

## Microbes Enhance Mobility of Arsenic in Pleistocene Aquifer Sand from Bangladesh

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**S** Supporting Information

**ABSTRACT:** Dissimilatory metal-reducing bacteria can mobilize As, but few studies have studied such processes in deeper orange-colored Pleistocene sands containing 1–2 mg kg<sup>-1</sup> As that are associated with low-As groundwater in Bangladesh. To address this gap, anaerobic incubations were conducted in replicate over 90 days using natural orange sands initially containing 0.14 mg kg<sup>-1</sup> of 1 M phosphate-extractable As (24 h), >99% as As(V), and 0.8 g kg<sup>-1</sup> of 1.2 M HCl-leachable Fe (1 h at 80 °C), 95% as Fe(III). The sediment was resuspended in artificial groundwater, with or without lactate as a labile carbon source, and inoculated with metal-reducing *Shewanella* sp. ANA-3. Within 23 days, dissolved As concentrations increased to 17 μg L<sup>-1</sup> with lactate, 97% as As(III), and 2 μg L<sup>-1</sup> without lactate. Phosphate-extractable As concentrations increased 4-fold to 0.6 mg kg<sup>-1</sup> in the same incubations, even without the addition of lactate. Dissolved As levels in controls without *Shewanella*, both with and without lactate, instead remained <1 μg L<sup>-1</sup>. These observations indicate that metal-reducers such as *Shewanella* can trigger As release to groundwater by converting sedimentary As to a more mobilizable form without the addition of high levels of labile carbon. Such interactions need to be better understood to determine the vulnerability of low-As aquifers from which drinking water is increasingly drawn in Bangladesh.

### INTRODUCTION

Shared deep wells, typically more than 150 m deep and installed by the government or nongovernmental organizations, have emerged over the past decade as an important means of reducing the exposure to As of the rural population of Bangladesh.<sup>1</sup> There are, however, concerns about the risk of contaminating the deep aquifer with As drawn down from shallow depths, especially if crop irrigation, which accounts for about 90% of all groundwater use in the country, were to tap the deep aquifer.<sup>2</sup> The concern is that, in addition to transporting As into the deep aquifer, intruding shallow groundwater could potentially stimulate biogeochemical transformations that release labile As from the sediment to groundwater.<sup>3</sup>

A number of microcosm experiments using natural soils and aquifer sands from West Bengal (India), Bangladesh, and Cambodia have demonstrated the release of As following the addition of electron donors in various forms, although typically starting from shallow gray sands containing at least partially reduced amorphous iron minerals.<sup>4–7</sup> A subset of these experiments paired with microbial community analysis has shown that metal-reducing bacteria such as *Geobacter* play a role in releasing As to groundwater either by reducing As-bearing Fe oxides or by reducing sediment-bound As(V) to As(III).<sup>4,7</sup> Column experiments using ferrihydrite-coated sand loaded with As have separated the role of Fe and As

reduction using modified strains of the genus *Shewanella* in the laboratory.<sup>8</sup> The need for substantial reduction of Fe oxides to mobilize As is fairly well established after a decade of research,<sup>9–11</sup> but whether reduction of As accompanies its release to solution or is a prerequisite for mobilization in natural systems remains unclear.

The need for Fe and As reduction is re-examined here on the basis of incubations carried out anaerobically over a period of three months using orange-colored sand from Bangladesh containing primarily Fe(III) and As(V). The choice of starting material reflects the importance of such less-reduced, but still anoxic, aquifers as a source of low-As groundwater.<sup>3,9</sup> Deeper wells tapping such aquifers have reliably supplied numerous villages of Bangladesh with low-As groundwater for over a decade, although a few failures due to increasing As concentrations have also been reported.<sup>12</sup> The cause of these failures is presently not known. A shallow break in a PVC pipe could result in high-As groundwater entering a well, but so could potentially a regional intrusion of shallow groundwater into a deeper aquifer. This study was designed to simulate certain aspects of a regional intrusion. The orange sands used in the

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incubations are representative of deeper aquifers in Bangladesh of likely Pleistocene age. Total As concentrations in the orange sands are close to its crustal abundance of 1–2 mg kg<sup>-1</sup> 13 and approximately an order-of-magnitude lower than in shallower gray sands associated with groundwater elevated in As.<sup>3,9</sup>

