm of water) and greenhouse conditions. The soil used for the experiment contained an average of 7.5 mg As kg\textsuperscript{-1} at the outset (low exposure level). Half of the plants from each cultivar received, after transplanting, 10 mg As in the form of As\textsuperscript{V} per pot (high exposure level). During growth the plants were regularly fertilized using Yoshida’s nutrient solution (Yoshida et al., 1976) and grown until the grains were mature without additional exposure to As. Just before harvesting, plant height, grain length (with husk), and the internodes stem diameter (mean of smaller diameter of the first internodes of all stems from a single plant) were measured by using a caliper ruler.

Sample preparation

Shoot lengths and all plant weights were determined after harvesting. Soil from each pot was collected and the individual plant parts (grains, shoots, and roots after removing the soil) were weighed. A portion of fresh roots (~25 g) were washed with tap water for 10 min and placed in 100 mM phosphate solution (Meharg and MacNair, 1992) to remove any surface-bound As.

Fresh root materials were used for the determination of the As–PC complexes and thiol compounds present. Sample preparation followed the procedure described in Raab et al. (2004) and Bluemlein et al. (2008). Briefly, roots were ground in liquid nitrogen and extracted on ice for 1 h with 1% (v/v) formic acid. After centrifugation, the supernatant was immediately injected into the HPLC–ICP-MS/ESI-MS. This method has been shown in plants to guarantee the integrity of the As–PC complexes and to prevent de novo synthesis of As–PC complexes during extraction and analysis (Bluemlein et al., 2008, 2009). The standards of DMA and N-acetylcysteine were daily prepared for the quantification of As and sulphur, respectively.

Plants and soil were individually ground and oven-dried at 50 °C for the determination of dry weight, total As (t-As), and speciation of i-As and o-As in grains. Samples for total As were digested with nitric acid/hydrogen peroxide in an open microwave system (MARS5, CEM). Three reference materials (NIST SRM 1568a, IAEA 140 TM, and NCZ2C73007, rice grains, algae, and soil, respectively) were used for quality control purposes. For i-As and
o-As determination, grains were extracted following the procedure of Sun et al. (2009). Briefly, ground grains were extracted with 1% (v/v) nitric acid using the microwave system above. The supernatant was mixed 1:5 v/v with hydrogen peroxide and stored overnight for complete oxidation of AsIII to AsV (Hansen et al., 2010). Then the samples were injected into a HPLC-ICP-MS.

**Conditions used during speciation**

**As-PCs and thiol-compounds.** These compounds were separated using a reversed-phase column (Eclipse, XDB-C18, 5 µm, 150 × 4.6 mm, Agilent) with a linear water/methanol (0.1% formic acid) gradient from 0–20% methanol (Acela HPLC system). The column was kept at 30 °C, the autosampler compartment at 4 °C and 0.1 ml sample was injected. The column effluent was split with one part going into the ICP-MS (Element 2) mixed via a T-piece with Ga as the internal standard. The ICP-MS was used in medium resolution with nickel cones and a PFA micro-nebulizer. The instrument was optimized using standard conditions. The rest of the column effluent was directed into an Orbitrap ESI-MS optimized daily and calibrated using standard conditions, and used in positive mode (capillary voltage 4.5 kV) with 30 000 resolution. MS² spectra were recorded when the signal intensity was above 50 000 counts. Then the influence of the gradient on the ICP-MS signal was calculated daily according to the procedure published by Amayo et al. (2011). Compounds were identified from the mass spectra and quantified using N-acetyl-cysteine and DMA, respectively (see Supplementary Table S3 available at JXB online). Inorganic and methylated As. The separation of i-As and o-As in grains was performed using a Hamilton PRP X-100 column (10 µm, 150 × 4.1 mm) and the column oven set to 30 °C. Mobile phase (1 ml min⁻¹ flow rate) consisted of 6.66 mM NH₄H₂PO₄ and 6.66 mM NH₄NO₃ at pH 6.2 (Williams et al., 2005). Sample volume injected was 50 µl. The outlet of the HPLC (Agilent 1100) was connected to the ICP-MS (Agilent 7500c) via a T-piece used for mixing the internal standard (Ga 10 µg l⁻¹) into the column effluent. The ICP-MS was optimized daily for maximum sensitivity and used in normal mode. DMA was used for calibration and quantification was done based on peak areas. DMA, AsV and MMA standard solutions were used for the identification of species (retention time).

**Arsenic transfer factor evaluation**

The soil/roots, soil/shoots, soil/grains, roots/shoots, roots/grains, and shoots/grains As transfer factors (TFs) were calculated following the procedure published by Raab et al. (2005), where the total As in the plant part of destination is divided by the total As in the origin sample.

**Statistical analysis**

The results were analysed by one-way ANOVA followed by Duncan’s multiple range test and critical ranges and correlations were tested by Pearson. Statistica (V 6.0) and SigmaStat (V 3.5) statistical software packages were used. The confidence interval used was set to higher than 95%.

