

# Mobilization of Arsenic During One-Year Incubations of Grey Aquifer Sands from Araihazar, Bangladesh

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Elevated As concentrations in shallow groundwater pose a major health threat in Bangladesh and similarly affected countries, yet there is little consensus on the mechanism of As release to groundwater or how it might be influenced by human activities. In this study, the rate of As release was measured directly with incubations lasting 11 months, using sediment and groundwater collected simultaneously in Bangladesh and maintained under anaerobic conditions throughout the study. Groundwater and gray sediment were collected as diluted slurries between 5 and 38 m in depth, a range over which ambient groundwater As concentrations increased from 20 to 100  $\mu\text{g L}^{-1}$ . Arsenic was released to groundwater in slurries from 5 and 12 m in depth at a relatively constant rate of  $21 \pm 4$  ( $2\sigma$ ) and  $23 \pm 6 \mu\text{g As kg}^{-1} \text{yr}^{-1}$ , respectively. Amendment with a modest level of acetate increased the rate of As release only at 12 m ( $82 \pm 18 \mu\text{g kg}^{-1} \text{yr}^{-1}$ ). Although the groundwater As concentration was initially highest at 38 m depth, no release of As was observed. These results indicate that the spatial distribution of dissolved As in Bangladesh and local rates of release to groundwater are not necessarily linked. Iron release during the incubations did not occur concurrently with As release, providing further confirmation that the two processes are not directly coupled. Small periodic additions of oxygen suppressed the release of As from sediments at all three depths, which supports the notion that anoxia is a prerequisite for accumulation of As in Bangladesh groundwater.

## Introduction

An estimated 100 million people are exposed to elevated levels of As by drinking untreated groundwater throughout

South Asia. In Bangladesh, one-third of all wells did not meet the Bangladesh drinking water standard of  $50 \mu\text{g L}^{-1}$  and one-half did not meet the World Health Organization guideline of  $10 \mu\text{g L}^{-1}$  before mitigation efforts began (1, 2). It is widely believed that elevated As concentrations in anoxic groundwater reflect desorption from Fe oxyhydroxide surfaces as Fe(III) is converted to dissolved Fe(II) or mixed Fe(II/III) surfaces (2, 3). Extensive field surveys show, however, that the relationship between dissolved As and Fe concentrations is highly scattered in Bangladesh, although their concentrations in the sediment are highly correlated (2, 4). More recently, detrital sulfide phases have been proposed as an alternative shallow source of As that is mobilized through redox cycling in near-surface sediments in Bangladesh (5). The importance of Fe reduction has been further brought into question by observations in model systems of significant As mobilization with only moderate reduction of Fe(III) minerals and before significant releases of Fe(II) into solution (6–8). The role of As reduction per se for mobilization is also still unclear. Studies using natural sediments and model sediments inoculated with Fe- or As-reducing microbes have shown enhancement of mobilization when arsenate is reduced to As(III) (6, 10). But other studies have shown that As(III) is not necessarily more mobile than As(V) under all environmental conditions, particularly when phosphate is present (11, 12).

It is generally accepted that microbial activity influences As release, either through direct interactions or indirectly via Fe oxyhydroxides, although the nature of this relationship remains poorly understood (2, 13). Organic substrate additions in both field and incubation studies, meant to enhance microbially mediated As mobilization have shown mixed results (5, 10, 14, 15, 17). In addition, the extent to which large substrate additions are representative of the process of As release under natural conditions is unclear since major shifts in the population of dominant groups of the microbial community in response to substrate additions have been observed (15). The provocative notion that irrigation pumping might have enhanced the supply of organic substrate to shallow aquifers in recent decades, hereby triggering the release of As, has also been put forth (14, 16).

The motivation for the present study is that some of these unsettled issues could be addressed by knowing the rate at which As is released in aquifers in different geological settings. Previous incubations using natural aquifer sediment have provided valuable mechanistic insights, but intentional or unintentional manipulations of the material have hampered the extrapolation of measured rates to natural conditions (5, 10, 15, 17). A distinguishing feature of the incubations described here is that groundwater and sediment were collected simultaneously using a device coined the needlesampler (18), placed immediately under anaerobic conditions in the field, rapidly transported to the laboratory, and maintained under such conditions for the 11-month duration of the experiment.