The orange sands were inoculated with *Shewanella* sp. strain ANA-3 which is a well-characterized As(V) and Fe(III) reducer that has been used for incubations of synthetic Fe(III)-oxide coated sands containing As.<sup>8,14–16</sup> Although *Shewanella* species are not major constituents of subsurface environments, there is considerable genomic evidence that the enzymes and genes for As(V) reduction are highly conserved among very diverse bacteria,<sup>10</sup> some of which have been detected in aquifer sediments of West Bengal and Cambodia.<sup>7,17,18</sup> *Shewanella* also uses enzyme pathways for Fe(III) reduction similar to *Geobacter* which, in contrast, has been shown to be ubiquitous in the subsurface.<sup>19</sup> *Shewanella* sp. strain ANA-3 contains, in addition, the plasmid pBBR1-MCS2 that provides resistance to the antibiotic kanamycin. The intention was to separate the effect of a well-studied organism such as *Shewanella* from that of the complex natural assemblage of micro-organisms present in the sediment by amending the incubations with kanamycin. Although the kanamycin did not fully suppress the activity of the native micro-organisms, the incubations clearly demonstrate that *Shewanella* amendments can trigger the release of As from orange sands without extensive reduction of Fe oxides driven by a large addition of metabolizable carbon.

## EXPERIMENTAL SECTION

**Sediment Collection and Preservation.** A core section from 43.3 to 43.9 m depth in Dari Satyabandi village (23.785°N, 90.603°E) in Araihasar, Bangladesh, was used in all experiments after manually homogenizing the sand inside an anaerobic chamber containing 95% N<sub>2</sub>:5% H<sub>2</sub> (Coy Lab. Prod. Inc., AALC). Both ends of the PVC liner containing the core were wax-sealed on site immediately after collection in January 2001.<sup>3</sup> The cores were shipped at ambient temperature and stored at 4 °C upon their arrival at Lamont-Doherty Earth Observatory's core repository in May 2001. When the liner was opened inside the anaerobic chamber in July 2005, the sediment was still orange and moist. The presence of water inside the core after four years of storage in a refrigerated and dry environment indicates that the wax seal was intact. Exposure of the core to atmospheric oxygen that slowly diffuses through the wax and the PVC liner was therefore limited.

**Shewanella Culture.** *Shewanella* sp. strain ANA-3 was previously isolated from an As-treated wooden pier piling in Woods Hole, MA.<sup>14,16</sup> The strain was initially grown aerobically and then anaerobically in TME media<sup>20</sup> on an incubator shaker (250 rpm) at 28–30 °C for 24 and 72 h, respectively. The TME media for aerobic growth was prepared by adding 1.5 g of NH<sub>4</sub>Cl, 0.6 g of NaH<sub>2</sub>PO<sub>4</sub>, 0.1 g of KCl, 0.5 g of yeast extract (50 mL of autoclave sterilized stock solution of 10 g L<sup>-1</sup>), 20 mmol L<sup>-1</sup> Na-lactate (20 mL of autoclave sterilized stock solution of 1 mol L<sup>-1</sup>), and 10 mL of trace metal (modified Wolf's mineral mix) solution<sup>20</sup> in Milli Q water (18.1 MΩ) to make 1 L of solution. The pH was adjusted to 7 by adding 1 g of Na-HEPES to 1 L of the media. For anaerobic growth of *Shewanella*, an anaerobic electron acceptor was added to TME media. In this case, 1.56 g of Na<sub>2</sub>HAsO<sub>4</sub>·7H<sub>2</sub>O was added to 500 mL aliquots of initial TME media to reach an As concentration of 10 mmol L<sup>-1</sup>.<sup>16</sup> The TME media was autoclave sterilized prior to use for aerobic and anaerobic growth. *Shewanella* sp. ANA-3 strain containing plasmid

pBBR1-MCS2 was cultured with 50 μg mL<sup>-1</sup> kanamycin (further details provided in Supporting Information).

**Incubations.** The orange sands were incubated anaerobically for 3 months with five different types of amendments. Three sets of controls were incubated without *Shewanella*: the first, labeled C<sub>0</sub>, contained only sediment and artificial groundwater; the second (C<sub>lac</sub>) was amended with lactate (Fisher, syrup 60% w/w) to a concentration of 20 mmol L<sup>-1</sup>; the third (C<sub>lac+kan</sub>) was also amended with the same amount of lactate and with 50 μg mL<sup>-1</sup> of the antibiotic kanamycin. Incubations inoculated with *Shewanella* sp. ANA-3 were all amended with 50 μg mL<sup>-1</sup> kanamycin, one with 20 mmol L<sup>-1</sup> lactate (S<sub>lac+kan</sub>) and the other without (S<sub>kan</sub>).

