Diversity and Ecophysiology of New Isolates of Extremely Acidophilic CS₂-Converting Acidithiobacillus Strains


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Biofiltration of industrial carbon disulfide (CS₂)-contaminated waste air streams results in the acidification of biofilters and therefore reduced performance, high water use, and increased costs. To address these issues, we isolated 16 extremely acidophilic CS₂-converting Acidithiobacillus thiooxidans strains that tolerated up to 6% (vol/vol) sulfuric acid. The ecophysiological properties of five selected strains (2Bp, Sts 4-3, S1p, G8, and BBW1) were compared. These five strains had pH optima between 1 (2Bp) and 2 (S1p). Their affinities for CS₂ ranged between 80 (G8) and 130 (2Bp) μM. Strains S1p, G8, and BBW1 had more hydrophobic cell surfaces and produced less extracellular polymeric substance than did strains 2Bp and Sts 4-3. All five strains converted about 80% of the S added as CS₂ to S⁰ when CS₂ was supplied in excess. The rate of S⁰ consumption varied between 7 (Sts 4-3) and 63 (S1p) nmol O₂ min⁻¹ ml culture⁻¹. Low S⁰ consumption rates correlated partly with low levels of cell attachment to externally produced S⁰ globules. During chemostat growth, the relative amount of CS₂ hydrolyase in the cell increased with decreasing growth rates. This resulted in more S⁰ accumulation during CS₂ overloads at low growth rates. Intermittent interruptions of the CS₂ supply affected all five strains. Strains S1p, G8, and BBW1 recovered from 24 h of starvation within 4 h, and strains 2Bp and Sts 4-3 recovered within 24 h after CS₂ was resupplied. We recommend the use of mixtures of Acidithiobacillus strains in industrial biofilters.

Carbon disulfide (CS₂) is a toxic, volatile, flammable, and explosive solvent widely used in, e.g., the viscose rayon industry (1). Because of its toxicity and the increasingly stringent rules governing the emission of harmful gases, it is necessary to treat CS₂-containing industrial waste gases. Biological treatment of CS₂ (and hydrogen sulfide, H₂S) with sulfur-oxidizing bacteria provides an attractive alternative to conventional treatment systems (e.g., active carbon, incineration, caustic scrubbing) (2, 3). Typical concentrations of CS₂ in contaminated air from viscose industries are around 4 to 20 nmol ml⁻¹ (100 to 500 ppm) (2).

The number of microorganisms known to be able to grow chemolithoautotrophically on CS₂ is limited to some Thiobacillus species, Thiothrix ramosa, Paracoccus denitrificans, and a Thiomonas sp. (4–8). All of these CS₂-utilizing bacteria grow at neutral pH. Thus far, only one CS₂-utilizing species (Thiobacillus sp. strain Tj330, DSM8985) capable of growth under acidic conditions (as low as pH 0.5) has been described (9). The only reported screening of 10 (Acidithiobacillus) strains showed that CS₂ conversion is not a general trait of (acid)thiobacilli (4). In that screening, only one strain, Thiobacillus thioparus TK-m, was found to be capable of CS₂ conversion. We recently discovered that CS₂ conversion is not limited to the domain Bacteria; the hyperthermophilic archaea Acididans sp. strain A1-3 and Sulfolobus solfataricus P2 can also grow on CS₂ as a main carbon and energy source (10). However, these archaea are not able to grow at the extremely low pH values that acidithiobacilli can cope with.

CS₂-converting sulfur oxidizers in operating biofilters are acidophilic bacteria (2, 11–13). They convert CS₂ via the two-hydrolysis reaction steps CS₂ + H₂O → COS + H₂S and COS + H₂O → CO₂ + H₂S and obtain their energy from the oxidation of H₂S via S⁰ and SO₂⁻ to SO₄²⁻ as follows: H₂S + SO₂⁻ → SO₄²⁻ + 2H⁺ (14). Therefore, an inherent result of CS₂ conversion is acidification of the biofilters, which can be limited only by flushing the trickling filters with fresh water. Operating at a pH as low as possible will considerably reduce the volume of fresh water used for neutralization. Water use would be further reduced if the H₂SO₄ produced could be reused in the viscose-rayon industry. This becomes economically feasible when the H₂SO₄ concentration in the reactor effluent is at least 10% (wt/vol) (5.6% [vol/vol]). However, the performance of biotrickling filters is compromised by severe acidification and buildup of elemental sulfur (S⁰) that can clog the filters. Therefore, we set out to isolate new CS₂-converting bacterial strains able to tolerate extremely low pH values with variable CS₂ loads without loss of CS₂ conversion efficiency and without the production of large amounts of elemental sulfur.

MATERIALS AND METHODS

Media and culture conditions. Strains were enriched and cultured in basal salt mineral medium (MM) with CS₂ as the sole carbon and energy source as described previously (15). Bacteria were grown at room temperature (RT, 22°C) in 120-ml bottles containing 20 ml MM acidified with sulfuric acid. Alternatively, bacteria were grown on MM plates solidified with 1% (vol/vol) Gelrite (16) and acidified with 0.1% (vol/vol) sulfuric acid. This was the maximum [H₂SO₄] at which plates could still be poured without the Gelrite solution immediately solidifying when brought into contact with the H₂SO₄-containing MM solution. In the case of Gelrite, H₂SO₄ increases solidification while the opposite occurs with agar(ose). Plates were incubated in an airtight jar. Unless stated differently, sulfuric acid concentrations are reported as percentages (vol/vol) (1% [vol/vol] is 1 July 2013 Accepted 25 August 2013
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equivalent to 1.8% [wt/vol] and 0.18 M sulfuric acid). Strains were also grown in minichemostat reactors as described previously (15).