## Materials and Methods

**Site Description.** The sampling site ( $23.8^\circ \text{N}$ ,  $90.6^\circ \text{E}$ ) is at the center of an expanse of rice paddies in Araihazar upazila, 30 km east of Dhaka. Grey aquifer sediments extend to a depth of 48 m at this location. The radiocarbon age of two wood fragments recovered during drilling of the site in January 2005 ( $6970 \pm 45$  and  $7180 \pm 55$   $^{14}\text{C}$  yr at 40 and 44 m depth, respectively) confirm that the aquifer is of Holocene age (19). The generally sandy section of gray material is interrupted by a clay layer between 27 and 34 m depth

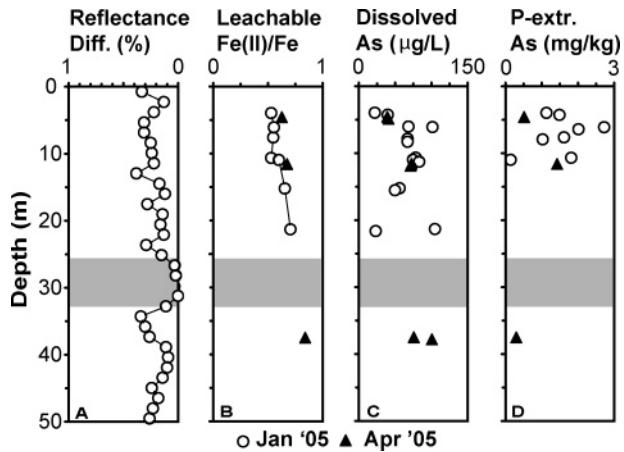
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**FIGURE 1.** Profiles of sediment and groundwater at the study site in Araihazar, Bangladesh: (a) difference in diffuse spectral reflectance at 530 and 520 nm measured on-site for cuttings recovered during drilling; (b) proportion of Fe(II) in the HCl-leachable Fe fraction of sediment samples; (c) dissolved As concentrations in groundwater; (d) concentration of P-extractable As in the sediment. With the exception of the cuttings used for reflectance measurements in January 2005, all sediment and groundwater samples were obtained with the needle-sampler in January (○) and April (▲) 2005. The gray shading indicates the extent of a thick clay layer encountered at the site.

(Figure 1). Numerous mechanized irrigation wells are distributed across the surrounding rice paddies and draw water through screens that span the entire shallow gray aquifer (20).

A set of monitoring wells (site F in ref 21) located 1 km northwest of the sampling site shows a gradual increase in As concentrations in groundwater of the area from <10 µg L<sup>-1</sup> at 6 m depth to ~50 µg L<sup>-1</sup> at 15 m depth (21). Over the same depth range, groundwater dating using the <sup>3</sup>H-<sup>3</sup>He technique indicates an increase in the age of groundwater from <1 to ~5 years at the same location (22). Between 15 and 20 m depth at site F, the concentration of As increases rapidly to 200 µg L<sup>-1</sup> and the age of groundwater increases by several decades to about 30 years.

**Sample Collection.** A preliminary survey of the area was conducted in January 2005. A single well was drilled using the traditional “sludger” method and a series of groundwater and sediment samples were collected to a depth of 21 m using the needle-sampler (18). The diffuse spectral reflectance of the cuttings collected every 1.5 m was measured minutes after collection in the field to indicate the Fe speciation in the solid phase (21).

The samples used for the long-term incubations were collected in April 2005 from a new well within 2 m of the first profile. The needle sampler was used to obtain slurry samples from three depths: 5, 12, and 38 m. The starting material from each depth consisted of groundwater and sediment contained in two needle sampler tubes collected within 0.3 m of each other vertically. Preferential uptake of groundwater relative to sediment with the needle-sampler increased the porosity of the collected slurries. The porosity of sandy aquifers typically varies between 0.2 and 0.3, while the material collected by the needle sampler had a porosity ranging from 0.7 to 0.8. The collected sediment may also have been biased toward the finer fraction because the coarsest particles are excluded by the entry holes of the needle, which are less than 2 mm in diameter (18).