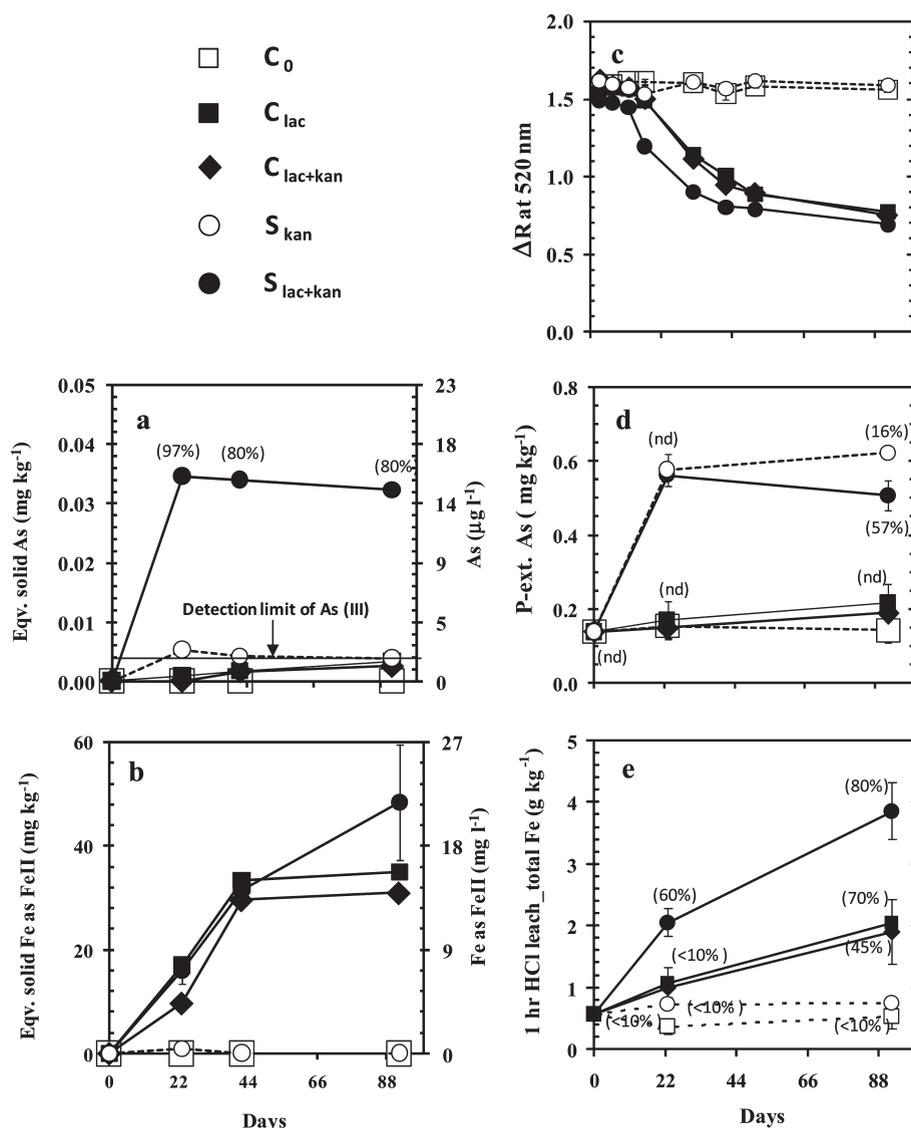
For all incubations, approximately 5 g of sand was added to sterile 17-mL culture tubes (Hungate 2047–16125) with 10 mL of artificial groundwater previously purged with ultrapure N<sub>2</sub> for 2 h. During preparation of culture tubes, tubes with sediment were also purged with ultrapure N<sub>2</sub> and capped immediately after addition of artificial groundwater. Artificial groundwater of composition similar to that of groundwater associated with the Pleistocene aquifer in Dari Satyabandi<sup>3</sup> was prepared by adding salts of Na<sup>+</sup>, Ca<sup>2+</sup>, K<sup>+</sup>, Mg<sup>2+</sup>, Cl<sup>-</sup>, HCO<sub>3</sub><sup>-</sup>, PO<sub>4</sub><sup>3-</sup>, and SO<sub>4</sub><sup>2-</sup> to Milli Q water (Table S1, Supporting Information) followed by filtration through a disposable sterile system (Corning, 09-761). The initial pH of 7.9 of the artificial groundwater was higher than for groundwater pumped from the orange aquifer in the village (pH 6.5 ± 0.1).<sup>21</sup>

The optical density at 600 nm of 0.05 of the *Shewanella* cultures indicate that the incubations were inoculated with ~5 × 10<sup>8</sup> cells per 10 mL culture tube.<sup>14</sup> The tubes were wrapped in black plastic and set horizontally on an orbital shaker (Cole-Parmer, 51401-15) gently rotating at 50–75 rpm over the entire period. Each set of incubations started with seven replicates that were sampled sacrificially on day 23 in triplicate, on day 42 in duplicate, and on day 92 in duplicate. The sampling intervals were chosen by monitoring the color change of the sand suspension. After measurement of the pH (Orion, model 260A), the supernatant from each sampled tube was filtered in the anaerobic chamber through a 0.23 μm filter.