The headspaces of the bottles and jars used were continuously flushed with a CS$_2$-containing air stream from a purpose-built distribution system (see the supplemental material).

**Enrichment and isolation.** Volumes of 0.5 to 1 ml of environmental or industrial samples were inoculated into 120-ml serum bottles with 20 ml acidified MM. The initial H$_2$SO$_4$ concentration was 0.5 to 1% (vol/vol) in the environmental samples and 2% in the samples from the biotrickling filters. CS$_2$ was supplied as the sole energy source via the distribution system. When a visually dense culture was observed, the enrichment was transferred to fresh MM. The maximum H$_2$SO$_4$ concentration at which growth occurred was determined by the subsequent transfer of enrichments to MM with higher H$_2$SO$_4$ concentrations. Pure cultures were obtained from enrichment cultures grown in 4 to 6% H$_2$SO$_4$ on Gelrite plates. Single colonies were serially transferred three times to fresh plates and checked microscopically for purity. Fungal contamination (present mainly in the enrichments from industrial samples) was eliminated by adding 150 µg ml$^{-1}$ chlorothalonil either to the plates or to liquid cultures. The H$_2$SO$_4$ tolerance of each isolated strain was confirmed by repeated subculturing at least three times in liquid MM containing 4 to 5% H$_2$SO$_4$.

**Screening of known Acidithiobacillus strains for CS$_2$ conversion capacity.** Five Acidithiobacillus strains were obtained from the Deutsche Sammlung von Mikroorganismen und Zellkulturen (DSMZ) and grown in batch cultures in the media suggested (DSMZ medium number in parentheses after the strain name) or as otherwise stated, i.e., A. ferrooxidans DSM41882 (882), A. caldus DSM8584 (150a), A. thiooxidans DSM14887 (71), A. thiobacillus DSM504 (MM plus 10 g liter$^{-1}$ sterile S$^0$), and A. albertensis DSM14366 (71). After growth for 1 week, the headspace of the cultures was supplemented with CS$_2$ (20 to 30 nmol ml$^{-1}$). The CS$_2$ concentration and the presence of intermediates (H$_2$S and COS) in the headspace were monitored over time by gas chromatography (17).

**PCR, cloning, and sequencing.** DNA was isolated from each strain by phenol extraction (18). The 16S rRNA gene and the 16S-23S intergenic spacer region (ISR) of 16 isolated CS$_2$-hydrolyzing bacterial strains were amplified by hot-start PCR with the Go Taq Green buffer system (Fermentas) with 2.5 mM MgCl$_2$, 0.2 mM deoxynucleoside triphosphates, 1 µl bovine serum albumin, 0.4 µM each primer (see Table S1 in the supplemental material), and 1 µl Taq polymerase (Fermentas). The PCR protocol consisted of 2 min at 95°C, 30 cycles of 1 min at 95°C, 1 min at 54 to 65°C, and 2 min at 72°C; and a final elongation step of 10 min at 72°C. PCR products were ligated into pGEM-T Easy (Promega) and transformed into Escherichia coli strain TOP10 (Invitrogen) according to the manufacturer’s instructions. Plasmids with a correct insert were Sanger sequenced by the sequencing facility at the Department of Human Genetics, University of Groningen. Sequences of the ribosomal operon of A. thiobacillus ATCC 19377 and A. caldus ATCC 51756 were kindly made available by Jorge Valdés and David Coble, CBGB, Santiago, Chile (21).

**Cryo-SEM.** Cryo-scanning electron microscopy (cryo-SEM) was used to study the morphology of colonies growing on 1% Gelrite plates acidified with 0.018 M H$_2$SO$_4$ by a method similar to that described in reference 22. The morphology of colonies was observed on electrocoative aqueous colloidal graphite (DAG; Agar Scientific) on a mounting stub, and quickly frozen by submersion in degassed liquid N$_2$ (−196°C). While under vacuum, the sample was transferred to the Gatan cryotransfer box. Sections through colonies were made with a razor blade at this stage when required. The temperature was subsequently increased to −100°C to sublime off water that had settled on top of the specimen by condensation for a period of 5 min. When all of the surface water had been removed, the temperature was reduced to between −100 and −150°C. The sample was then sputter coated for 45 s with a mixture of 60% gold and 40% palladium and transferred to a JEOL 6330F scanning electron microscope.

