Modulation of Metabolism and Switching to Biofilm Prevail over Exopolysaccharide Production in the Response of Rhizobium alamii to Cadmium

Mathieu Schue1,2,3, Agnes Fekete4, Philippe Ortet1,2,3, Catherine Brutesco1,2,3, Catherine Santaella1,2,3, Thierry Heulin1,2,3, Wafa Achour4,5, M H Kvetan6

1 CEA, Lab EcoL MicrobiennE Rhizosphere & Environmental Extre, IEB, DSV, Saint-Paul-lez-Durance, France, 2 CNRS, Unité Mixte Rech Biol Végétale & Microbiol Environ, UMR 6191, Saint-Paul-lez-Durance, France, 3 Université Aix Marseille, Saint-Paul-lez-Durance, France, 4 Helmholtz-Zentrum München, German Research Center for Environmental Health, Institute for Ecological Chemistry, Department of BioGeochemistry and Analysis, Neuherberg, Germany, 5 Department for Chemical-Technical Analysis Research Center Weihenstephan for Brewing and Food Quality, Technische Universität München, Freising-Weihenstephan, Germany

Abstract

Heavy metals such as cadmium (Cd2+) affect microbial metabolic processes. Consequently, bacteria adapt by adjusting their cellular machinery. We have investigated the dose-dependent growth effects of Cd2+ on Rhizobium alamii, an exopolysaccharide (EPS)-producing bacterium that forms a biofilm on plant roots. Adsorption isotherms show that the EPS of R. alamii binds cadmium in competition with calcium. A metabolomics approach based on ion cyclotron resonance Fourier transform mass spectrometry has showed that cadmium alters mainly the bacterial metabolism in pathways implying sugars, purine, phosphate, calcium signalling and cell respiration. We determined the influence of EPS on the bacterial response to cadmium, using a mutant of R. alamii impaired in EPS production (MSΔGT). Cadmium dose-dependent effects on the bacterial growth were not significantly different between the R. alamii wild type (wt) and MSΔGT strains. Although cadmium did not modify the quantity of EPS isolated from R. alamii, it triggered the formation of biofilm vs. planktonic cells, both by R. alamii wt and by MSΔGT. Thus, it appears that cadmium toxicity could be managed by switching to a biofilm way of life, rather than producing EPS. We conclude that modulations of the bacterial metabolism and switching to biofilms prevail in the adaptation of R. alamii to cadmium. These results are original with regard to the conventional role attributed to EPS in a biofilm matrix, and the bacterial response to cadmium.

Introduction

The exposure of bacterial cells to heavy metals in their environment mediates biological effects, usually through the direct or indirect action of reactive oxygen species [1,2]. In fact, non-redox-reactive metals, such as cadmium, show a high degree of reactivity towards sulfur, nitrogen and oxygen atoms in biomolecules. Cadmium may bind sulfur in essential enzymes, and alter their functions. Many studies have focused on the molecular mechanism of bacterial cell tolerance to cadmium, mainly for the case of species that are resistant to high metal concentrations, such as Stenotrophomonas [3] or Capnocytophaga [4]. However, cadmium concentrations and its availability in metal contaminated soils are generally low. At low cadmium concentrations, Dedieu et al. [5] studied the interactions of Sinorhizobium meliloti extracellular compounds on cadmium speciation and availability, and Pagès et al. [6] reported on the completely different adaptation mechanisms of phenotypic variants of Pseudomonas brassicacearum in the presence of cadmium. Varied mechanisms account for cadmium detoxication in bacteria, including exclusion, binding and sequestration. Cadmium is removed from cells by metal efflux transporters [7,8,9], reduced as cadmium sulfide [10], precipitated as insoluble salts [11], immobilized within the cell walls [12], or linked to chelating agents [13,14]. Cell exudates, such as proteins, siderophores and to a minor extent polysaccharides, play a role in the short-term interaction between Sinorhizobium meliloti and cadmium [5,15]. Because the adsorption of cadmium as well as of other metals can be associated with the secretion of exopolysaccharide (EPS) or capsular material [2,16,17], EPSs are considered as potential metal transporters in soil [18,19].