**Sediment Reflectance.** Changes in the color of the sediment were recorded over the course of the incubations with a Minolta CM2500D diffuse spectral reflectance spectrophotometer through the round-bottom of the glass tubes. The difference in reflectance (ΔR) between wavelengths of 530 and 520 nm was used as a measure of conversion of Fe(III) to Fe(II) within the solid phase.<sup>22</sup>

**As and Fe Concentrations and Speciation.** Concentrations of As(III) in the filtered but unacidified supernatant were determined immediately after opening a culture tube by differential pulse cathodic stripping voltammetry (DPCSV).<sup>23</sup> The detection limit of this method is 0.2 μg L<sup>-1</sup> (3 nmol L<sup>-1</sup>). Dissolved Fe(II) was also measured immediately in the filtered unacidified supernatant by colorimetry (detection limit of 30 μg L<sup>-1</sup>) using ferrozine.<sup>24</sup> The concentrations of total As and Fe in the filtered supernatant acidified to 1% Optima HCl were analyzed by high-resolution inductively coupled plasma spectrometry (HR ICP-MS).<sup>25</sup> The detection limit of the method for As is 0.1 μg L<sup>-1</sup> (0.001 μmol L<sup>-1</sup>).

To analyze the sediment, one 0.5 g aliquot of wet sediment was leached in 10 mL of 1.2 mol L<sup>-1</sup> HCl at 80 °C for 1 h, after which the concentration of Fe(II) and Fe(III) in the leachate was measured with ferrozine.<sup>22</sup> Another 0.5 g of wet sediment from each tube was leached for 24 h in 10 mL of 1 M Na<sub>2</sub>HPO<sub>4</sub> (pH 5) and 0.1 M ascorbic acid at room temperature in the anaerobic chamber. This extraction has been shown to release As strongly bound to goethite, presumably by competitive desorption.<sup>26</sup> This procedure, previously applied to Bangladesh sediment by Harvey



**Figure 1.** Variations in As and Fe concentrations and speciation in the dissolved and the solid phase over the course of incubations of orange sand from Bangladesh in artificial groundwater. The five sets of incubations include a control without lactate ( $C_0$ ), a control with lactate ( $C_{lac}$ ), a control with lactate and kanamycin ( $C_{lac+kan}$ ), an amendment with *Shewanella* in kanamycin without lactate ( $S_{kan}$ ) and amendment with *Shewanella* in kanamycin with lactate ( $S_{lac+kan}$ ). Liquid phase (a) As and (b) Fe measurements are expressed as dissolved concentrations (right scale) as well as equivalent concentrations in the solid phase (left scale) by normalizing to sediment weight. Error bars indicate the standard deviation of triplicates on day 22 and duplicates on day 45 and day 90. Some error bars are covered by the symbol. “nd” indicates As(III) was not detected by DPCSV with a detection limit  $2 \mu\text{g L}^{-1}$  ( $0.004 \text{ mg kg}^{-1}$ ). (c) Changes in sediment color expressed as the difference in reflectance between 530 and 520 nm ( $\Delta R$ ). Also shown are (d) solid phase P-extractable total As, with symbols labeled with the proportion of as As(III) and (e) solid phase 1 h HCl-leachable Fe, with the proportion of Fe(II).

et al.,<sup>27</sup> relies on a phosphate concentration that is 10-fold higher than the one used to extract As from aquifer sands in, for instance, Vietnam.<sup>28,29</sup> Subsequent experiments have shown that the addition of ascorbic acid preserves the oxidation state of adsorbed As without causing significant reductive dissolution of Fe oxides.<sup>30</sup> The concentration of As(III) in the phosphate extract was also measured by DPCSV<sup>23</sup> and the concentration of total As by HR ICP-MS.<sup>25</sup>