Before sampling, each needle-sampler tube was flushed with N<sub>2</sub> and then evacuated to <300 mbar. To minimize O<sub>2</sub> exposure, the headspace of each tube was also flushed with N<sub>2</sub> immediately after collection. Aliquots of the collected

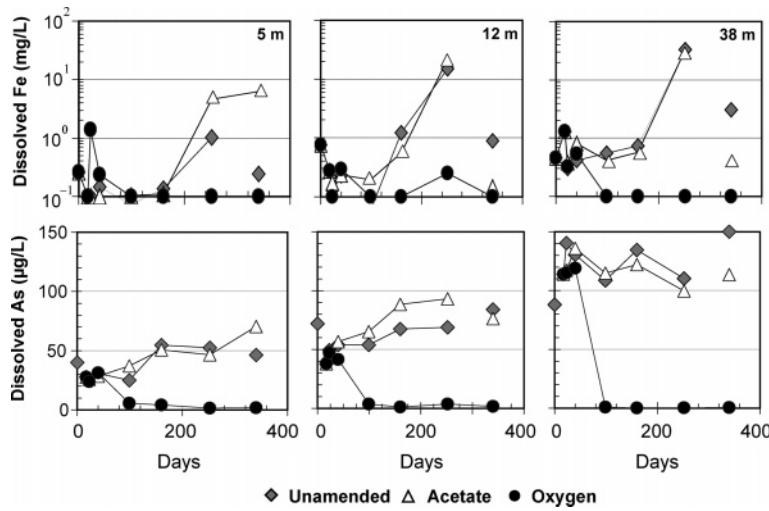
groundwater were gently pressure-filtered in the field, also under N<sub>2</sub>. The needle-sampler tubes were then stored in plastic, N<sub>2</sub>-flushed, airtight anaerobic boxes (Becton-Dickinson) containing pouches that generate H<sub>2</sub> and consume O<sub>2</sub> with a catalyst. Three days after collection, the needle-sampler tubes were transferred to an anaerobic chamber (Coy Laboratories) where they remained for the duration of the experiment. The chamber was kept under a N<sub>2</sub> atmosphere containing 1–2% H<sub>2</sub>.

**Amendments.** The two needle sample slurries of roughly equal proportion from each depth were combined, homogenized, and divided into five 60-mL amber glass bottles in the anaerobic chamber 17 days after collection (Supporting Information). Each microcosm bottle contained between 8 and 15 mL of groundwater and between 5 and 11 g of sediment. The amendments could not be replicated because of the limited quantities of sediment and groundwater that were collected. The sample bottles were sealed with a butyl rubber stopper and an aluminum crimp. The bottles were gently agitated over the next 11 months and kept at room temperature. To minimize the potential effect of H<sub>2</sub> contained in the anaerobic chamber on the incubations, the bottles were flushed with ultrahigh purity N<sub>2</sub> for several minutes after every sampling. The porosity of the different slurries was gradually reduced from an initial range of 0.6–0.9 to a final 0.4–0.8 by sampling water seven times over the course of the experiment.

The slurries from each depth were treated in three different ways: without amendment, with the addition of acetate, and with periodic, small oxygen additions, hereafter referred to as unamended, acetate, and oxygen incubations. The acetate addition (14 µmole per sample, which corresponds to ~2 µmole g<sup>-1</sup> sediment or ~1.5 µmole L<sup>-1</sup> groundwater) supplied enough electron donors to potentially reduce approximately 10% of the acid-leachable Fe present in the sediment. The oxygen additions occurred at each sampling point by adding approximately 1.2 mL of air to the 60 mL bottles (i.e., 10.3 µmole of O<sub>2</sub>, equivalent to ~0.21 mg O<sub>2</sub> L<sup>-1</sup> groundwater). The oxygen addition could in principle oxidize 18 µg of Fe after each sampling, which would be sufficient to remove more than 1.5 mg L<sup>-1</sup> Fe from solution. In addition, two amendments (autoclaving and addition of antibiotics) were prepared in an attempt to inhibit microbial activity and distinguish between biotic and abiotic processes. The results are not reported as autoclaving dramatically altered the sediment geochemistry and the antibiotic mix was not effective in reducing microbial activity.