**Yield determinations.** For total carbon measurements, 3- to 4-ml reactor samples were centrifuged and the pellets were resuspended in 2 ml 1 M HCl, pH 3. The washed cells were dried overnight under vacuum at 70°C. The C/N ratio of the dried material (0.3 to 0.4 mg) was determined by elemental analysis with a Thermo Fisher Scientific EA 1110 CHN element analyzer coupled to a Finnigan DELTA plus mass spectrometer. For protein determination, 2-ml reactor samples were centrifuged at 4°C for 30 min at 18,000 × g. The pellets were resuspended in 0.5 ml 1 M NaOH, boiled for 5 min, and neutralized with 0.5 ml 1 M HCl. Alternatively, 0.5 ml 1 M NaOH was added directly to 200-µl reactor samples, and after boiling, the mixture was neutralized with 0.3 ml 1 M HCl. Protein concentrations were determined with the Bio-Rad protein assay kit according to the manufacturer’s instructions. Culture density, determined by both cell density and the presence of S$^0$, was measured spectrophotometrically as the optical density at 600 nm (OD$_{600}$).

**Determination of pH optima.** pH optima were determined by a floating-filter method. Samples taken from the steady-state chemostats (1% H$_2$SO$_4$ measured pH, 0.72) were diluted 10$^6$-fold in MM at pH 2, and 1 ml of this dilution was filtered through a sterile, 0.2-µm, 25- or 47-mm diameter Cyclopore polycarbonate filter (Whatman). Filters were then floated on 20 ml MM acidified with sulfuric acid to pHs of 0.5 to 6. They were placed in airtight jars and incubated at RT for 16 days with a continuous flow of 45 ml min$^{-1}$ air containing 10 mmol CS$_2$ ml$^{-1}$. Growth at different pHs was determined by measuring colony diameters and counting colonies. The growth of strain BBW1 was quantified visually, as cells had spread over the filter and over the surface of the medium.

**Cell surface hydrophobicity.** Cell surface hydrophobicity was determined by a modified form of the method of Rosenberg et al. (23). The pH of culture samples from the minichemostats was adjusted to 3 or 7 with PUM buffer (23). The suspensions were diluted to an OD$_{600}$ of 0.45 (A$_{600}$) in PUM buffer with the appropriate pH. In a test tube, a 1-ml suspension was mixed with 200 µl n-octane or n-hexane. The mixture was incubated at RT for 10 min, mixed vigorously for 1 min, and left to stand at RT for at least 25 min. OD$_{600}$ was measured (A$_{600}$), and the percent adherence to the solvent was calculated with the equation $(1 - A_{600})/A_{600} 	imes 100$. The use of OD$_{600}$ to measure suspension turbidity yielded similar results.

**Preparation of cell extracts.** Cell extracts from steady-state reactor-grown bacterial cells were prepared as follows. Thirty to 50 ml was removed from the reactors and centrifuged at 4°C for 30 min at 12,000 × g. The cell pellets were washed with 15 ml sterile distilled H$_2$O and resuspended in 0.5 ml 20 mM KP, at pH 7. Approximately 350 µM glass beads (80- to 110-µm size) were added, and the cells were broken by bead beating for 2 × 2 min at 30 Hz (Retsch) with intermittent cooling on ice. The broken cell mixtures were centrifuged for 5 min at 16,000 × g, and the supernatants were stored at −20°C with a final concentration of 10% glycerol.

**Enzyme kinetics based on H$_2$S measurements.** The Michaelis-Menten constants $K_m$ and $V_{max}$ were determined for the CS$_2$ conversion rates of each cell extract of the different strains by measuring the H$_2$S production rate with an H$_2$S microsensor (Unisense) in 20 mM HEPES (pH 7) as described previously (15). Experiments were performed at pH 7, as the CS$_2$ hydrolyase is predicted to reside in the cytoplasm of the cell because of the absence of a signal sequence at the N terminus of the enzyme. The $K_m$ and $V_{max}$ values were calculated from Michaelis-Menten plots by nonlinear regression with the Michaelis-Menten equation $V = V_{max} \times [S]/(K_m + [S])$. Experiments were repeated at least three times, and average values of three independent experiments ± the standard errors of the means were calculated. The method was used to determine $V_{max}$ values (with 600 µM CS$_2$ as the substrate) for steady-state cells from the genome.
minichemostats \((D = 0.02)\) diluted 200× in reactor medium (MM containing 1% \(\text{H}_2\text{SO}_4\)). Rates were normalized to the reactor OD \(600\text{nm}\).

**Sulfur (S\(^0\)) determination.** Sulfur was determined by a modified form of the method of Sorbo (24) to reduce interference from medium components and loss of sulfur during processing. Poly carbonate filters (0.1-µm Whatman Cyclopore track etched membrane 7060-2501 or Millipore Isopore VCTP filters) were rinsed with MilliQ water and placed on a vacuum filter unit. Liquid samples from batch and continuous cultures, as well as bacteria removed from Gelrite plates and resuspended in 0.5 ml sterile demineralized water, were immediately filtered to prevent the bacteria from metabolizing the S\(^0\) before processing. Up to 5 ml culture was carefully loaded directly onto the membrane to prevent S\(^0\) from sticking to the glass of the vacuum unit and filtered under vacuum. Controls consisted of 3 ml MM containing 1% \(\text{H}_2\text{SO}_4\). The filters were rinsed with 1 ml MilliQ water and inserted into a 2-ml Eppendorf tube, and 1.5 ml 0.1 M KCN was added immediately to stop the cells from metabolizing the S\(^0\). The samples were incubated at 90°C for 10 min and cooled to RT, and 200 µl 0.75 M Fe(NO\(_3\))\(_2\) in 20% HNO\(_3\) was added. Samples were centrifuged for 5 min at 16,000 × g to pellet precipitate cell debris and filters, and the supernatant absorbance was measured immediately at a wavelength of 460 nm. Standard curves were prepared in the same manner, with a solution of S\(^0\) in acetone (259 mg in 100 ml) diluted in MilliQ water to a final amount of up to 75 µg S\(^0\) per filter.