**Results**

**Total As concentration and agronomic parameters**

The recovery of certified reference materials varied between 49% for soil and 105% for plant materials. Soil As concentrations were 7.0 and 14.9 mg kg⁻¹ for low and high exposure levels, respectively (Fig. 1). The soil As availability to plants depends on factors such as mineral composition (Fe, P), organic matter content, redox potential and others (Tu and Ma, 2003; Cao and Ma, 2004). Since these factors were similar for all plants, it was enough to confirm at which level the different groups were exposed to As. The high exposure level was achieved by spiking the soil used for low exposure levels with AsV. At low exposure levels, total arsenic (t-As) in roots varied from 6.9 mg kg⁻¹ in YRL-1 to 92.1 mg kg⁻¹ in Kitrana 508, in shoots from 6.7 mg kg⁻¹ in 9524 to 22.3 mg kg⁻¹ in Italica Carolina, and in grains from 0.4 mg kg⁻¹ in 9524 to 1.0 mg kg⁻¹ in Italica Carolina (Fig. 1). At high exposure levels, Lemont and Italica Carolina presented the lowest and highest concentrations, respectively (roots: 64.8 and 373.0 mg kg⁻¹; shoots: 13.5 and 38.2 mg kg⁻¹; grains: 0.7 and 2.1 mg kg⁻¹, respectively; Fig. 1). The comparison between low and high exposure samples per cultivar showed significant differences (P < 0.05) between t-As concentrations in roots, shoots, and grains in five cultivars. Lemont did not show a significant difference upon exposure to different As levels in soil (Fig. 1).

Generally, all plants per cultivar exposed to high As levels or not, flowered at the same time and the number of shoots were not different between low and high exposure plants (see Supplementary Table S1 available at JXB online). High As concentrations in soil significantly reduced plant height only for Italica Carolina (~11.7%) which also showed a reduced total plant mass. Lemont was the only cultivar where the shoot diameters were reduced (~21.4%) (see Supplementary Table S1 available at JXB online). High As treatment reduced grain yield by 13.9–54.7% in all six cultivars. Exposure to elevated As reduced the grain size for Italica Carolina, 9524, and Kitrana 508 (see Supplementary Table S1 available at JXB online).

Positive strong correlations were observed between all agronomic parameters (flowering, plant height, shoots’ diameters, weight of roots, shoots, and whole plant). Grain weights were strongly correlated with the weight of roots, shoots, and, consequently, the whole plant (see Supplementary Table S2 available at JXB online). Total As influenced all these parameters. Significant negative correlations were found between (i) t-As in: soil, roots or shoots versus grain weights; and (ii) t-As in shoots versus stem diameters, weight of roots, shoots, and grains (see Supplementary Table S2 available at JXB online).

**Arsenic transfer factors**

The TF is the measurement of the plant’s potential to take up a specific element/compound from soil, transferring it to roots, shoots and grains (Carbonell et al., 1998; Raab et al., 2007b). All cultivars, except Lemont, showed significant difference in TFsoil-root depending on As exposure, varying between 2.7 and 28.5 (Fig. 2A). TFroot-shoot were only significantly different for 9524 and YRL-1 (Fig. 2B). YRL-1 had the highest TFroot-shoot compared with other cultivars (Fig. 2B). As exposure did not significantly affect TFshoot-grain ratios (P > 0.05) to any cultivar (Fig. 2C). High exposure to As significantly reduced TFsoil-grain for Lemont and Kitrana 508 (Fig. 2D). Exposure to elevated As reduced TFsoil-grain for all cultivars (Fig. 2E), whereas TFsoil-shoot were independent of exposure level (Fig. 2F).
Fig. 1. Total arsenic concentration on a dry weight basis in soil and rice tissue from six cultivars exposed to low and high arsenic concentrations. Results presented as mean ± standard deviation; n=4; IC, Italica Carolina; DS, Dom Sofid; 9, 9524; K, Kitrana 508; Y, YRL-1; L, Lemont; (L), low exposure; (H), high exposure; *, statistical differences between low and high (P <0.05).

Fig. 2. Arsenic transfer factors between soil and plant parts from the cultivars studied. Results presented as mean ± standard deviation. IC, Italica Carolina; DS, Dom Sofid; 9, 9524; K, Kitrana 508; Y, YRL-1; L, Lemont; (L), low exposure; (H), high exposure; statistical differences between low and high exposure samples, *P <0.05; **P <0.01.
Arsenic species in grains

The species identified in grains were i-As as AsV, MMA, and DMA. No cationic species such as tetramethylarsonium, as found by Hansen et al. (2010), were present in grains. NIST 1568a rice flour was used as the reference material for extraction efficiency (~85%) and species quantification, which was similar to other studies reported in the literature (Batista et al., 2011).