Gram-negative soil bacteria belonging to the commonly named rhizobia are able to produce EPSs with a large diversity of chemical structures [20,21]. These EPSs are the main contributors in legume-rhizobia interactions, leading to nodulation and nitrogen fixation. Eventually, much of what we know about rhizobia and their EPSs arises from studies of their symbiotic interactions with legume plants, whereas their interactions with...
non-legumes have been neglected. However, rhizobia associate with the roots of non-legumes such as Arabidopsis thaliana [22,23], Helianthus annuus [24] and Brassica napus [25]. Rhizobial EPSs also have functions beyond specific recognition in the nodulation process, such as plant growth promotion of non-legumes [24] or evasion from the defense response of plant legumes during crack entry in roots [26]. We should consider communities of rhizobia and their EPSs as integral and functionally important partners of a diverse plant rhizosphere.

For that purpose, we addressed the question of how Rhizobium alamii [27], an EPS-producing bacterium, responds to the toxic impact of cadmium. R. alamii is a rhizobacterium, isolated from the rhizophere of the sunflower, producing a mucoid and an adhesive EPS [24]. This bacterium colonizes the root system of the sunflower, A. thaliana, and rapeseed, and together with its production of EPS in the rhizosphere, it improves the physical structure of the root adherent soil and plant growth under conditions of hydric stress [23,24]. We studied the mechanisms through which EPS could contribute to the tolerance of R. alamii to cadmium, by using a mutant strain impaired in EPS production (MSDGT) [23].

Results and Discussion

Effect of cadmium on R. alamii growth and EPS production

In this study, the minimal inhibitory concentration (MIC) of cadmium (Cd²⁺) in R. alamii cells was assessed at 44 μM (5 mg.L⁻¹ Cd²⁺) in a tenfold diluted Tryptic Soy Broth (TSB/10) and as illustrated in Figure 1. The increase of Cd²⁺ concentration from 1 mg.L⁻¹ to 2 mg.L⁻¹ almost doubled the lag phase. We studied the mechanisms through which EPS could contribute to the tolerance of R. alamii to cadmium, by using a mutant strain impaired in EPS production (MSDGT) [23]. We investigated cadmium adsorption on the EPS, in the absence or in competition with calcium. Ion cyclotron resonance Fourier transform mass spectrometry (ICRFT/MS) was used to monitor the metabolic perturbations after exposure of R. alamii cells to cadmium, which slowed down, but did not inhibit cell growth. We discuss the means by which the bacterial cells adjusted their entire metabolic processes, and their way of life, to limit cadmium damage. We have for the first time shown that cadmium promotes the formation of a biofilm by R. alamii, and that this change occurs independently of the presence of EPS. These results are original with regard to the conventional role attributed to EPS in a biofilm matrix, and the bacterial response to a heavy metal.