S\(^0\), \(\text{H}_2\text{S}\), and \(\text{SO}_3\)\(^2-\) accumulation in bacterial cells. The accumulation of intermediates during CS\(_2\) respiration was measured simultaneously but in separate reaction chambers as follows. Samples from the minichemostats were diluted 10-fold with O\(_2\)-saturated MM containing 1% \(\text{H}_2\text{SO}_4\). The diluted cultures were used to fill three glass cuvettes containing stir bars, after which the cuvettes were closed and placed in a 22°C water bath for 35 min with vigorous stirring. CS\(_2\) (17 µM) was added to each cuvette, and simultaneous measurements were then made of (i) \(\text{O}_2\) respiration with an \(\text{O}_2\) sensor (Unisense), (ii) \(\text{H}_2\text{S}\) production with an \(\text{H}_2\text{S}\) sensor (Unisense), and (iii) S\(^0\) production by following the change in OD \(480\) with an Agilent 8453 spectrophotometer. In addition and simultaneously, a 100-ml glass-syringe containing stir bars was filled with diluted culture and incubated in a 22°C water bath for 35 min with continuous stirring. CS\(_2\) was also added to the syringe, and at intervals during the respiration and \(\text{H}_2\text{S}\) production curves, samples were pushed out of the syringe into an Eppendorf tube and frozen immediately for subsequent S\(^0\) determination.

In order to study \(\text{H}_2\text{S}\) and S\(^0\) formation kinetics during CS\(_2\) respiration in more detail, these parameters were also simultaneously measured in one cylindrical cuvette with three entry ports, which was positioned in a single-beam spectrophotometer. Two ports were used for \(\text{O}_2\) and \(\text{H}_2\text{S}\) sensors, and the third served for the addition of CS\(_2\).

The percentage of S\(^0\) accumulated was calculated as follows. The total \(\text{O}_2\) consumption is represented by the formula \(\text{H}_2\text{S} + 2\text{O}_2 \rightarrow \text{H}_2\text{SO}_4\). In a respiration curve, the total \(\text{O}_2\) consumed \((A)\) will depend on the total amount of \(\text{H}_2\text{S}\) (produced from CS\(_2\)), and so, \(A/2\) represents the amount of sulfide oxidized to sulfate. Sulfur S\(^0\) is an intermediate and is oxidized according to the formula \(\text{S}^0 + 1.5\text{O}_2 + \text{H}_2\text{O} \rightarrow \text{H}_2\text{SO}_4\). At the sulfur peak \((A_{\text{peak}})\), only S\(^0\) is present and the amount of oxygen consumed \((a)\) from then until the end represents sulfur oxidation, and so, \(a/1.5\) represents the amount of S\(^0\) present at the peak. This gives a \(S^0/\text{total sulfide ratio of} \left|\frac{a/1.5}{A/2}\right| = 1.33 \times a/\text{A}\), where \(\text{A}\) is the fraction of the oxygen consumption after the sulfur peak compared to the total consumption, determined from the respiration curve.

**Sulfite (SO\(_3\)\(^2-\)) determination.** Sulfite (SO\(_3\)\(^2-\)) was measured by the method described by Truper and Schlegel (25) but with reduction of the amount of \(\text{H}_2\text{SO}_4\) in the assay to take into account the amount of \(\text{H}_2\text{SO}_4\) that is present in the samples (in these experiments, 1% [vol/vol]). Dilutions of an anaerobic 1 M stock of Na\(_2\)SO\(_4\) were used as standards.

**Starvation experiments.** Starvation experiments were performed in steady-state chemostats by stopping the influent and effluent pumps and removing the CS\(_2\) supply. One-milliliter samples were removed at regular intervals from the reactors during starvation and recovery, diluted 10× in MM containing 1% [vol/vol] \(\text{H}_2\text{SO}_4\), and incubated in a 7.63-m1 double-port cuvette at 22.0 to 22.2°C. Ten minutes after the sample was removed from the chemostat, a 10 µM CS\(_2\) pulse from a 6 ml stock bottle (see above) was injected into the cuvette. \(\text{H}_2\text{S}\) production and removal, as well as respiration, were measured simultaneously with an \(\text{H}_2\text{S}\) microsensor (Unisense) and an oxygen sensor (Strathkelvin Instruments), respectively.

**Nucleotide sequence accession numbers.** The 16S ISR sequences determined in this research have been deposited in the GenBank database under accession numbers KC902816 to KC902829 and KC902831.