Regarding the low exposure level, i-As concentrations ranged from 221 ng g⁻¹ for Lemont to 772 ng g⁻¹ for Italica Carolina and o-As from 49 ng g⁻¹ for YRL-1 to 238 ng g⁻¹ for Kitrana 508. Under these conditions, the main species found in grains was i-As for all cultivars (Fig. 3). By contrast, exposure to high As levels increased the amount of t-As (Fig. 1) mainly due to o-As species in grains (Fig. 3). In this case, i-As levels ranged from 237 ng g⁻¹ for Lemont to 888 ng g⁻¹ for YRL-1 and o-As from 71 ng g⁻¹ for Lemont to 725 ng g⁻¹ for Italica Carolina. Considering all the experiments, t-As, i-As, and o-As were positively correlated with each other (P < 0.05).

As–PC complex and thiol compound concentrations

Rice, similar to most grasses, produces a variety of PCs. By using MS and MS² data members of the following groups were identified: PCs, γ(GluCys)nGly n=2–4; Ser–PCs, γ(GluCys)nSer n=2–4; Des–PCs, γ(GluCys)n n=2–3; and Glu–PCs, γ(GluCys)nGlu n=2–3. Roots exposed to high As levels also contained, in addition to free-PCs, a variety of As–PC complexes in the extract. Dominant among these were As–PC₃, As–PC₄, As–Des–PC₃, EC–As–EC–PC₂, As–(Des–PC₂)₂, As–Des–PC₄, As–Ser–PC₃, As–Ser–PC₄, and As–Glu–PC₃. The mixture of PCs such as (γ-GluCys)nGlu, (γ-GluCys)nGly, (γ-GluCys)nSer, and (γ-GluCys)n is characteristic of the Poaceae family (grasses) (Klapheck et al., 1994; Cobbett and Goldsborough, 2002) (see Supplementary Table S3 available at JXB online; Fig. 4). Non-complexed As was
the dominant form of As in roots independent of exposure level and rice cultivar. In roots exposed to low levels of As, between 0% and 27% of t-As was complexed by PCs. On the other hand, at high As exposure levels, this increased from 45 to 38%, depending on cultivar. The particular action of phytochelatins on As transport and storage explains this slight increase which is discussed in the next section.

GSH and its oxidized form GSSG were the dominant thiol-containing peptides in all analysis (Table 1). GSH concentration was reduced at high As exposure in Kitrana 508, Dom Sofid, YRL-1, and Lemont. PCs were expressed by all plants. Dom Sofid produced the highest amount of PCs when compared with other cultivars.

Different As levels in soil did not have a significant influence on PCs concentration in YRL-1 (high grain As accumulator and medium TFroot–grain) and Lemont (low grain As accumulator and low TFroot–grain) (Figs 1, 2), showing that PCs are important for some cultivars while other varieties seem to have different mechanisms for As detoxification.

**Correlations between As–PCs complex, thiol compounds, arsenic species, and translocation factors of arsenic**

Total As levels in roots were positively correlated with some non-complexed PCs for all cultivars to low exposure ($n=24$ where DesGly–PC$_2$, $r=0.648$, $P<0.01$; PC$_{oxi}$, $r=0.535$, $P<0.01$; Glu–PC$_3$, $r=0.691$, $P<0.01$, and PC$_3$, $r=0.514$, $P<0.05$). Considering high exposure situations, the correlations were weaker ($n=24$ where DesGly–PC$_2$, $r=0.480$, $P<0.05$; PC$_{oxi}$, $r=0.365$, $P<0.05$; Glu–PC$_2$, $r=0.273$, $P>0.05$; and PC$_3$, $r=0.364$, $P<0.05$). By contrast, As–PC complexes and other thiol compounds did not correlate with t-As in roots.

TF$_{soil–root}$ were weakly correlated with the sum of non-complexed PCs ($r=0.359$, $P<0.05$), whereas TF$_{root–grain}$ and TF$_{shoot–grain}$ showed strong negative correlations ($r=-0.739$ and $-0.541$, respectively, $P<0.05$). Significant positive correlations were observed between o-As (MMA and DMA) and As in grains and GSSG and non-complexed PCs in roots of high exposure samples (Table 2). Concentrations of i-As in grains versus PCs in roots (non-complexed and complexed) showed a negative correlation whereas i-As levels in grains were positively correlated with GSH in roots. Grain t-As correlated negatively with As–PC$_4$ and As–Glu–PC$_3$. GSH did not correlate with PCs. GSSG correlated significantly with low molecular weight PCs but otherwise only weakly with DMA and As–DesGly–PC$_3$ (Table 2; for molecular weights see Supplementary Table S3 available at JXB online).

Correlations were observed between TF$_{root–grain}$ or TF$_{root–shoot}$ and t-As in grains and some As–PCs (Table 2). Finally, a correlation was observed between TF$_{root–grain}$ and the sum of PCs further correlations between all PCs (free PCs versus free PC and As–PCs; Table 2; Fig. 5).