## RESULTS

**Initial Conditions.** Before the incubations, the hot HCl-leachable Fe concentration was  $0.8 \text{ g kg}^{-1}$  and the P-extractable As concentration was  $0.14 \pm 0.03 \text{ mg kg}^{-1}$  ( $n = 3$ ), mostly

(>95%) in the form of Fe(III) and As(III), respectively. The Fe(II)/Fe ratio <0.05 in the leachable Fe fraction and the diffuse spectral reflectance are consistent with previous analyses of Pleistocene sediment from the same area.<sup>3,22</sup> The initial difference in reflectance at 530 and 520 nm ( $\Delta R$ ) of 1.5 (Figure 1c) is higher than previously measured values of  $\sim 1.0$  for freshly collected orange sands, probably because the reflectance was measured through the bottom of the culture tube instead of through Saran Wrap in direct contact with the sediment. In this study, changes in reflectance therefore provide an effective, albeit qualitative, indication of the redox state of acid-leachable Fe oxides over the course of the experiments.

**Aqueous As and Fe.** The most significant changes in As concentrations occurred within the first 23 days of the incubations

Table 1. Aqueous and Postincubated Sediment Properties and As, Fe, Mn Concentrations after 23 Days

		Liquid Phase (solid equivalent)				
		controls			<i>Shewanella</i> sp. ANA-3	
AGW (artificial groundwater)		$C_0$	$C_{lac}$	$C_{lac+kan}$	$S_{kan}$	$S_{lac+kan}$
As $\mu\text{g L}^{-1}$ ( $\text{mg kg}^{-1}$ )	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>	$2.6 \pm 0.01$ (0.005)	$17.3 \pm 3.7$ (0.035)
As(III) $\mu\text{g L}^{-1}$ ( $\text{mg kg}^{-1}$ )	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>	$16.1 \pm 1.9$ (0.032)
Fe $\text{mg L}^{-1}$ ( $\text{mg kg}^{-1}$ )	nd <sup>a</sup>	nd <sup>a</sup>	$7.6 \pm 0.6$ (17)	$4.4 \pm 0.4$ (10)	$0.4 \pm 0.04$ (1)	$7.3 \pm 1.1$ (16)
Fe(II) $\text{mg L}^{-1}$ ( $\text{mg kg}^{-1}$ )	nd <sup>a</sup>	nd <sup>a</sup>	$7.6 \pm 0.7$ (17)	$4.3 \pm 0.4$ (9)	$0.4 \pm 0.06$ (1)	$7.2 \pm 1.1$ (16)
Mn $\text{mg L}^{-1}$ ( $\text{mg kg}^{-1}$ )	nd <sup>a</sup>	nd <sup>a</sup>	$7.4 \pm 0.9$ (17)	$8.1 \pm 0.8$ (18)	$0.8 \pm 0.08$ (2)	$7.9 \pm 0.2$ (18)
pH	7.88	$8.13 \pm 0.02$	$7.41 \pm 0.02$	$7.41 \pm 0.02$	$7.89 \pm 0.02$	$7.13 \pm 0.01$

		Solid Phase				
		controls			<i>Shewanella</i> sp. ANA-3	
pre-incubated sediments		$C_0$	$C_{lac}$	$C_{lac+kan}$	$S_{kan}$	$S_{lac+kan}$
color	orange	orange	orange	orange	orange	gray
$[\text{As}_{\text{solid}}]/[\text{As}_{\text{liquid}}]$ (L/kg)		>320 <sup>b</sup>	>320 <sup>b</sup>	>320 <sup>b</sup>	218	33
P-extractable As ( $\text{mg kg}^{-1}$ )	$0.14 \pm 0.03$	$0.15 \pm 0.01$	$0.17 \pm 0.05$	$0.15 \pm 0.03$	$0.58 \pm 0.04$	$0.56 \pm 0.01$
P-extractable As(III) ( $\text{mg kg}^{-1}$ )	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>	nd <sup>a</sup>
24 h HCl leachable Fe ( $\text{g kg}^{-1}$ )	$10.6 \pm 1.2$	$12 \pm 0.6$	$12 \pm 0.8$	$11 \pm 0.7$	$11 \pm 1.6$	$12 \pm 0.3$
1 h HCl leachable Fe ( $\text{g kg}^{-1}$ )	$0.79 \pm 0.15$	$3.43 \pm 0.69$	$5.63 \pm 0.63$	$5.11 \pm 0.83$	$2.94 \pm 0.12$	$5.4 \pm 0.04$
1 h HCl leachable Fe(II) ( $\text{g kg}^{-1}$ )	$0.04 \pm 0.002$	$0.2 \pm 0.02$	$0.41 \pm 0.01$	$0.33 \pm 0.02$	$0.2 \pm 0.03$	$3.2 \pm 0.02$
HCl leachable Fe(II)/Fe ratio	0.051	0.061	0.073	0.065	0.068	0.593
HCl leachable Mn ( $\text{g kg}^{-1}$ )	$0.2 \pm 0.05$	$0.2 \pm 0.09$	$0.3 \pm 0.03$	$0.2 \pm 0.04$	$0.2 \pm 0.04$	$0.2 \pm 0.02$