Figure 1. Growth curves of Rhizobium alamii in ten-fold diluted tryptic soy broth (TSB/10) at 30°C in the presence of cadmium nitrate (0 to 10 mg.ml⁻¹ of Cd²⁺ (1 to 89 μM). Dark circle 0 mg.L⁻¹; Open circle 1 mg.L⁻¹; Grey triangle 2 mg.L⁻¹; Cross 5 mg.L⁻¹; Lozenge 10 mg.L⁻¹. Mean of 3 replicates ± SD. doi:10.1371/journal.pone.0026771.g001
of restricted metabolite possibilities for a given organism, using the KEGG pathway database: http://www.genome.jp/kegg/pathway.html [29]. By considering the set of *R. alamii* metabolites detected in positive and negative ESI of MS, in the experiments with or without cadmium, 1897 putative compounds were common to these conditions, and respectively 936 and 653 probable components were exclusively expressed, in the presence or absence of the metal. Cadmium induced metabolic alterations in major pathways, as summarized in Tables 1 and 2. Cadmium increased the number and level of enzymes and metabolites involved in the sugar metabolism (potentially fructose, glucose, mannose, galactose, cellobiose, inositol, starch and sucrose), phosphorylated intermediates of glycolysis compounds (likely glucose 6-phosphate, fructose 6-phosphate, glucose 1-phosphate), ABC transporters of sugars (potential methyl-galactoside, D-fructose, fructose, cellobiose) and the phosphotransferase system which, in bacteria, is the major carbohydrate transport system for incoming sugar substrates, through translocations across the cell membrane. Cadmium ions are admitted into sensitive bacterial cells by the energy-dependent manganese transport systems, which cause rapid cessation of respiration by binding to sulfhydryl groups in proteins [30,31]. Glycolysis therefore appeared to be an alternative pathway for energy. Under cadmium stress, hexoses were potentially channelled towards a pentose phosphate pathway. In the presence of Cd\(^{2+}\), the nucleotide metabolism revealed a modulation of the purine metabolism, with an accumulation of purine-based nucleosides (adenosine, deoxyadenosine, guanine and deoxyguanine) at the same time as a decrease in purine-based nucleotides (AMP and dGMP) as illustrated in Figure 3. This result was confirmed by ultra performance liquid chromatographic (UPLC) analysis (Figure 4). The decrease in nucleotide contents and the recovery of pyrimidine and purine compounds could be consistent with an adaptation process, through slowing of cell division in cells likely to adapt to the metal toxicity [32]. In the presence of cadmium, the citric acid cycle and tryptophan metabolism were shut off (Table 1) suggesting the cessation of aerobic sugar respiration and amino acid synthesis. A glutathione precursor, \(\gamma\)-L-Glutamyl-L-cysteine, was identified under cadmium stress. In response to cadmium, *Pseudomonas brassicacearum* is found to switch from the citric acid cycle to an anaerobic metabolism [6], and the inhibition of protein and glutathione syntheses are described in the response of *R. leguminosarum* to cadmium [14,33]. Cadmium activated the biosynthesis of lipopolysaccharide and capsular polysaccharide, which are the outer membrane components of Gam-negative bacteria and are potential metal binding sites [6,12,34]. Pyrophosphate was detected following Cd\(^{2+}\) treatment only. Pyrophosphate can form crystals with cadmium [35]. The accumulation of inorganic phosphate is a detoxification mechanism reported in *Klebsiella aerogenes* [36] and is also involved in the regulation of biofilm formation in *Pseudomonas fluorescens* [37]. Cadmium is also activated the inositol phosphate metabolism that regulates cytoplasmic Ca\(^{2+}\) and communication, and the cyclic ADP-ribose pathway, which is a

### Table 1. Modification of KEGG pathway metabolites induced by cadmium in *Rhizobium alamii*.

| KEGG pathway                                           | Number of identified metabolites | Cadmium concentration (mg.L\(^{-1}\)) |
|--------------------------------------------------------|----------------------------------|--------------------------------------|
| Glycolysis/Gluconeogenesis                              | 0                                | 2                                    |
| Citrate cycle (TCA cycle)                               | 3                                | 0                                    |
| Pentose phosphate pathway                               | 0                                | 6                                    |
| Pentose and glucuronate interconversions                | 1                                | 4                                    |
| Fructose, mannose, inositol and galactose metabolism   | 12                               | 44                                   |
| Purine, pyrimidine metabolism                          | 30                               | 26                                   |
| Lysine biosynthesis                                    | 10                               | 2                                    |
| Glutathione metabolism                                 | 1                                | 1                                    |
| Starch and sucrose metabolism                          | 6                                | 21                                   |
| Nucleotide sugars metabolism                           | 0                                | 2                                    |
| Lipopolysaccharide biosynthesis                        | 0                                | 1                                    |
| Peptidoglycan biosynthesis                             | 1                                | 2                                    |
| Inositol phosphate metabolism                          | 0                                | 7                                    |
| ABC transporters                                       | 13                               | 20                                   |
| Two-component systems                                  | 0                                | 2                                    |
| Phosphotransferase system                              | 9                                | 17                                   |

*Rhizobium alamii* cells were grown up to a late exponential phase, with 2 mg.L\(^{-1}\) of cadmium, as compared to the absence of cadmium. The potential metabolites identified were matched in KEGG pathway using MassTRIX (http://masstrix.org). doi:10.1371/journal.pone.0026771.g002
secondary messenger for Ca\(^{2+}\) mobilisation [38]. The structural homology between calcium and cadmium ions might account for the activation of pathways needed to maintain calcium homeostasis and signal transduction.

Altogether, these results show that the adaptation of \textit{R. alamii} cells to cadmium could imply a multifaceted scheme using metal binding, precipitation or export, changes in respiration process, and the retention of cadmium-like metal homeostasis.