**Analyses.** Groundwater samples collected from each incubation on days 17, 23, 40, 99, 160, 252, and 340 provide a detailed time series of variations in the concentrations of dissolved constituents, including As, Fe, Mn, S, P and the major cations, Na, K, Mg, and Ca (Supporting Information). Aliquots of 1.2 mL were drawn from the slurries, filtered through 0.45 µm syringe filters, acidified to 1% HCl (Optima), and analyzed by high-resolution inductively coupled plasma mass spectrometry (HR ICP-MS) (23).

Aquifer sediment was analyzed on day 17 after collection, when the slurries were divided into amendments, and again on day 340 at the completion of the experiment. Two extractions were used to assess the availability of As in the solid phase. A 1 M phosphate solution was used to estimate the amount of adsorbed As (24, 25). A hot 1.2 M HCl solution was used to release As more tightly bound within amorphous Fe minerals (21). The extracts were analyzed for As using HR ICP-MS (23). Iron speciation of the HCl leachates were quantified on day 17 and day 340 using the ferrozine method (21). The abundance of organic substrate in each amendment was not assessed.



**FIGURE 2.** Variations in dissolved Fe and As concentrations over the course of the 11 month long incubations of sediment and groundwater slurries collected in Bangladesh at three different depths in April 2005. Note the logarithmic scale for dissolved Fe. Disconnected symbols indicate samples exhibiting a large drop in dissolved Fe at the completion of the incubations.

## Results

**Site Profile.** Dissolved As concentrations generally increased with depth from 20 to  $100 \mu\text{g L}^{-1}$  in gray aquifers at the study site, although there is considerable variability between some closely spaced samples (Figure 1). Detectable levels of dissolved Fe and Mn indicate that groundwater was suboxic over the entire depth range (Supporting Information). The profile of sediment reflectance documents the increasingly reduced state of the aquifer as a function of depth and is consistent with a rise from 0.6 to 0.8 in the proportion of Fe(II) to total Fe in the sediment leachates. The concentration of P-extractable As in the sediment was highly variable in the upper 15 m ( $0.1\text{--}2.7 \text{ mg kg}^{-1}$ ). Two samples from three depths were used for the incubations. Differences in As concentrations between samples collected 0.3 m apart were small and initially averaged 40, 72, and  $88 \mu\text{g L}^{-1}$  in the dissolved phase and 0.5, 1.4, and  $0.3 \text{ mg kg}^{-1}$  P-extractable As in the sediment at 5, 12, and 38 m depth, respectively. Dissolved Fe concentrations were more variable for samples collected 0.3 m apart and ranged from 0.1 to  $0.9 \text{ mg L}^{-1}$  (Supporting Information).

**Iron.** Concentrations of dissolved Fe declined from an average of 0.3 to  $0.1 \text{ mg L}^{-1}$  and 0.8 to  $0.3 \text{ mg L}^{-1}$  at 5 and 12 m depth, respectively, between the time of collection and day 17 when slurries from nearby depths were combined in the laboratory (Supporting Information). Over the same period, dissolved Fe concentrations in material from 38 m depth increased from 0.5 to  $1.3 \text{ mg L}^{-1}$ . These early variations are small compared to subsequent 10–85-fold increases in dissolved Fe concentrations for all unamended and acetate incubations until day 252, after which an almost equally large loss of Fe was observed on day 340 in the majority of samples (Figure 2). Over 90% of the total Fe released occurred after day 99 of the incubations. The proportion of Fe (II) to total Fe in the acid leachates remained nearly unchanged for the anoxic samples (with the exception of the unamended sample from 12 m depth for which the Fe(II)/Fe ratio decreased from 0.7 to 0.5). After day 40, the oxygen addition maintained dissolved Fe concentrations  $0.1 \text{ mg L}^{-1}$  at all three depths (with exception for the sample on day 252 at 12 m that measured  $0.25 \text{ mg L}^{-1}$ ). The leachable Fe(II)/Fe ratio in the sediment decreased from 0.6–0.8 to 0.4–0.5 in the incubations amended with oxygen.