**RESULTS**

**Enrichment and isolation of CS\(_2\)-utilizing bacteria.** To obtain extremely acidophilic CS\(_2\)-converting microorganisms, samples from naturally acidic, sulfur-rich environments and from industrial biotrickling filters were incubated in acidified MM and with CS\(_2\) as the sole energy source. Significant CS\(_2\) conversion was observed within a few days, and dense cultures were obtained within 2 to 4 weeks of incubation with 1 to 4% \(\text{H}_2\text{SO}_4\). Ultimately, some enrichment cultures showed growth at 6% \(\text{H}_2\text{SO}_4\) (Table 1). This concentration is equal to a theoretical pH of ~0.05. Enrichments containing 4 to 6% \(\text{H}_2\text{SO}_4\) (Table 1) were used to isolate 16 pure cultures on Gelrite plates (0.1% \(\text{H}_2\text{SO}_4\)). To reconfirm their acid tolerance, all isolates were successfully transferred to 4% \(\text{H}_2\text{SO}_4\) medium.

Descriptions of growth characteristics on plate cultures and in liquid cultures are given in the supplemental material and shown in Fig. S1 and S2 in the supplemental material. The 16 isolated strains showed distinctly different colony morphologies, ranging from large, dry, white colonies to smooth, shiny, compact colonies. Stationary-phase liquid cultures showed white or yellow aggregates or little aggregation. From the cryo-SEM analysis of surfaces and cross sections of colonies, we conclude that on solid medium, the compactly growing strains produce more extracellular polymeric substance (EPS) than the spreading strains do.

**Phylogenetic analysis.** All of the new CS\(_2\)-converting isolates were identified as *Acidithiobacillus thiooxidans* strains by conventional 16S rRNA gene analysis (data not shown). Improved discrimination was achieved by analysis of the 16S-23S ISR sequences, containing two tRNA genes and three intergenic transcribed spacers (ITS). The total length of the ISR sequences of the CS\(_2\)-converting isolates varied between 456 and 460 bp, with one exception; isolate BDW2 from the Oy Visko reactor had an ISR of 439 bp (see Table S2 in the supplemental material). Within the ISR, most of the variation in nucleotide composition was observed in the third of the three ITS. Strains BAW3 and BBW1, which originated from two parallel reactors operating at the Loudon factory, were distinctly different with respect to their nucleotide composition, differing by 57 bp and harboring three insertions and three deletions compared with reference strain G8. In a neighbor-joining phylogenetic tree constructed from the 16S ISR sequences of the CS\(_2\)-converting strains and other *A. thiooxidans* strains, strains BAW3 and BBW1 form a distinct cluster within the *A. thiooxidans* strains (Fig. 1). Three more clusters could be distinguished, i.e., cluster 2, which did not contain any of the CS\(_2\)-converting strains; cluster 3, comprising most of the CS\(_2\)-converting strains and also *A. albertensis*, described as a distinct species but phylogenetically indistinguishable from *A. thiooxidans* (26); and cluster 4, containing strain S1p from the Solfatara.
The OD of strain Sts 4-3 was consistently lower than that of the other four strains, probably because of the observed smaller amount of elemental sulfur in the culture.

The \( \mu_{\text{max}} \) values of the strains growing under these conditions were determined by increasing the medium flow and simultaneously increasing the \([\text{CS}_2]\) supplied to the reactors with the same factor (resulting in a constant \([\text{CS}_2]\) load per milliliter of medium input), up to and beyond the point where \( S^0 \) formation started becoming visible in the reactors. Strains Sts 4-3 and G8 had slightly higher \( \mu_{\text{max}} \) values than the other three strains (Table 2).

**Cell surface hydrophobicity.** The differences in growth characteristics on plates and in liquid culture suggested that there may be differences in the surface properties of the strains. To test this, cell surface hydrophobicity was measured by using adherence to octane and hexane (23). Strains 2Bp and Sts 4-3 had less hydrophobic cell surfaces than strains Sts 4-3, G8, and BBW1 (see Fig. S3 in the supplemental material). On plates, strains 2Bp and Sts 4-3 produced compact, smooth, and opaque colonies versus the dry and spread-out colonies of strains Sts 4-3, G8, and BBW1. Similar results were obtained when experiments were performed at pH 7 (see Fig. S3A and B) or 3 (see Fig. S3C and D); therefore, a different response to high-pH shock is not the cause of the differences in surface hydrophobicity observed.

**pH optimum and pH tolerance.** The pH optima of the five selected strains were determined with a floating-filter assay on media with pHs ranging from 0.5 to 6. The pH optima varied among 1 (2Bp), 1.5 (Sts 4-3 and G8), and 2 (Sts 4-3) (Table 2). Strain BBW1 did not form colonies but showed spreading surface growth on the filter and the medium as well. On the basis of visual observations, the pH optimum of this strain was between 1 and 1.5 (Table 2).

Tolerance of different pH levels was estimated by counting the colonies that appeared on the filters after the transfer of cells growing at 1% \( \text{H}_2\text{SO}_4 \) (measured pH, 0.72) to media with lower or higher pHs. Survival rapidly decreased when cells were exposed to low pH.
more acid (pH 0.5) and more gradually decreased when cells were exposed to higher-than-optimum pHs. Strain BBW1 was the most sensitive to high-pH stress; no growth was observed in medium with a pH higher than 3.5.