Cadmium induces the formation of a biofilm by \textit{R. alamii} independently of EPS synthesis

\textit{R. alamii} wt and MS\textsc{AGT} strains were grown statically in the presence of cadmium (from 0 to 15 mg.L\(^{-1}\) of Cd\(^{2+}\)), in mineral RCV medium supplemented with glucose (for composition see materials and methods) to promote EPS synthesis. Figure 5A shows staining of biofilms formed on the tube walls using crystal violet [39], and Figure 5B represents the quantity of planktonic

| Activation of pathways                              | Hypothesis                                                                                     |
|-----------------------------------------------------|------------------------------------------------------------------------------------------------|
| Sugar metabolism                                    | The cessation of respiration by binding of cadmium to                                          |
| ABC transporters of sugars                          | sulfhydryl groups in proteins induces the glycolysis or                                         |
| Phosphorylated intermediates of glycolysis          | the shift towards the pentose phosphate pathway as an alternative pathway for energy           |
| Phosphotransferase system                           |                                                                                                 |
| Pentose phosphate pathway                           |                                                                                                 |
| Biosynthesis of lipopolysaccharide and capsular polysaccharide | Outer membrane components of Gram-negative bacteria are metal binding sites                  |
| Pyrophosphate                                        | Cadmium binding to pyrophosphate                                                               |
| Inositol phosphate metabolism                       | Regulation of cytoplasmic Ca\(^{2+}\) and communication                                        |
| \(\gamma\)-L-Glutamyl-L-cysteine                     | Glutathione precursor                                                                         |

| Inhibition or decrease in pathways                  | Hypothesis                                                                                     |
|-----------------------------------------------------|------------------------------------------------------------------------------------------------|
| Citric acid cycle                                   | The cessation of aerobic sugar respiration and amino acid synthesis                           |
| Tryptophan metabolism                               | acid synthesis                                                                                 |
| Nucleotides                                          | Cell division is slowed to accommodate to the toxicity                                         |

KEGG pathways activated or inhibited in the response of \textit{Rhizobium alamii} to cadmium (2 mg.L\(^{-1}\)).

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Figure 3. Analysis of purine and pyrimidine metabolism from \textit{R. alamii} cells grown with no cadmium (white bars) or 2 mg.L\(^{-1}\) of cadmium as cadmium nitrate. Masses of the purine based cyclic nucleotides were additionally detected from the Cd\(^{2+}\) treated samples. 2,3\&3,5-cyclic-AMP from the wt and 2,3\&3,5-cyclic-GMP were detectable according to the FTMS measurements and use of MassTrix for data interpretation. Generally, increase in the intensity of the purine based nucleosides and decrease of the detected nucleotides were observed in presence of Cd\(^{2+}\). doi:10.1371/journal.pone.0026771.g003
cells monitored by means of optical density measurements, and the quantification of surface-attached cells stained by crystal violet coloration as a function of cadmium concentration. Increasing cadmium concentrations significantly decreased the population of the planktonic cells, and induced the attachment of cells as a biofilm \((p<0.05)\).

A biofilm is a lifestyle able to resist various environmental stressors such as antibiotics [40], or metals such as copper, zinc and lead [41], and nickel [42]. In the present study, we reveal for the first time cadmium-induced biofilm formation by \textit{R. alamii} cells. Interestingly, wt and EPS-mutant strains showed the same change in growth, from free-swimming to biofilm mode, in response to cadmium, suggesting that biofilm formation, rather than EPS, was a remediation to metal toxicity.