**Discussion**

**Total As in rice and agronomic parameters**

According to the literature (Norton et al., 2012; Tripathi et al., 2012) different genotypes of rice cultivated under the same conditions exhibit different As uptake. Here, Italica Carolina and Lemont were the highest and the lowest grain As accumulators, respectively. The low grain t-As of Lemont
matches data reported by Norton et al. (2009a, b; 2012) where Lemont was identified as a low As cultivar in multiple field experiments. The cultivars which accumulated more As in roots were those that had a higher As uptake and, consequently, transported more As to aerial parts (Fig. 1). Finally, the t-As in roots of Lemont shows that the iron plaques (Chen et al., 2005) were effectively removed by applying the washing method, since no statistical difference was observed between low and high exposure levels (Fig. 1).

Arsenic had a strong negative influence on plant development as was clearly shown by the negative correlations with agronomic parameters and t-As (see Supplementary Tables S1 and S2 available at JXB online, confirming previous studies (Carbonell-Barrachina et al., 1995, 1998; Knauer et al., 1999; Zhang et al., 2002, Shri et al., 2009). Italica Carolina appeared to be the most sensitive to As and YRL-1 the least sensitive cultivar.

**Thiol compounds and As–PC complexes in roots**

Exposure to elevated As levels led to the increased formation of As–PC complexes compared with that at low exposure (Table 1). Bluemlein et al. (2008) showed that the same proportion of As complexed to biothiols was determined in plant extracts using the low temperature extraction, followed by the online HPLC-ICPMS/ESI-MS detection, as determined using direct speciation without sample preparation using XANES/EXAFS. The low-temperature extraction and the online separation/detection limit the de novo synthesis of As–PCs and their degradation during extraction and chromatographic separation (Bluemlein et al., 2009). However, this was only shown for the plant Thunbergia alata and not for rice plants. This needs to be considered when the occurrence of the As–PC complexes is discussed.

In this regard, all cultivars contained PCs at low As exposure, but only Dom Sofid and Kitrana 508 contained As bound to PCs under these conditions. Both cultivars contained, on average, the same amount of thiols in the form of GSH as in the form of PCs at low As exposure whereas, in other cultivars, the amount of thiols from GSH were higher than that present as PCs. Exposure to elevated As levels led to increased PCs’ production. All cultivars, except Dom Sofid and Kitrana 508, increased their total PCs concentrations by at least a factor of three (Dom Sofid and Kitrana 508 by about 2) upon exposure to elevated As levels.

Some PCs produced by rice in general increased in the cultivars during exposure to high As levels. These were, in most cases, bound up in the form of As–PC complexes and were not present as free PCs. Only the amount of free PC₂ and DesGly–PC₂ increased in some instances with elevated As. Tripathi et al. (2012) also found that GSH and GSSG concentrations varied according to As concentration and the

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**Table 1. Concentration of GSH, GSSG, and As–PCs complexes found in fresh roots from six different rice cultivars**

Results expressed as nmol PC kg⁻¹ of fresh roots ±standard deviation; *statistical differences between low (L), n=4 and high (H), n=4 exposure levels (P <0.05); ND, not detected; IC, Italica Carolina; DS, Dom Sofid; 9, 9524; K, Kitrana 508; Y, YRL-1; L, Lemont.