<sup>a</sup> Not detectable. <sup>b</sup> Liquid phase As was not detectable. Limit of detection (LOD) for As and Fe/Mn by HR ICPMS are  $0.10 \mu\text{g L}^{-1}$  ( $0.0002 \text{ mg kg}^{-1}$ ) and  $1 \mu\text{g L}^{-1}$  ( $0.002 \text{ mg kg}^{-1}$ ), respectively. As(III) by DPCSV is  $2 \mu\text{g L}^{-1}$  ( $0.004 \text{ mg kg}^{-1}$ ) and Fe is  $0.03 \text{ mg L}^{-1}$  ( $0.06 \text{ mg kg}^{-1}$ ) by Ferrozine method.

(>Figure 1a; Table 1). In the amendments with lactate inoculated with *Shewanella* sp. strain ANA-3 ( $S_{lac+kan}$ ), concentrations of dissolved As rose to  $17 \pm 4 \mu\text{g L}^{-1}$  ( $n = 3$ ), which is equivalent to a release of  $0.035 \text{ mg kg}^{-1}$  from the sand (Table 1). The increase in dissolved As over the same period was smaller ( $2.6 \pm 0.01 \mu\text{g L}^{-1}$ ; equiv to  $0.005 \text{ mg kg}^{-1}$ ), but easily detectable by HR ICP-MS for the inoculated incubation without lactate ( $S_{kan}$ ). No increase above the detection limit of  $0.1 \mu\text{g L}^{-1}$  dissolved As was observed on day 23 in the supernatant of the three incubations without *Shewanella* strain ( $C_0$ ,  $C_{lac}$ ,  $C_{lac+kan}$ ), two of which were amended with lactate. Aqueous As remained undetectable over three months in the control without amendment ( $C_0$ ), whereas aqueous As in the other two controls amended with lactate ( $C_{lac}$ ,  $C_{lac+kan}$ ) gradually increased to  $1.7$  and  $1.3 \mu\text{g L}^{-1}$  As ( $0.003$  and  $0.002 \text{ mg kg}^{-1}$ ) (Figure 1a; Table S2, Supporting Information).

Most of the As released from the orange sand in the amendments with lactate and *Shewanella* ( $S_{lac+kan}$ ) was in the form of As(III) on day 23 and remained in that form for the duration of incubations (Figure 1a). The proportion of dissolved As(III) in the incubation with *Shewanella* without lactate ( $S_{kan}$ ) and all the controls could not be determined because concentrations were below the detection limit  $<2 \mu\text{g L}^{-1}$  by DPCSV.

In contrast to As, the incubations of orange sands amended with lactate systematically released more Fe than the incubations without lactate. Concentrations of dissolved Fe(II) in all three lactate amendments ( $C_{lac}$ ,  $C_{lac+kan}$ ,  $S_{lac+kan}$ ) rose above  $4 \text{ mg L}^{-1}$  by day 23 (Table 1) and stabilized to  $\sim 15 \text{ mg/L}$  by day 42 (Figure 1b; Table S2, Supporting Information). The maximum of Fe released to solution by the end of the incubations corresponds to  $<1\%$  of the

total Fe in the acid-leachable fraction in the original sand. Dissolved Fe(II) remained below the detection limit of  $0.03 \text{ mg L}^{-1}$  throughout the incubations in both incubations without lactate ( $C_0$ ,  $S_{kan}$ ).

**Sediment As and Fe.** The concentrations of P-extractable As in the sediment rose 4-fold to  $0.6 \text{ mg/kg}$  by day 23 in all six tubes inoculated with *Shewanella*, with or without lactate ( $S_{lac+kan}$ ,  $S_{kan}$ ), and remained constant thereafter (Figure 1d). In marked contrast, concentrations of P-extractable As remained below  $0.23 \text{ mg kg}^{-1}$  throughout the experiment for all the other amendments without *Shewanella*, including those amended with lactate ( $C_0$ ,  $C_{kan}$ ,  $C_{lac+kan}$ ).