Enzyme kinetic analysis. To assess the CS2 removal efficiencies of the five new CS2-converting *A. thiooxidans* isolates, the CS2 affinity constants ($K_m$) and maximum CS2 conversion rate ($V_{max}$) of crude protein extracts from continuous cultures were determined by adding a pulse of CS2 to diluted protein extracts and measuring the rate of H2S production. Table 2 shows that the $K_m$ values of the five strains are of the same order of magnitude, ranging between 81 and 130 μM CS2 for strains G8 and 2Bp, respectively. Comparison of the $V_{max}$ values revealed crude extracts from strain S1p to have a consistently higher $V_{max}$ under these conditions than crude extracts from the other four strains tested, resulting in a 2-fold higher $k_{cat}/K_m$ value for S1p. In accordance, the $V_{max}$ for CS2 of steady-state, minichemostat-grown, intact S1p cells was consistently slightly higher than the $V_{max}$ of the other strains (Table 2).

Sulfur production and consumption. (i) Sulfur production. Production of $S^0$ by CS2-converting acidithiobacilli as an intermediate in the oxidation of $H_2S$ to $SO_4^{2-}$ can block biotrickling filters and subsequently decrease performance. Therefore, we compared the $H_2S$ and $S^0$ production and consumption of the five selected *A. thiooxidans* strains upon CS2 pulses. When concentrations as low as 1.8 μM CS2 (equivalent to 36 ppm CS2 in the gas phase) were added to a cuvette containing diluted samples from the minichemostats, there was an almost immediate increase in $S^0$, just above the detection level. Peak concentrations of $H_2S$ ranged from 0.5 to 1.0 μM under these conditions, and so, $S^0$ formation already started at $H_2S$ concentrations below this value. In subsequent experiments, $H_2S$ and $S^0$ production was measured simultaneously in one cuvette. Addition of 17 μM CS2 resulted in an immediate and large accumulation of $H_2S$ for all strains. Also, $S^0$ formation started almost without delay and continued until virtually all of the $H_2S$ had been consumed (Fig. 2). The respiration curves show an initial period of fast respiration when $H_2S$ is still present, followed by a sudden decrease in the respiration rate when the cells start respiring solely on $S^0$. This decrease in the respiration rate indicates that the processing of the $S$ intermediates in the cell, and not respiration, is the rate-limiting step. This was observed in all of the strains, but it was most prominent in strain S1p (Fig. 2). The chemical reaction of $H_2S$ with O2 was less than 2 μM $H_2S / O_2$ and therefore did not significantly contribute to the observed oxygen consumption rates.

During the period of $S^0$ respiration, we often observed a temporary reduction in the respiration rate. Although it is possible that an intermediate temporarily accumulated that inhibited res-
TABLE 2 Comparison of CS₂-converting A. thiooxidans strains 2Bp, Sts 4-3, S1p, G8, and BBW1