**Arsenic.** From the time of collection to day 17, the concentration of dissolved As declined from 40 to  $28 \mu\text{g L}^{-1}$  and from 72 to  $38 \mu\text{g L}^{-1}$  in the slurries from 5 and 12 m depth

and increased from 88 to  $114 \mu\text{g L}^{-1}$  at 38 m depth. Dissolved As then gradually increased to  $50\text{--}90 \mu\text{g L}^{-1}$  in the unamended and acetate samples from 5 and 12 m depth (Figure 2). At 38 m, the amended and acetate samples showed no systematic trend or slightly decreased. The loss of dissolved Fe on day 340 was accompanied by a decrease in dissolved As in the unamended incubation from 5 m and in the acetate incubation from 12 m. There was no systematic difference in As concentrations between the unamended and acetate incubations at 5 m, while at 12 m the acetate amendment roughly doubled the amount of As released. After day 99 of incubation, the oxygen additions maintained dissolved As concentrations  $<6 \mu\text{g L}^{-1}$  at all depths.

The concentration of P-extractable As in the sediment was determined on day 17 and 340; HCl-extractable As was determined only on day 340 (Supporting Information). At 5 m, the P-extractable As concentration remained steady at  $0.5 \text{ mg kg}^{-1}$ . Samples from 12 m had the highest level of P-extractable As of  $1.4 \text{ mg kg}^{-1}$ , which decreased to about  $0.6 \text{ mg kg}^{-1}$  at the end of the experiment. At 38 m, the P-extractable fraction decreased slightly from 0.30 to 0.24  $\text{mg kg}^{-1}$  between day 17 and 340. The observed decreases in P-extractable As concentrations in the solid phase cannot be explained by the measured releases of As to groundwater because the releases are equivalent to only 2–6% of the total present in P-extractable form. The HCl-extractable As concentration in the solid phase at the end of the experiment was comparable at the three depths ( $1.3\text{--}1.5 \text{ mg kg}^{-1}$ ).

**Sulfur.** Dissolved S concentrations in the unamended and acetate incubations show no discernible trend. In contrast, dissolved S increased in the oxygen incubations to maximum concentrations of 2.2, 3.8, and  $53.5 \text{ mg L}^{-1}$  in the 5, 12, and 38 m samples, respectively (Supporting Information).

## Discussion

**Iron Release and Precipitation.** Few, if any, studies have documented the release of Fe from natural aquifer solids to actual groundwater under anoxic conditions and over an extended period of time. In comparison to subsequent changes, the variations in dissolved Fe concentrations in slurries between the time of collection and the onset of the actual incubations on day 17 day were modest, suggesting that the system was not excessively perturbed by sampling, transport, and manipulation. By day 99, oxygen additions had already drastically decreased dissolved Fe concentrations to  $<0.1 \text{ mg L}^{-1}$  due to oxidation and precipitation of Fe(II)

initially present in the groundwater. The higher dissolved Fe concentrations maintained throughout for other incubations suggest that anoxic conditions were successfully maintained in the absence of oxygen additions.

It is likely that the dilution of aquifer particles in ambient groundwater increased the exposure of surface sites available for adsorption, and this may be the reason for the 30% reduction in dissolved Fe observed at 5 and 12 m depth between days 0 and 17. The increase in dissolved Fe at 38 m depth cannot be similarly explained, but there are other indications that the geochemistry of the aquifer material was different at that depth. The increase in dissolved S (most likely in the form of sulfate) concentrations observed in all oxygen incubations was much greater at 38 m depth than at 5 or 12 m (Supporting Information). This suggests that aquifer material from 38 m depth, and not in the shallow material, contained a considerable proportion of reduced and labile S, which has been observed in Bangladesh aquifer material (14). A simple calculation shows that labile S could be bound to an equivalent of 0.8% of the total acid-leachable Fe at 38 m depth.