**Cadmium binds to the EPS of \textit{R. alamii}**

The chemical structure of \textit{R. alamii} EPS contains carboxylic and hydroxyl functions [43] that may bind metals. The biosorption of metal ions on the EPS of \textit{R. alamii} has been measured in a MOPS buffer and in a calcium-containing medium, and has been described using the Langmuir and Freundlich isotherms [44]. The parameters calculated from these two models are summarized in Table 3. The Langmuir model, based on the assumption of a monolayer adsorption onto a solid surface with a defined number of identical sites, gave the best fit of equilibrium adsorption data measured in a MOPS buffer. The Freundlich isotherm, which is an empirical model used to describe heterogeneous systems, best fitted the biosorption of cadmium in the presence of an excess of calcium ions. Both models showed that the EPS of \textit{R. alamii} actually bound cadmium. An apparent distribution coefficient of 3480 L.mg\(^{-1}\) was found at an EPS concentration of 0.125 g.L\(^{-1}\) and a maximum biosorption capacity of 11 mg of cadmium per g of EPS was determined. Both models showed that calcium, an abundant ion in soil solutions, competed with cadmium and reduced its sorption. At the highest EPS concentration tested (1 g.mL\(^{-1}\)), the EPS of \textit{R. alamii} showed the lowest ability to bind Cd\(^{2+}\) (apparent distribution coefficient of 387 L.mg\(^{-1}\) and maximum biosorption capacity of 6 mg of cadmium per g of EPS), showing that polysaccharide chain-chain interactions could preclude the binding of cadmium.

**Cadmium and biofilm matrix imaging by confocal laser scanning microscopy**

In Figure 6, the structure of cell organization, single cells, microcolonies and biofilms, from \textit{R. alamii} wt and the MS\textit{AGT} mutant, were examined after 3 days and 5 days of incubation at 30°C. The constitutive expression of GFP allowed the bacterial cells to be imaged. EPS was labelled with a fluorescent lectin (left lane only). Figure 6 shows typical pictures of the development of \textit{R. alamii} wt and MS\textit{AGT} mutant cells on a membrane, in the
The metal uptake, $X$ (mol.kg$^{-1}$), is determined using: $X = V.(C_0 - C)$, where $C_0$ and $C$ are the concentrations of initial and equilibrium cadmium in the solution (mol.L$^{-1}$), respectively; $V$ is the solution volume (L); $M$ is the mass of sorbent (kg). The error bars represent the confidence intervals (95%) of triplicates.

### Table 3. Isotherm parameters.

| R. alamii EPS % w/v | Medium | Langmuir Model | Freundlich model |
|---------------------|--------|---------------|-----------------|
|                     |        | $Q_m$ (mg/g)  | $K_L$ (L/mg)   | $r^2_L$ | $n$ | $K_f$ [(mg/g)(L/mg)$^{1/n}$] | $r^2_f$ |
| 0.0125              | MOPS   | $10^{-3}$     | $3\times10^{4}$ | 0.997   | 1.01 | 2.32 | 0.987 |
|                     | Calcium nitrate | $4\times10^{-4}$ | 47,868 | 0.680 | 2.00 | 1.03 | 0.991 |
| 0.1                 | MOPS   | $6\times10^{-3}$ | 3,870 | 0.999 | 1.13 | 0.60 | 0.987 |
|                     | Calcium nitrate | $3\times10^{-4}$ | 4,976 | 0.983 | 1.14 | 0.37 | 0.987 |

Isotherms parameters were determined from Langmuir and Freundlich models for Cd$^{2+}$ adsorption on the EPS of Rhizobium alamii in a MOPS buffer or calcium nitrate (10$^{-4}$ M) at pH 6.5. The reaction was maintained at 25°C under stirring, pH 6.5±0.2 for 24 h and 48 h to ensure that the equilibrium was reached.

The view of the role of rhizobia and their EPS being restricted mainly to symbiotic interactions with legumes is insufficient to gain insight into their physiology, and ability to adapt to environmental fluctuations. Altogether, our data has revealed some unique results, related to the metabolic response and lifestyle of R. alamii, and to the function of the EPS in response to cadmium. The belief that EPSs are the main players in metal tolerance of bacteria is persistent in microbiology, mainly in environmental science literature. Our experiments show that the bacterial polysaccharide (EPS) of R. alamii can bind cadmium, although this is not the way that these bacteria use to adapt to the metal toxicity. Changes of lifestyle, from planktonic to biofilm growth, and altering the metabolism, are the means to escape metal toxicity by wt cells as well as by mutant cells that no longer produce EPS. Further studies are being designed to determine the role of R. alamii and its EPS on plant cadmium uptake in the rhizosphere.