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>2Bp</th>
<th>Sts 4-3</th>
<th>S1p</th>
<th>G8</th>
<th>BBW1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Colony morphology</td>
<td>Smooth, domed</td>
<td>Small, smooth, domed</td>
<td>White, dry, compact</td>
<td>White, dry, spreading</td>
<td>White, dry, spreading</td>
</tr>
<tr>
<td>A. thiooxidans 16S ISR cluster (Fig. 1)</td>
<td>3</td>
<td>3</td>
<td>4</td>
<td>3</td>
<td>1</td>
</tr>
<tr>
<td>Maximum acid tolerance (% [vol/vol] sulfuric acid)</td>
<td>5</td>
<td>5</td>
<td>5</td>
<td>4</td>
<td>6</td>
</tr>
<tr>
<td>pH optimum</td>
<td>1–6</td>
<td>0.5–6</td>
<td>1–6</td>
<td>0.5–6</td>
<td>0.5–3.5</td>
</tr>
<tr>
<td>Cell surface hydrophobicity</td>
<td>Low</td>
<td>Low-intermediate</td>
<td>High</td>
<td>High</td>
<td>High</td>
</tr>
<tr>
<td>Reactor dilution rate at point of S₀ formation</td>
<td>0.09</td>
<td>0.08</td>
<td>0.1</td>
<td>0.1</td>
<td>0.09</td>
</tr>
<tr>
<td>(approaching p_max)</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Mean cell extract Kₘ (µM CS₂) ± SEM</td>
<td>130 ± 22</td>
<td>97 ± 10</td>
<td>100 ± 5</td>
<td>81 ± 7</td>
<td>116 ± 4</td>
</tr>
<tr>
<td>Mean cell extract V_max (µmol H₂S min⁻¹ mg protein⁻¹) ± SEM</td>
<td>38 ± 3</td>
<td>17 ± 0</td>
<td>48 ± 7</td>
<td>23 ± 1</td>
<td>28 ± 5</td>
</tr>
<tr>
<td>k_cat/K_m (µM⁻¹ s⁻¹)</td>
<td>3.5</td>
<td>2.1</td>
<td>6.5</td>
<td>3.4</td>
<td>3.3</td>
</tr>
<tr>
<td>Mean whole-cell V_max (µmol H₂S min⁻¹ ml culture⁻¹ for OD₆₀₀ of 1) ± SEM</td>
<td>11 ± 0</td>
<td>10 ± 0</td>
<td>12 ± 0</td>
<td>8 ± 0</td>
<td>11 ± 0</td>
</tr>
<tr>
<td>Mean calculated S₀ accumulated (% of total S added) ± SEM</td>
<td>76 ± 4</td>
<td>83 ± 1</td>
<td>82 ± 1</td>
<td>83 ± 1</td>
<td>ND a</td>
</tr>
<tr>
<td>Mean measured S₀ accumulated (% of total S added) ± SEM</td>
<td>67 ± 2</td>
<td>63 ± 2</td>
<td>65 ± 1</td>
<td>61 ± 1</td>
<td>ND</td>
</tr>
<tr>
<td>Mean respiration rate on H₂S and S₀ combined after a 10 µM CS₂ pulse (nmol O₂ min⁻¹ ml culture⁻¹ for OD₆₀₀ of 1) ± SEM</td>
<td>45 ± 9</td>
<td>43</td>
<td>87 ± 10</td>
<td>101 ± 1</td>
<td>66 ± 7</td>
</tr>
<tr>
<td>Mean respiration rate on S₀ only after 10 µM CS₂ pulse (nmol O₂ min⁻¹ ml culture⁻¹ for OD₆₀₀ of 1) ± SEM</td>
<td>41 ± 3</td>
<td>7</td>
<td>63 ± 3</td>
<td>44 ± 1</td>
<td>49 ± 1</td>
</tr>
<tr>
<td>Mean % of S₀ globules attached to cells after 10 17 µM CS₂ pulses ± SEM</td>
<td>87 ± 7</td>
<td>17 ± 4</td>
<td>9 ± 2</td>
<td>52 ± 2</td>
<td>22 ± 2</td>
</tr>
<tr>
<td>Mean % of cells attached to S₀ globules after 10 17 µM CS₂ pulses ± SEM</td>
<td>35 ± 10</td>
<td>13 ± 3</td>
<td>7 ± 2</td>
<td>30 ± 6</td>
<td>13 ± 0</td>
</tr>
<tr>
<td>Resistance to CS₂ stress</td>
<td>Low</td>
<td>Medium</td>
<td>Medium</td>
<td>Medium</td>
<td>Medium</td>
</tr>
<tr>
<td>Speed of recovery after CS₂ starvation</td>
<td>Slow</td>
<td>Slow</td>
<td>Slow</td>
<td>Fast</td>
<td>Fast</td>
</tr>
</tbody>
</table>

a Strains were grown in continuous culture (D = 0.02) with CS₂ as the sole energy source. Samples were taken from these reactors for the comparison experiments. The Kₘ and V_max of each strain were calculated from Michaelis-Menten plots by nonlinear regression with the Michaelis-Menten equation V = V_max × S/K_m + S. Mean values from at least three independent experiments are shown.

ND, not determined.

piration on S₀, at this point, the S₀ consumption rate did not decrease. Also, although detectable levels of the potentially inhibiting intermediate SO₃²⁻ were present throughout the experiment, there was no increase at the reduced respiration “bump.” Therefore, the apparent temporary reduction in respiration is more likely due to the sensor being affected by an intermediate at that point in the experiment. At the point where S₀ peaks, the H₂S and CS₂ had been completely consumed and only 30 to 40% of the total oxygen was consumed. Therefore, the remaining O₂ consumption resulted entirely from S₀ oxidation, if the accumulation of any other intermediates is excluded. With this O₂ consumption, and taking into account that this is an underestimate of the actual O₂ consumption because of a small baseline drift upward of the O₂ sensor during the experiment, we calculated that at least 80% of the S added as CS₂ accumulated as S₀. This was similar for all of the strains tested (Table 2), and the percentage also remained identical when CS₂ concentrations as low as 1.8 µM were added. Below this value, H₂S formation could not be observed any more, making such calculations impossible.

The rates of S₀ formation and consumption were calculated from the slopes before and after the apex of the OD₄₈₀ trace (Fig. 2). Consistently, S₀ production (corrected for consumption) was between four times (strains 2Bp, S1p, G8, and BBW1) and nine times (strain Sts 4-3) faster than S₀ consumption. This matches with a buildup of S₀ of at least 80% of the total S added, as calculated above from the oxygen respiration. In separate experiments, chemical analysis for S₀ at the moment OD₄₈₀ reached its maximum value consistently showed S₀ recovery between 60 and 67% (Table 2). Therefore, chemical analysis appeared to underestimate the amount of S₀ produced during CS₂ respiration. However, both methods of determining S₀ accumulation upon the application of CS₂ pulses indicated that all of the strains tested produced about equal amounts of extracellular S₀ under conditions of excess CS₂.

(ii) Sulfur consumption. Although all of the strains tested behaved similarly in terms of S₀ production, there were some obvious differences in subsequent S₀ respiration (Fig. 2); strain Sts 4-3 respired on S₀ five to eight times more slowly than the other four strains. The H₂S combined with S₀ respiration rates (the first part of the respiration curve) showed a maximum 2-fold difference between the strains (Fig. 2 and Table 2). Although strain Sts 4-3 respired more slowly on S₀, the respiration rate did not decline until all of the S₀ had been depleted.
This was in contrast to the other strains, especially strain S1p, which showed a continuous decline in the rate of respiration and concurrent S0 depletion.