The quantity of As extracted from the solid phase in 1 M phosphate is 20-fold greater than the total released to the supernatant over the same period for the incubations inoculated with *Shewanella* and lactate ( $S_{lac+kan}$ ). Whereas As released to the supernatant was primarily in the form of As(III), the P-extractions of the sediment in both sets of inoculated incubations contain no detectable As(III) through day 23 ( $S_{kan}$ ,  $S_{lac+kan}$ ) (Figure 1d). The proportion of As(III) in the P-extractions rose to 20 and 60% in the  $S_{kan}$  and  $S_{lac+kan}$  amendments by the end of the incubations, respectively.

The orange sediment amended with lactate and *Shewanella* ( $S_{lac+kan}$ ) turned gray by day 17, followed by the two incubations with lactate but without *Shewanella* ( $C_{lac}$ ,  $C_{lac+kan}$ ) on day 23. Evidently, the kanamycin did not suppress Fe reduction by the indigenous population of microbes. The remaining amendments without lactate ( $S_{kan}$ ,  $C_0$ ), including the one with *Shewanella*, remained orange for the entire period. The timing of the color changes is consistent with the changes in reflectance (Figure 1c). The proportion of Fe(II)/Fe in the 1 h HCl-leachable Fe fraction rose to 60% by day 23 in the incubation with lactate and

*Shewanella* ( $S_{\text{lac+kan}}$ ) (Figure 1e). In all other incubations with lactate or *Shewanella*, the Fe(II)/Fe ratio remained <10% (Table 1). The leachable Fe(II)/Fe ratio continued to rise as the concentration of total leachable Fe in the incubations amended with lactate ( $C_{\text{lac}}$ ,  $C_{\text{lac+kan}}$ ,  $S_{\text{lac+kan}}$ ) further increased over the course of the incubations (Table S3, Supporting Information). In incubations without lactate ( $C_0$ ,  $S_{\text{kan}}$ ), neither the HCl-leachable Fe concentration nor the HCl-leachable Fe(II)/Fe ratio changed over the entire incubation period. The  $\Delta R$  of  $\sim 0.7$  for gray sediment at the end of the incubations with lactate was again 0.4 unit higher than that observed in the field with freshly collected gray sediment.<sup>22</sup> The reflectance of the incubations without lactate did not change over the course of the experiments.

## DISCUSSION

**Decoupling of As and Fe Release.** Incubations of sediment from South and Southeast Asia have to date relied primarily on additions of acetate or lactate to stimulate the reduction of Fe oxides and the transfer of As from the solid to the dissolved phase. A limited number of studies have relied on natural organic matter and have attempted to characterize the role of bacteria by including abiotic controls and characterizing microbial populations.<sup>4,5,7,31</sup> The incubations presented here differ from previous work because of the nature of the starting material, orange as opposed to gray sands, and because well-characterized bacteria were added with and without lactate. The most surprising result is that the additions of *Shewanella* alone led to a substantial increase in the concentrations of P-extractable As in the solid phase as well as a release of As to solution, whereas the additions of lactate without *Shewanella* caused only a marginal increase in the concentrations of P-extractable As in the solid phase. The additions of lactate triggered the anticipated conversion of Fe (III) to Fe(II) within the solid phase as well as reductive dissolution, but without a concomitant release of As in the absence of *Shewanella*. This extent of decoupling between the release of As and Fe has been reported for model systems but, to our knowledge, rarely for natural sediment.

**Underlying Mechanisms.** The interpretation of results from this pilot-study is limited by a number of factors. It appears that extended storage of the sediment core had little impact on the speciation of As and Fe, although confirmation by X-ray spectroscopy<sup>6,7</sup> would have been useful. The initial predominance of As(V) in the orange sands is unlikely to be a storage artifact because it has also been reported on the basis of X-ray spectroscopy for freshly collected and preserved orange sands from Bangladesh.<sup>32</sup> However, no information is available to determine to what extent the original microbial population was affected by storage and how it evolved in response to the various treatments. *Shewanella* has not been reported in natural sediment of South and Southeast Asia, although the presence of bacteria with similar properties has been documented.<sup>17–19</sup> The pH of the experiments was high compared to natural conditions, even in the incubations that were amended with lactate (pH 7.1–7.4 at day 23 vs pH 6.5 in the local groundwater). It is also not known to what extent the sulfate initially contained in the artificial groundwater was reduced and whether this conversion could have resulted in precipitation of As-sulfides and, therefore, could have limited the increase in dissolved As.<sup>7</sup> Finally, the amount of kanamycin added was insufficient to inhibit fully microbial activity, particularly that of Fe reducers, in the presence of lactate.