### Materials and Methods

#### Bacterial strains and plasmids

*Rhizobium alamii* wt (described in [24] and available at the Institut Pasteur CNCM I-1809P) and MSAGT (described in [23] and available upon request), were grown at 30°C in a tenfold diluted tryptic soy broth (DIFCO Laboratories, Detroit, USA), or in a minimal RCV medium [23], at pH 6.8, supplemented with glucose (2 g.L$^{-1}$) as a carbon source to favour EPS synthesis or in a minimal M9 medium (NaH$_2$PO$_4$ 6 g.L$^{-1}$, KH$_2$PO$_4$ 3 g.L$^{-1}$, NH$_4$Cl 1 g.L$^{-1}$, NaCl 0.5 g.L$^{-1}$, MgSO$_4$ 1 mM, Thiamine 50 mg.L$^{-1}$, L-Leucine 0.25 g.L$^{-1}$, L-Proline 0.25 g.L$^{-1}$, and glucose 2 g.L$^{-1}$). Nalidixic acid, kanamycin and tetracycline were respectively used at 50 μg.mL$^{-1}$, 25 μg.mL$^{-1}$ and 15 μg.mL$^{-1}$ for the appropriate antibiotic selection of R. alamii strains.

*Rhizobium alamii* wild type and its EPS-deficient mutant MSAGT labelled with green-fluorescent protein (GFP) [23,47] constitutively expressed the GFP for almost three months in soil, even in the absence of the antibiotic selection pressure.
Figure 6. Confocal laser-scanning micrographs of *Rhizobium alamii* wt and MSΔGT mutant. mg.L\(^{-1}\) of Cd\(^{2+}\) in the form of cadmium nitrate, 3 days (left lane) and 5 days (right lane). Bacteria were grown on polycarbonate membranes in a minimal M9 medium supplemented with 2 g.L\(^{-1}\) of glucose, with 0 or 2 mg.L\(^{-1}\) of cadmium following inoculation of the medium with 2.10\(^5\) cell.ml\(^{-1}\). The bacterial cells were localized by constitutive expression of GFP (in green); EPS was labeled by binding with fluorescent Alexa660-Concanavalin A (in red) in observations performed at 3 days of growth. 5-day biofilms were unlabeled for EPS. A) *R. alamii* wt after 3 days of growth in the absence of cadmium nitrate. Projections of Z-sections (1 \(\mu\)m) through 5 \(\mu\)m. The scale bar corresponds to 20 \(\mu\)m. Few cells were attached to the membrane surface as a small core. The EPS can be seen in red, surrounding a single cell. B) After 5 days of growth in the absence of cadmium. Projections of Z-sections (1 \(\mu\)m) through 5 \(\mu\)m. The scale bar corresponds to 50 \(\mu\)m. Bacterial cells colonized the membrane, in the form of a loose spherical biofilm. The EPS can be seen in red in some areas of the biofilm (see picture insert projection of Z-sections through 15 \(\mu\)m, the scale bar corresponds to 20 \(\mu\)m). C) *R. alamii* wt after 3 days of growth in the presence of cadmium nitrate (Cd\(^{2+}\) 2 mg.L\(^{-1}\)). Projections of Z-sections (1 \(\mu\)m) through 5 \(\mu\)m. The scale bar corresponds to 20 \(\mu\)m. The cells grew as a sphere. The EPS was localized inside the biofilm. D) *R. alamii* wt after 5 days of growth in the presence of cadmium nitrate (Cd\(^{2+}\) 2 mg.L\(^{-1}\)). Projections of Z-sections (1 \(\mu\)m) through 15 \(\mu\)m. The scale bar corresponds to 20 \(\mu\)m. The cells were organized in the form of fused, expanded spheres. E) MSΔGT mutant cell development on a membrane surface after 3 days of growth, in the absence of cadmium nitrate. Projection of Z-sections (1 \(\mu\)m) through 6 \(\mu\)m. The scale corresponds to 20 \(\mu\)m. The cells adhered to the membrane surface, although as single cells. F) MSΔGT mutant after 5 days of growth in the absence of cadmium. Projections of Z-sections (1 \(\mu\)m) through 17 \(\mu\)m. The scale bar corresponds to 50 \(\mu\)m. The bacterial cells formed a flat, loose biofilm architecture. G) MSΔGT mutant after 3 days of growth in the presence of cadmium nitrate (Cd\(^{2+}\) 2 mg.L\(^{-1}\)). Projection of Z-sections (1 \(\mu\)m) through 5 \(\mu\)m. The scale bar corresponds to 50 \(\mu\)m. Biofilms developed in the yz plane. 2 h. MSΔGT mutant after 5 days of growth in the presence of cadmium nitrate (Cd\(^{2+}\) 2 mg.L\(^{-1}\)). Projection of Z-sections (1 \(\mu\)m) through 17 \(\mu\)m. The scale bar corresponds to 50 \(\mu\)m. The cells formed a distributed, dense biofilm.