<table>
<thead>
<tr>
<th>Cultivar</th>
<th>[GSH]</th>
<th>[GSSG]</th>
<th>[OH–Me–PC₂]</th>
<th>[PC₂Red]</th>
<th>[DesGly–PC₂]</th>
<th>[PC₂Oxi]</th>
<th>[Iso–PC₂–Glu]</th>
</tr>
</thead>
<tbody>
<tr>
<td>IC(L)</td>
<td>109±69*</td>
<td>10±7.7</td>
<td>2±0.2*</td>
<td>3±0.7*</td>
<td>3±0.6*</td>
<td>4±1.0*</td>
<td>7±1.5*</td>
</tr>
<tr>
<td>IC(H)</td>
<td>1452±201</td>
<td>6±3.4</td>
<td>5±1.2</td>
<td>6±2.7</td>
<td>21±9.3</td>
<td>9±3.7</td>
<td>13±4.0</td>
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<tr>
<td>DS(L)</td>
<td>2914±1671</td>
<td>26±9.3</td>
<td>44±16.2</td>
<td>40±21</td>
<td>33±13*</td>
<td>27±8.4*</td>
<td>58±16.4</td>
</tr>
<tr>
<td>DS(H)</td>
<td>625±573</td>
<td>32±19.1</td>
<td>32±6.3</td>
<td>42±16</td>
<td>50±12</td>
<td>42±3.5</td>
<td>73±24.4</td>
</tr>
<tr>
<td>9(L)</td>
<td>137±52*</td>
<td>8±6.7</td>
<td>4±1.7</td>
<td>5±1.9</td>
<td>5±2.4</td>
<td>5±2.2</td>
<td>3±1.7*</td>
</tr>
<tr>
<td>9(H)</td>
<td>1254±466</td>
<td>5±2.5</td>
<td>4±2.2</td>
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<td>8±3.5</td>
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<td>K(L)</td>
<td>944±495*</td>
<td>8±2.4</td>
<td>5±1.6</td>
<td>6±2.7</td>
<td>13±2.8*</td>
<td>9±3.1</td>
<td>16±2.1*</td>
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<tr>
<td>K(H)</td>
<td>174±78</td>
<td>8±2.9</td>
<td>6±2.5</td>
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<td>23±6.1</td>
<td>13±5.1</td>
<td>24±3.4</td>
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<tr>
<td>Y(L)</td>
<td>2296±2198</td>
<td>5±2.2</td>
<td>3±0.7</td>
<td>3±1.0</td>
<td>6±3.9</td>
<td>5±2.3</td>
<td>7±4.8</td>
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<tr>
<td>Y(H)</td>
<td>733±123</td>
<td>4±9.9</td>
<td>3±0.6</td>
<td>2±0.9</td>
<td>5±1.8</td>
<td>2±0.4</td>
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<tr>
<td>L(L)</td>
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<td>4±1.5</td>
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<td>13±5.1</td>
</tr>
<tr>
<td>L(H)</td>
<td>355±192</td>
<td>5±1.2</td>
<td>6±0.9</td>
<td>5±1.1</td>
<td>7±0.6</td>
<td>6±1.5</td>
<td>9±2.1</td>
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<tr>
<th>Cultivar</th>
<th>[PC₃]</th>
<th>[As-PC₃]</th>
<th>[As-Iso-PC₃-Glu]</th>
<th>[As-DesGly-PC₃]</th>
<th>[As-PC₄]</th>
<th>[Iso-PC₃-Glu]</th>
<th>[As-OH-Me-PC₃]</th>
</tr>
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<tbody>
<tr>
<td>IC(L)</td>
<td>2±0.8*</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>2±0.5*</td>
<td>ND</td>
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<tr>
<td>IC(H)</td>
<td>5±0.6</td>
<td>0.2±0.1</td>
<td>0.2±0.03</td>
<td>0.4±0.3</td>
<td>1.4±0.7</td>
<td>12±5.0</td>
<td>0.2±0.1</td>
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<tr>
<td>DS(L)</td>
<td>27±16.7</td>
<td>3.7±2.2</td>
<td>3.7±2.3</td>
<td>3.8±1.6*</td>
<td>12.1±5.1*</td>
<td>27±11.7*</td>
<td>2.4±1.5</td>
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<tr>
<td>DS(H)</td>
<td>28±4.5</td>
<td>5.3±2.4</td>
<td>4.4±3.4</td>
<td>16.0±7.4</td>
<td>24.3±8.4</td>
<td>74±17.2</td>
<td>2.6±1.0</td>
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<tr>
<td>9(L)</td>
<td>2±0.6*</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
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<tr>
<td>9(H)</td>
<td>4±1.2</td>
<td>5.2±1.9</td>
<td>6.1±3.5</td>
<td>26.0±5.9</td>
<td>3±1.6</td>
<td>6.2±5.4</td>
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<tr>
<td>K(L)</td>
<td>8±3.8</td>
<td>3.0±1.4</td>
<td>9.9±3.5</td>
<td>18.0±6.0</td>
<td>13±6.0*</td>
<td>2.4±2.3</td>
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<tr>
<td>K(H)</td>
<td>14±12.3</td>
<td>4.9±2.4</td>
<td>9.0±5.4</td>
<td>17.4±7.3</td>
<td>23.9±11.0</td>
<td>46±26.2</td>
<td>7.2±10.4</td>
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<tr>
<td>Y(L)</td>
<td>1±0.8</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
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<tr>
<td>Y(H)</td>
<td>2±0.5</td>
<td>1.5±0.6</td>
<td>1.3±0.1</td>
<td>3.7±2.7</td>
<td>13.2±6.4</td>
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<td>1.3±0.1</td>
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<tr>
<td>L(L)</td>
<td>1±0.5</td>
<td>ND</td>
<td>ND</td>
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<tr>
<td>L(H)</td>
<td>1±0.6</td>
<td>2.8±1.4</td>
<td>ND</td>
<td>5.2±2.7</td>
<td>14.5±6.3</td>
<td>2.1±0.6</td>
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Table 2. Correlation between t-As, DMA, MMA, and i-As of grains and GSH, GSSG, and As–PCs of roots and transfer factors (TF) from high exposure (n=24)