This could have been because of resistance of the indigenous population to kanamycin and limits our ability to distinguish biotic and abiotic processes regulating dissolved As levels in the experiments.

Despite these limitations, our results show that the additions of *Shewanella* with lactate and only the additions of *Shewanella* resulted in a significant conversion of As from a relatively immobile to an exchangeable phase. The extraction method used to quantify this exchangeable phase has previously been shown to effectively release As adsorbed onto ferrihydrite and goethite into solution.<sup>30</sup> Because the concentration of phosphate is high, the extraction could potentially overestimate the pool of mobile arsenic by releasing As that is incorporated into the mineral lattice. On the other hand, previous work using natural sediments from the Bengal Basin has shown that the phosphate extraction procedure applied to the incubations provides a useful operational definition of a pool of As in the solid phase that is related in a fairly systematic way to dissolved concentration of As in groundwater.<sup>3,33</sup>

It is presently unclear which properties of *Shewanella* led to the conversion to an exchangeable form of As or whether the inoculations provided a source of metabolizable carbon that was utilized, instead, by other resident bacteria that were not inhibited by kanamycin. For comparison with the amount of the lactate added (600  $\mu\text{mol}$  of C per tube), the quantity of reduced carbon in the form of bacterial cells added was on the order of 1  $\mu\text{mol}$  C per tube, assuming a carbon content of the natural bacterial assemblage of 20 fg of C cell<sup>-1</sup>.<sup>34</sup> The quantity of reduced carbon added in the form of bacteria was clearly insufficient to release as Fe(II) the equivalent of 50 mg kg<sup>-1</sup> (Figure 1b), i.e. 5  $\mu\text{mol}$  Fe per tube, but considerably larger than the maximum of 0.03 mg kg<sup>-1</sup>, i.e. 0.002  $\mu\text{mol}$  per tube, of As transformed to a more exchangeable form.

It is worth pointing out that the importance of As reduction for As release from natural orange sands remains uncertain in these experiments. On one hand, the incubation with *Shewanella* and lactate ( $S_{\text{lac+kan}}$ ) showing the highest degree of As reduction in the solid phase at the end of the incubations resulted in the largest release of As concentrations to the dissolved phase, and the solution speciation was dominated by As(III) for this incubation (Figure 1a,b). Taken at face value, this could imply that reduction enhances As release to solution. On the other hand, the marked difference in the extent of reduction of sedimentary As(V) at the end of the experiment between  $S_{\text{kan}}$  and  $S_{\text{lac+kan}}$  did not affect the total increase in P-extractable As. It is not clear which of the two processes, the short-term release of As to solution or the conversion to P-extractable As, is ultimately responsible for increasing groundwater As levels under natural conditions. Another reason the role of As reduction is difficult to ascertain from these experiments is that As can be sequestered in remineralized solids, by analogy to the impact of transformations from ferrihydrite to magnetite documented in model systems.<sup>35</sup>

**Implications for Mitigation.** The solid to water ratio in the culture tubes was about 12 times lower than in the sandy aquifer, assuming a typical porosity of 0.3. All other things being equal, the impact of introducing a similar dose of *Shewanella* in the field could therefore conceivably have increased dissolved As concentrations to 31  $\mu\text{g L}^{-1}$  without further amendment and 210  $\mu\text{g L}^{-1}$  with lactate or an equivalent source of metabolizable carbon. The lower concentrations of dissolved As, which still exceeds the WHO guideline for As in drinking water of 10  $\mu\text{g L}^{-1}$ , is probably more realistic because it was observed without any major change in the redox state of Fe on the sediment surface.

Our results suggest that, in the presence of a significant hydraulic head gradient that favors downward transport of groundwater as in Dari Satyabandi,<sup>3</sup> shallow groundwater with a different resident population of bacteria and, potentially, higher levels of metabolizable carbon could intrude into the deeper low-As aquifer and trigger the release of As. There is also growing evidence of preferential transport of bacteria in sandy aquifers,<sup>36</sup> suggesting that imperfect sealing between the shallow and deep aquifer could be conducive to transferring bacteria and therefore release of As even from orange sands. As villagers of Bangladesh increasingly rely on deep community wells to meet their needs for safe drinking water, the potential vulnerability of low-As aquifers clearly deserves further study.

## ■ ASSOCIATED CONTENT

**S Supporting Information.** Additional information as noted in the text. This material is available free of charge via the Internet at <http://pubs.acs.org/>.

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