The large release of Fe observed in the six incubations under anaerobic conditions between days 99 and 252 suggests a significant departure from initial conditions in the aquifer. The enhanced release may reflect that new and/or more Fe(III) surface sites became available for microbial reduction because of dilution of initially more tightly packed particles, as well as a potential thinning of the diffuse-boundary layer caused by gentle agitation of the slurries. Laboratory studies using model solids have shown that Fe(II) is not necessarily released during the early stages of microbial Fe(III) reduction, but can instead adsorb onto the oxide surface or form secondary minerals (9, 26–28). Readsorption of Fe(II) on newly available sites, until they became saturated is, therefore, a plausible explanation for the observed delay in the large increase in dissolved Fe(II) concentrations until after day 160 of the incubations (Figure 2).

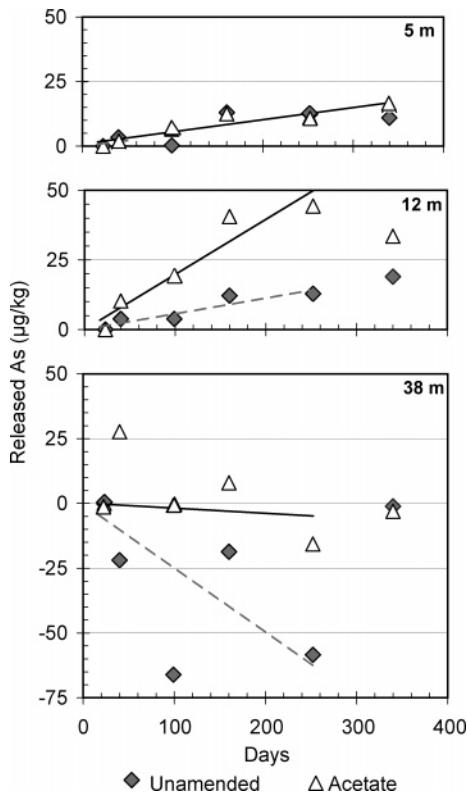
The maximum Fe concentrations reached during the anoxic incubations are higher than typically observed in Bangladesh groundwater, but similar levels have been observed in Vietnam (29). The rise in Fe concentrations corresponds to a release of 0.2–50 mg Fe kg<sup>-1</sup> of sediment and corresponds to only 0.01–0.8% of HCl-extractable Fe. The limited impact of the acetate amendment suggests that organic substrate availability does not limit microbial Fe reduction in these shallow aquifers (30). We cannot exclude the possibility that the observed Fe releases were driven by abiotic desorption rather than microbial Fe reduction (5, 12, 28, 31–33). In that case, however, a gradual release over the entire course of the experiment toward a new steady-state might have been expected instead of a delay followed by an increasing release of Fe.

The large drop in dissolved Fe concentrations for 5 out of 6 anoxic incubations between days 252 and 340 suggests that saturation with respect to a solid-phase had been reached (27). Geochemical calculations were performed using Geochemist's Workbench (Rockware, Inc.) with the Lawrence Livermore National Laboratories combined database with published alterations (34, 35). Analysis of Fe precipitation by day 340 was conducted using concentrations of all elements measured in the unamended samples at day 252 (Supporting Information). Solubility calculations and a concurrent 2–40 µg L<sup>-1</sup> reduction in dissolved P concentrations suggest the formation of Fe-sulfides and Fe-phosphate minerals, such as pyrite and vivianite. It is noteworthy that the decline in dissolved As in response to significant Fe precipitation was modest, possibly due to competition with other groundwater constituents for binding sites, particularly on Fe sulfide minerals (36), and limited coprecipitation.

**Initial As Adsorption.** In the case of As, we use a formulation equivalent to a Langmuir isotherm to estimate the potential effect of increasing the availability of surface site due to dilution of the particles in the slurries upon collection. Adsorption of As is assumed to be controlled by a simple mass-action law of the form:  $S + A = SA$  and an adsorption coefficient  $K_{ads} = [SA]/([S][A])$ , where  $S$  represents the concentration of surface sites in the slurry and  $A$  is the concentration of dissolved As. The concentration of dissolved As ( $A$ ) was measured in the sample filtered in the field right after collection and 17 days later; the concentration of P-extractable As is used as a proxy for adsorbed As ( $SA$ ). For each of the three depths, the change in available surface sites,  $S$ , was calculated assuming equilibrium at collection and that the change in dissolved As from day 0 to day 17 was entirely due to adsorption (Supporting Information). Using this model, the decline in As concentrations observed in slurries from 5 and 12 m depth could be explained entirely by a 50 and 90% increase, respectively, in surface sites (as in the case of Fe, no effect on dilution was observed in slurries from 38 m depth for As).