<table>
<thead>
<tr>
<th>Analytes</th>
<th>SoP</th>
<th>t-As</th>
<th>DMA</th>
<th>MMA</th>
<th>i-As</th>
<th>GSH</th>
<th>GSSG</th>
<th>OHMe</th>
<th>PC2</th>
<th>DesGly</th>
<th>PC2</th>
<th>OXI</th>
<th>Iso-PC2</th>
<th>OMe-PC3As</th>
<th>PC3</th>
<th>PC2As</th>
<th>Iso-PC3</th>
<th>DesGly</th>
<th>PC4As</th>
<th>Iso-PC3</th>
<th>TFRS</th>
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<tbody>
<tr>
<td>t-As</td>
<td>0.174</td>
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<tr>
<td>DMA</td>
<td>0.699** 0.643*</td>
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<tr>
<td>MMA</td>
<td>0.713** 0.570** 0.888**</td>
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<tr>
<td>i-As</td>
<td>-0.057 0.726* 0.317 0.302</td>
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<tr>
<td>GSH</td>
<td>0.020 0.170 0.214 0.105 0.442*</td>
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<tr>
<td>GSSG</td>
<td>0.749** 0.041 0.417* 0.408 -0.252 0.086</td>
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<tr>
<td>OHMe PC2</td>
<td>0.731** -0.026 0.314 0.292 -0.262 0.257 0.761**</td>
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<tr>
<td>PC2 Red</td>
<td>0.809** 0.160 0.508* 0.483* -0.178 0.049 0.719* 0.712**</td>
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<tr>
<td>DesGly</td>
<td>0.856** 0.139 0.569** 0.491* -0.214 0.293 0.699** 0.718* 0.775**</td>
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<tr>
<td>PC2 OXI</td>
<td>0.800** 0.116 0.540* 0.486* -0.317 0.008 0.797* 0.757* 0.902* 0.891**</td>
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<tr>
<td>iso-PC2</td>
<td>0.812** 0.066 0.517* 0.486* -0.302 0.042 0.800** 0.691* 0.794* 0.902* 0.921**</td>
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<tr>
<td>Glu</td>
<td>0.455* -0.288 0.210 0.224 -0.161 -0.152 0.275 0.240 0.253 0.114 0.170 0.292</td>
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<tr>
<td>PC3 As</td>
<td>0.555** -0.322 0.238 0.296 -0.184 -0.083 0.405 0.294 0.345 0.220 0.246 0.378 0.928* 0.332</td>
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<tr>
<td>iso-PC3</td>
<td>0.643** -0.425* 0.130 0.386 -0.392 -0.162 0.416 0.289 0.310 0.139 0.299 0.440 0.776* 0.153 0.756**</td>
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<tr>
<td>Glu As</td>
<td>0.668** -0.185 0.283 0.456* -0.241 -0.221 0.500 0.321 0.432* 0.418* 0.419* 0.504* 0.763* 0.326 0.791* 0.741*</td>
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<tr>
<td>PC4 As</td>
<td>0.575** -0.485 0.111 0.225 -0.244 -0.210 0.292 0.155 0.284 0.168 0.229 0.321 0.853* 0.311 0.910* 0.756* 0.833*</td>
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<tr>
<td>iso-PC4</td>
<td>0.767** 0.091 0.363 0.483* -0.419 0.103 0.777* 0.853* 0.743* 0.814* 0.819* 0.782* 0.118 0.824* 0.238 0.319 0.329 0.052</td>
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<tr>
<td>Glu TFRS</td>
<td>-0.297 0.244 -0.177 -0.086 0.186 -0.326 -0.027 0.052 -0.097 -0.222 -0.018 -0.043 -0.268 -0.015 -0.277 -0.301 -0.178 -0.159 0.262</td>
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<tr>
<td>TFRG</td>
<td>0.547** 0.468 -0.053 -0.079 0.444* -0.184 -0.241 -0.221 -0.324 -0.410 -0.277 -0.296 -0.402 -0.204 -0.440 -0.445* -0.451* -0.384 -0.118 0.842*</td>
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</table>

* t-As, Total arsenic; DMA, dimethyl-arsenic; MMA, monomethyl-arsenic; i-As, inorganic arsenic; SoP, sum of PCs concentrations; TF, transfer factor; TFRS, TF roots–shoots; TFRG, TF roots–grains; *P <0.05; **P <0.01.
Phytochelatins and arsenic transfer in rice given long-term exposure

cultivar (genotype) studied, especially in susceptible ones. The authors found that susceptible rice cultivar IET-4786 showed more pronounced oxidative stress compared with tolerant Triguna. They attributed this tolerance to the activity of enzymes such as serine acetyltransferase, cysteine synthase, $\gamma$-glutamyl cysteine synthase, arsenate reductase, PC synthase, and others.

As reported by Tripathi et al. (2012) As translocation depends on the cultivar sensitivity and the synthesis of PCs, which is non-dose-dependent. The authors stated that the synthesis of PCs increase with As$^V$ concentration. However, above a certain concentration of As, there is a reduction in the synthesis of PCs. As previously observed, the present study indicated that a specific cultivar with different As TFs showed singular sensitivity to As. However, the total concentration of PCs did not give an indication of TFs.

Inorganic and methylated As in grains

Total As levels in grains were dose- and cultivar-dependent (Figs 3, 5). Despite the fact that the soil originally contained predominantly i-As and was spiked with i-As, high amounts of methylated As-species were found in grains. This was most likely the result of soil bacteria, especially As-methylating ones, thriving under flooded culture conditions. It has been shown (Raab et al., 2007b) that methylated As (especially DMA) is better translocated to aerial parts of rice than i-As, potentially due to (i) a lower ability to bind to PCs and be sequestered into root vacuoles and (ii) efficient transporters as shown by Li et al. (2009). The increased As levels in grains of most cultivars, except Lemont and YRL-1, at elevated As levels resulted from increased amounts of DMA and not of i-As.