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Cadmium adsorption isotherms

EPS from *R. alamii* was isolated as described in [23], dialyzed and lyophilized prior to use. Cadmium binding to EPS of *R. alamii* was performed as described by Nelson et al. [48]. EPS solutions (0.5 to 4 g L⁻¹) in a MOPS buffer (10⁻² M), or Ca(NO₃)₂ (10⁻⁴ M), were placed inside dialysis tubing (Cellu-Sep® T3 Membranes, 12-14000 MWCO, Interchim, Montluçon, France) and equilibrated in 10 mL Cd(NO₃)₂ solutions (10⁻¹ to 10⁻¹¹ M) doped with ¹⁰⁰CdCl₂ (830 cpm mL⁻¹, 0.62 nCi final concentration). The pH was adjusted to 6.5±0.2, with NaOH 1N or HCl 1N. Triplicate samples were placed on a rotator at 22°C. Dialysis tubing with no EPS in solution were taken as blanks. Aliquots of the external solutions were taken at 24 and 48 h, and the γ-emissions were measured with a gamma counter.

Tolerance of *Rhizobium alamii* to cadmium

The wt and MSAGT strains were grown in a ten-fold diluted tryptic soy broth (TSB/10), or in a RCV supplemented with glucose (2 g L⁻¹) and various concentrations of cadmium (0 to 133.4 μM; 0 to 15 mg L⁻¹ of Cd²⁺). The OD₆₅₀ nm was measured over time from triplicate samples and reflected planktonic cell concentrations.

The *R. alamii* wt cells were grown up to an early stationary phase in TSB/10 supplemented with 0 or 2 mg L⁻¹ of cadmium nitrate. The cells were collected at OD₆₅₀ nm near to 0.07. Cytosolic cellular extracts were extracted in 50/50 methanol/water, in an ultrasonic bath for 15 min. The pellets were centrifuged at 10,000 rpm for 15 min, and the supernatant was analyzed using a Bruker-Daltonics APEX-Q 12 Tesla ICR-FT mass spectrometer (Bremen, Germany). The samples were introduced by macrospray infusion, at a flow rate of 120 μL h⁻¹, ionised in negative and positive electrosprays (ESI), and 256 scans were accumulated over a broadband range of masses (m/z 150-2000). The instrument was externally calibrated on clusters of arginine every measurement day, and the mass spectra were internally calibrated with phthalate diesters in positive ESI, and with fatty acids in negative ESI, thus ensuring a maximum error of 100 ppb. For the ICR-FT MS experiments, three replicates were analyzed for each condition (Figure S2), and the masses common to all three measurements were selected in order to generate a list of potential compounds. From the full set of detected compounds, we focused on those having a suggested attribution for the chemical structure with a difference between the detected and calculated masses of less than ±3 ppm, with detected peaks preferentially confirmed by the presence of ¹⁰⁸O, ¹⁵N or ¹³C isotopes. The peaks exceeding a threshold signal-to-noise ratio of 3 were exported to peak lists, and were submitted to a metabolite-annotation web interface, MassTRIX (http://masstrix.org). MassTRIX processes the submitted mass peak list by comparing the input experimental masses with all of the compounds recorded in the Kyoto Encyclopedia Genes and Genome (KEGG) chemical compound database, using *Rhizobium leguminosarum* as the model organism. The MassTRIX annotation of metabolites was used to highlight differences in metabolic pathways between the *R. alamii* cells incubated with or without Cd²⁺ 2 mg L⁻¹. Moreover, the compounds identified by ICR-FT/MS were matched with the KEGG reaction components. A full table of metabolites was added as supporting information Tables S1, S2, S3, S4. The raw data and the position of the potential metabolites in KEGG pathways can be consulted at http://metabolomics.helmholtz-muenchen.de/masstrix2, section “job status”, job access numbers 1003171000016906 and 10031709591316561 for wild type cells [Cd²⁺] 2 or 0 mg/L in negative mode, and 1003170954915503 and 10031709560715018, for wt cells [Cd²⁺] 2 or 0 mg/L in positive mode.