**Implications of Observed Arsenic Release.** The noticeable impact of oxygen additions on the incubations of aquifer material from all three depths confirms that the formation of fresh Fe(III) adsorption sites is sufficient to significantly limit the release of As. Anoxia, therefore, appears to be a prerequisite for accumulating As in groundwater of Bangladesh. This does not mean that anoxia alone is sufficient to cause As mobilization or that releases of As and Fe to groundwater are necessarily coupled. Indeed, the time course of the anoxic incubations provides new evidence of decoupling between As and Fe mobilization. The release of As at 5 and 12 m was evident by day 40, while Fe release did not begin until day 99 and rapidly increased after day 160. The lack of Fe release could be due to Fe adsorption and secondary mineral formation, as described above. The large release of Fe to groundwater by day 252, however, did not cause a concomitant increase in As concentrations. This decoupling could be related to the formation of various secondary Fe minerals, including vivianite and magnetite, which can incorporate As into the new minerals (17, 28). Alternatively As may not be distributed uniformly within Fe oxyhydroxide surfaces (10). In either case, the observations suggest that the processes regulating As and Fe release may have little in common once an aquifer becomes anoxic.

Average rates of As mobilization were calculated for each anoxic incubation. The rate of As release for each sample was calculated beginning with day 23, 6 days after amendment. The entire 340 days were considered for the acetate incubation at 5 m. Only the releases until day 252 were included for the unamended incubation at the 5 m depth and both anoxic incubations from 12 and 38 m due to the large loss of Fe by this time and the uncertain influence this had on As release (Figure 3). The amount of As released was calculated by determining the change in mass of dissolved As between each sampling and normalized this to the total mass of sediment. Rates of release were determined by least-squares regression; the uncertainty is listed as two times the standard deviation ( $\sigma$ ) of the slope. The combined rate of As release was  $21 \pm 4 \mu\text{g As kg}^{-1}$  sediment per year ( $r^2 = 0.91$ ,  $n = 9$ ) for the two anoxic incubations from 5 m depth; the calculated rates were not statistically different between the types of amendment. At 12 m, the acetate addition increased the rate of As release nearly 4-fold from  $23 \pm 6 \mu\text{g As kg}^{-1} \text{yr}^{-1}$  ( $r^2 = 0.93$ ,  $n = 4$ ) to  $82 \pm 18 \mu\text{g kg}^{-1} \text{yr}^{-1}$  ( $r^2 = 0.95$ ,  $n = 4$ ) for the unamended and acetate incubations, respectively. The 5 m samples may have a higher natural abundance of organic substrate, due to their younger age and closer proximity to surface organic inputs, thus rendering the acetate addition superfluous (14, 37). The combined set of observa-



**FIGURE 3.** Amount of As released to groundwater for anoxic incubations normalized to the quantity of sediment in each slurry of Bangladesh aquifer material. The rates of release were calculated by least-squares regression fit over the time span, as indicated by the regression lines. The regression for the 5 m depth combined the unamended and acetate incubations, while the regressions were calculated separately at 12 and 38 m.

tions indicates that the process of As mobilization in Bangladesh is potentially microbially mediated, although not necessarily limited by the availability of organic substrate.