Arsenic translocation in rice cultivars

The transfer factor (TF) is defined as the long-range transport of a specific compound from one part into another under specific condition of cultivation (Carbonell et al., 1998; Raab et al., 2007b). TFs vary considerably between plant species and within cultivars of a single species (Zhao et al., 2010). The most important TF of rice with regard to human nutrition is the TF$_{soil-grain}$ (Fig. 2D). In these experiments it varied between 0.05 and 0.22 with Italica Carolina having the highest TF$_{soil-grain}$ at low exposure. TF$_{soil-shoot}$ values (Fig. 2F) found here for rice were similar to those found by Williams
et al. (2007), being about 50 times higher than reported for wheat and barley. Comparison of the different exposure levels showed that TF$_{\text{soil-root}}$ increased with the amount of As present in soil for all cultivars, but Lemont (Fig. 2A). By contrast, TF$_{\text{roots-shoot}}$ decreased in all cultivars with increasing soil As content, except Lemont. Similarly, all other TFs (soil–grain, root–grain, and soil–shoot) either remained constant or decreased with elevated As levels in soil.

Considerable differences in TF factors were identified between cultivars. Kitrana 508 showed the highest As transfer from soil to roots, but TF into shoots and grains was lower than for YRL-1 and Lemont. The mean roots to shoots/grains and shoots to grains TFs for low exposure level for YRL-1 (Fig. 2B, C, E) were higher than the TFs for the other cultivars, showing a predilection for As transfer from roots to shoots and roots to grains. However, TF$_{\text{shoot-grain}}$ does not seem to be a good parameter to evaluate since no statistical differences were observed between low and high exposure samples (Fig. 2C).

Role of thiol compounds and As–PCs on As transfer into grains

To understand the correlations between PCs it is necessary to study the biosynthesis of those compounds. Initially, the enzyme $\gamma$-glutamylcysteine synthetase (GCS) produces $\gamma$-GluCys and glutathione synthetase (GS) completes the synthesis of GSH by adding Gly to the structure, producing $\gamma$-GluCysGly (Fig. 6). GSH, by negative feedback, controls the GCS activity (Cobbett, 2000; Cobbett and Goldsbrough, 2002). The synthesis of peptides ($\gamma$-GluCys)$_n$Gly in plants is stimulated by toxic elements (such as As and Cd) being catalysed by the constitutive enzyme PC synthase (PCS), which transfers the glutamyl-cysteine fraction ($\gamma$-GluCys) from GSH to an acceptor molecule, GSH or peptide ($\gamma$-GluCys)$_n$Gly, producing ($\gamma$-GluCys)$_{n+1}$Gly, the PCs (Fig. 6). In this step the highest producer is Dom Sofid, followed by Kitrana 508 and Italica Carolina and, interestingly, the lowest are YRL-1 and 9524, cultivars that showed increased i-As in grains to high exposure level (Fig. 3; Table 1). PCs with no Gly at the

Fig. 6. Probable pathways of phytochelatins (PCs) biosynthesis in roots of rice: data from literature, correlation between the PCs found here and the PCs related to the cultivars studied. Cultivars: L, Lemont; DS, Dom Sofid; IC, Italica Carolina; Y, YRL-1; 9, 9524; K, Kitrana 508. LMW, low molecular weight; HMW, high molecular weight; PCS, PC synthase; GCS, $\gamma$-glutamylcysteine synthetase; GS, glutathione synthetase; hPCS, homo-PC synthases; 1, As uptake by roots; 2, production of LMW PCs; 3, As–PC complexes (As$^{3+}$+2PC$_2$, As$^{3+}$+PC$_3$, for example) stored in the vacuole; 4, transport to vacuoles; 5, release of PCs; 6, efflux of As up-taken; ?, unknown information; >, high concentration; <, low concentration; thicker dotted line, PC highly effective. Scheme elaborated according to descriptions from Klapheck et al. (1994); Cobbett (2000); Chassaigne et al. (2001); Cobbett and Goldsbrough (2002); Oven et al. (2002); Meharg and Hartley-Whitacker (2002); Rey et al. (2004); Vatamaniuk et al. (2004); Song et al. (2010); and Wood and Feldmann (2012).
PCs provide an important mechanism to decrease the concentration of i-As in grains and suggests that PC bound As was less likely to be transported into shoots and grains. Similar observations were made by Liu et al. (2010) in Arabidopsis thaliana. By contrast with the negative correlation between i-As in grains and PCs in roots, the GSH concentration was positively correlated with the amount of i-As in grains. GSH participates not only in production of PCs (Cobbett and Goldsbrough, 2002) and in antioxidant defence mechanisms, but also in the conversion of As$^V$ in As$^{III}$ (Raab et al., 2004; Bleeker et al., 2006; Duan et al., 2011), which might be the reasons for the positive correlation.