The analyses of pyrimidine and purine metabolites, with or without cadmium culture conditions, were performed by UPLC-MS on three independent replicates, which were characterised by similar profiles, indicating a high reproducibility of the three experiments (Figure S3 and Figure S4).

Adhesion of bacterial cells

The *R. alamii* wt and MSAGT mutant were statically grown to a stationary phase, in polycarbonate tubes containing RCV-medium, supplemented with 2 g L⁻¹ glucose and cadmium nitrate (0 to 133.4 μM; 0 to 15 mg L⁻¹ of Cd²⁺). Bacterial adhesion to solid surfaces and the formation of biofilms were monitored using a crystal violet staining-based protocol adapted from [39]. Briefly, the planktonic bacteria were removed from the tubes. Washing was performed with sterile water. After careful remove of the water, staining was performed with 2 mL of 1% crystal violet solution, 15 min at room temperature. The crystal violet solution was gently removed and successive washings were performed with water. Each tube was inverted and gently tapped on paper towels to remove any excess liquid and allow tubes to air-dry. 2 mL of 100% ethanol was added to each tube and OD₆₀₅ was measured in a cuvette on a spectrophotometer.

Microscopy

Biofilms of *R. alamii* wt and MSAGT mutant were statically grown on a polycarbonate membrane (Millipore 0.2 μm) incubated in a modified M9 medium ([Na₂HPO₄] 6 g L⁻¹, KH₂PO₄ 3 g L⁻¹, NH₄Cl 0.5 g L⁻¹, MgSO₄ 1 mM, thiamine 50 mg L⁻¹, L-Leucine 0.25 g L⁻¹, L-Proline 0.25 g L⁻¹ and glucose 20 g L⁻¹), supplemented with 0 or 2 mg mL⁻¹ of cadmium nitrate; 30 mL of M9 medium were inoculated with 2.10⁵ cells of *R. alamii* wt or MSAGT mutant, grown to half of the exponential phase in TSB/10, and washed with sterile ultrapurp water. EPS labelling with concanavalinA and confocal laser scanning microscopy were performed on an Olympus confocal microscope, as described in Sántaella et al. [23].

Data treatment

Statistical analyses (ANOVA) were made using version XV of the Statgraphics Centrion software.

Supporting Information

**Figure S1** Amount of EPS isolated from 5-day old *R. alamii* cultures (5 mL) in RCV medium (white bars) and in M9 modified medium (grey bars) supplemented with glucose (2 g L⁻¹). Mean of 3 replicates ± SD. (DOC)

**Figure S2** Examples of off-line FTMS measurements. (DOC)

**Figure S3** Examples of UPLC chromatograms of cell extracts of *R. alamii* grown without cadmium to the beginning of the stationary phase.UPLC-PDA, HSS column. Eluent: A: 10 mM ammonium acetate pH 6.4, B: 20% 10 mM ammonium acetate in acetonitrile. Flow: 0.9 mL min⁻¹ (pressure drop 600-900 bar), Column temperature: 40°C. Injection: 5 μL, partial loop with needle overfill. Wavelength detection 256 nm. Illustration of the reproducibility of the analyses from three repetitions of independent cell cultures and extracts. (DOC)
Table S1 Potential metabolites detected by ICR-FT MS negative ESI in R. alamii grown with no cadmium nitrate. (XLS)

Table S2 Potential metabolites detected by ICR-FT MS positive ESI in R. alamii grown with no cadmium nitrate. (XLS)

Table S3 Potential metabolites detected by ICR-FT MS negative ESI in R. alamii grown with 2 mg.L\(^{-1}\) of cadmium nitrate. (XLS)

Table S4 Potential metabolites detected by ICR-FT MS positive ESI in R. alamii grown with 2 mg.L\(^{-1}\) of cadmium nitrate. (XLS)

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Author Contributions
Conceived and designed the experiments: CS MS WA. Performed the experiments: MS CS AF CB. Analyzed the data: CS MS PO AF PS-K WA. Contributed reagents/materials/analysis tools: PO. Wrote the paper: CS WA.

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