The incubations unambiguously demonstrate that As is not necessarily released at higher rates in shallow gray aquifers of Bangladesh where dissolved As concentrations are elevated. Whereas groundwater As concentrations increased with depth at the sampling site (Figure 1), As release was only observed in aquifer material from 5 and 12 m depth. The sample from 38 m had the highest initial As concentrations in groundwater and yet no release was observed, although the release of Fe was comparable at all depths. The low level of P-extractable As contained in aquifer material from 38 m suggests that As release was limited by the lack of a sufficient pool of mobilizable As in the solid phase. Elevated groundwater As concentrations observed at 38 m and the absence of As release, therefore, may indicate transport of As released at shallower depths, as suggested previously (5, 38). Readsorption of As released elsewhere at 38 m depth may also have been inhibited by the lower adsorption capacity of Fe-sulfide minerals, whose presence is suggested by the effect of the oxygen amendment (36). According to the rates of As release measured at 5 and 12 m depth, an initial pool of  $\sim 10 \text{ mg kg}^{-1}$  mobilizable As (2, 39) could be lost from an aquifer due to groundwater flow in as little as 1000 years assuming no replenishment of transported As. Such a flushing mechanism has previously been invoked to explain low groundwater As concentrations in older gray sediment (19, 37).

**Reinterpreting Spatial Patterns of As in Groundwater.** Estimating the increase in dissolved As in the aquifer corresponding to the measured release rates requires con-

sideration of the increased surface site availability as well as the increased porosity of the incubations. If we assume As mobilization is related to surface site availability, greater surface exposure to the groundwater should lead to more As release. We postulate here that adsorption and release are affected similarly by changes in surface site availability, and therefore, we assume that the measured rates of As release overestimate the rate of release in the aquifer by 50–90% (depending on depth), as determined from the observed As adsorption that occurred upon sample collection. This is partially offset by the lower porosity of aquifers which causes the release (as measured in As released per weight of sediment) to have a greater effect on dissolved As concentrations than observed in the incubations. Considering these two competing effects, the observed release rates of As would be equivalent to increases in dissolved As of  $110 \mu\text{g L}^{-1}$  per year at 5 m and  $90 \mu\text{g L}^{-1}$  per year at 12 m.

The observed rates suggest a relatively constant release of As from aquifer sediments; however, this does not imply that dissolved As concentrations are necessarily increasing in the aquifer over time. Indeed, a recent study of a large number of wells in Araihazar has shown that dissolved As concentrations remained relatively constant over several years (40). Steady-state concentrations could reflect a balance between the release of As from gray aquifer solids and continuous replacement of groundwater with surface water that is initially low in As (22). The short time scale of As release does suggest that the distribution of As in shallow aquifers could change over time, particularly in a region like the present study area where irrigation pumping probably drastically affects flow patterns (20).

A recent study in this region of Bangladesh provides evidence that the observed rise of As concentrations in groundwater over time indicated by these long-term incubations is realistic. Shallow groundwater was dated by the  $^3\text{H}$ - $^3\text{He}$  technique at a set of six nests of monitoring wells in the same general area of Araihazar (22). Whereas the age of groundwater increased with depth at all six sites, the rate of increase varied considerably from one site to the other, most likely in response to variations in local recharge. Comparison of groundwater ages ranging from <1 to over 30 years with groundwater As concentrations of  $1\text{--}520 \mu\text{g L}^{-1}$  measured in the same set of wells, however, results in a relationship that is fairly consistent across all sites and corresponds to a release rate of  $19 \pm 2 \mu\text{g L}^{-1}$  As per year. This entirely independent measure of the rate of As accumulation in groundwater is within a factor of 5 of the rates measured in the unamended incubations, after correction for the estimated effect of increased exposure of surface sites.

Our incubations of simultaneously collected Bangladesh sediment and groundwater over an extended period provide new evidence that the concentration of dissolved Fe in groundwater is not necessarily linked to the rate of reductive dissolution of Fe oxyhydroxides. This may be one reason why there is no simple relationship between concentrations of Fe and As in anoxic groundwater (2, 5, 10, 21, 41). The limited impact of acetate on incubations of natural aquifer material suggests that the release of both Fe and As is not necessarily limited by the supply of organic substrate. Flow patterns and groundwater age may, therefore, play a dominant role in setting the highly variable distribution of As in shallow groundwater of Bangladesh.

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## Supporting Information Available

Dissolved concentrations of Fe, As, Mn, S, P and the major cations (Na, Mg, K, and Ca) are reported for both the site profiles and the incubations. Fe and As sediment concentrations from the HCl and phosphate extractions are given. The values used for the Langmuir calculations of increased surface site availability are reported. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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