No correlation was found between GSH and any of the individual PCs quantified, but individual PCs concentrations were correlated with the amount of GSSG. Strong correlations existed between individual PCs especially within a family, supporting the fact that chain elongation is taking place (Fig. 6). The correlations between the sum of PCs and individual PC concentrations were stronger for free PCs than for those bound to As. The correlations between DesGly–PC$_2$ and other PCs may indicate that, similar to microorganisms and yeast (Klapheck et al., 1994; Cobbett and Goldsbrough, 2002; Zhao et al., 2010), rice roots cells can also polymerize γ-GluCys directly with or without the addition of Glu or other amino acids. The cultivars showed individual differences not only in PC production but, potentially more importantly, in the different PC-families produced in response to As. Italicca Carolina responded to high As exposure especially with increases of PC$_{2oxi}$ and PC$_3$ besides DesGly–PC$_2$. This PC was also strongly elevated by As in cultivars YRL-1 and 9524. Both of these cultivars also increase the amount of Glu–PC$_2$ while 9524 had lower Glu–PC$_2$ than other cultivars. In these cultivars the PCs were not only positively correlated with the As concentration in roots but also with reduced i-As transfer to grains. YRL-1 did not produce many different PCs and was generally less sensitive to As compared with Italicca Carolina. Lemont, despite its low total As uptake into the roots (independent of exposure), produced PCs in response to As which appear to reduce As transfer to grains even further.

Some cultivars, especially those which produced significant amounts of PCs upon exposure to elevate i-As, might also respond to exposure to other soil contaminants such as Cd, which stimulates PC synthesis. Hence, a co-exposure of As and Cd will probably influence the TFs of i-As. This might explain the site to site variation of As accumulation in rice grains observed by Norton et al. (2009a, b, 2012). Those authors found that Lemont, as well as CT9993-5-10-1-M, Azucena, and Teqing cultivars, was the lowest in grain As concentration, with potential use on breeding seeds. In six different fields in Bangladesh, China, and India, Lemont was one of the lowest As grain accumulator. This cannot be explained by the high amount of PCs or thiols production, but must
be the result of another mechanism. The agreement between that study and this suggests Lemont is indeed a promising cultivar for breeding low grain As, and the results here suggest that its low TF$_{\text{soil-shoot}}$ is the main reason for a low grain $i$-As. Exactly why its TF$_{\text{soil-shoot}}$ is low is still not clear.

Conclusion

In general, t-As in grains reflected the concentration in roots. Cultivars with higher As uptake (all species) from soil translocate As more efficiently to the grains. Elevated As concentrations in roots, shoots, and grains negatively affected plant development and, consequently, grain yield, especially in cultivar Italica Carolina.

Cultivars with higher As-uptake into roots, such as Dom Sofid and Italica Carolina, produced higher concentrations of PCs than other cultivars. All rice cultivars produced a range of different PCs belonging to the Des–Gly–PC$_n$, PC$_n$, Ser–PC$_n$, and Glu–PC$_n$ families with $n$ varying from 2 to 4. Correlations between different PCs proved that chain elongation by one $\gamma$-Glu-Cys unit is used by rice for the production of PCs.

With the exception of cultivar Lemont, t-As in grains increased with increasing As exposure. Speciation of As in grains revealed that elevated As exposure resulted in increased concentration of DMA and also MMA in grains. PCs’ production and the formation of As–PC complexes in roots reduced the transfer of $i$-As into shoots and grains, but had no influence on DMA and MMA transfer.

Transfer factors for As from the soil/root to grains were lower when the plants were exposed to elevated As levels due to the stimulation of more extensive production of PCs and, consequently, reduced As transport to aerial parts of the plant. Results also showed that the uptake of methylated As species is different between cultivars, probably due to soil microbial methylation of As and As efflux from roots.

This study showed clearly that As-uptake and transfer into grains strongly depends on rice cultivar and bioavailability of As in soil. Some cultivars like Dom Sofid, Kitrana 508, and Italica Carolina are better able to mobilize As from soil than others, but efficiently trap the most toxic form, As$^{III}$, in the roots. Knowing the characteristics of each genotype (cultivar) and concentration (availability) of arsenic in soil, different cultivars can be used to prevent or reduce the arsenic contamination of grains and, consequently, the toxicological/nutritional risk, especially of $i$-As. Finally, this research suggests that modifications of PCs in plant roots will have implications on grain As levels and more research is need to investigate if this can be exploited in crop improvement targeting the reduction of As exposure risk.

Supplementary data

Supplementary data can be found at JXB online. Supplementary Table S1. Description of agronomic parameters. Supplementary Table S2. Correlations between agronomic parameters and t-As ($n=$48, low and high exposure levels).

Supplementary Table S3. Theoretical, experimental, and accurate masses ($\Delta$), retention time (RT), and molecular formula of GSH, free PCs, and PCs-As identified and quantified in the present study.

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References


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