We hypothesized that different strains produce differently bioavailable S0; strains 2Bp, G8, and BBW1 produce S0 that can be efficiently metabolized again, whereas strains Sts 4-3 and S1p produce S0 that is difficult to remove. To test this hypothesis, we examined sulfur accumulation by different strains upon the application of repeated CS2 pulses to undiluted chemostat samples by counting S0 globules under a light microscope (Table 2). We counted 160 to 240 cells of each strain, and a representative picture of strains 2Bp, Sts 4-3, S1p, and G8 is shown in Fig. S4 in the supplemental material. All five of the strains tested showed the accumulation of S0 globules of similar sizes, either attached to the bacteria or loose in the medium (see Fig.S4). There were large differences in the observed cell attachment to S0 globules. Although a correlation between the attachment of cells to S0 globules and cell surface hydrophobicity might be expected, we did not observe this despite the fact that all of the cells had identical physiological backgrounds. Strain S1p clearly had the fewest S0 globules that were attached to cells (9% ± 2%), while in cultures of strain 2Bp, most of the S0 globules were attached to cells (87% ± 7%, Table 2). This supports our hypothesis that S1p produces S0 that appears less bioavailable; attachment of S0 globules to bacteria is required for rapid S0 consumption upon H2S depletion. Strain Sts 4-3, which consumes S0 very slowly, also had a low percentage (13%) of cells attached to S0 globules, and 17% of the S0 globules were attached to cells (Table 2). This percentage was not as low as that of strain S1p, and it therefore cannot entirely explain the very low observed S0 consumption rate of this strain. However, strain Sts 4-3 produced more H2S than the other strains when pulsed with CS2 (25 versus 20 μM, Fig. 2), which could have had an inhibitory effect on overall respiration.

**CS2 stress and starvation.** The effect of fluctuations in CS2 concentrations in biofilters can be simulated by applying CS2 pulses to undiluted culture samples from CS2-limited steady-state chemostats in MM with 1% H2SO4. Therefore, chemostat samples of the newly isolated *A. thiooxidans* strains were incubated and subjected to repeated 17 μM CS2 pulses. Each new pulse was given when the H2S produced from the previous pulse had been

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**FIG 2** Conversion of CS2 by five *A. thiooxidans* strains. The arrows indicate the times when 17 μM CS2 was injected into a cuvette containing cells from a continuous culture growing at D = 0.02, diluted 6× in MM containing 1% H2SO4. Respiration (O2 consumption, black dashed line), H2S production (black solid line), and S0 production (gray line) were monitored. The production and consumption rates in these graphs were used to determine the percentages of S0 accumulation in Table 2. Note the different scale on the x axis of the graph for strain Sts 4-3.
consumed by the cells (after about 1.5 to 3 min). The H$_2$S consumption rates after the first CS$_2$ pulse varied between 1.9 (strain BBW1) and 3.4 μmol H$_2$S liter$^{-1}$ min$^{-1}$ unit of OD$_{600}^{-1}$ (strain G8) and decreased for all of the strains after repeated CS$_2$ pulses, with strain 2Bp appearing to be the most strongly affected (1.0 μmol H$_2$S liter$^{-1}$ min$^{-1}$ unit of OD$_{600}^{-1}$, 41% of the initial rate) and strain BBW1 appearing to be the least strongly affected (1.5 μmol H$_2$S liter$^{-1}$ min$^{-1}$ unit of OD$_{600}^{-1}$, 76% of the initial rate, Fig. 3). This indicates that the cells were becoming increasingly stressed when subjected to CS$_2$ loads above the concentration the cells were adapted to in the chemostat. This is probably due to the toxic effect of the repeated buildup of H$_2$S in the cells.

To test the effect of intermittent interruption of the factory CS$_2$ supply, the five strains growing in chemostats as described above were subjected to CS$_2$ starvation by temporarily shutting down the medium and CS$_2$ supply for 24 h. During this period of starvation and the subsequent 24-h recovery period when the CS$_2$ supply was reconnected again, samples were taken from the chemostats and subjected to a CS$_2$ pulse. Subsequent H$_2$S production and consumption rates and respiratory activity were measured. Results indicated that the strains cope reasonably well with short periods of starvation. After 4 h of CS$_2$ starvation, there was only modest to no reduction in respiratory activity when the cells were pulsed with 10 μM CS$_2$. Of the four strains tested at this time point, G8 appeared to be the most strongly affected, with a 40% reduction of its initial respiratory activity, when both H$_2$S and S$^0$ were present and a 22% reduction in S$^0$ respiratory activity (Fig. 4). After 24 h of CS$_2$ starvation, all of the strains showed reduced respiratory activity. Although not tested extensively, the reduction was largest for strain 2Bp, where H$_2$S/S$^0$ respiratory activity was reduced by 80% and S$^0$ respiration was reduced by 85% below the steady-state rates. Strain 2Bp also seemed to recover slowly from starvation, whereas the other four strains showed a doubling of initial respiratory activity back to steady-state levels after only 4 h of resumed CS$_2$ supply, the activity of strain 2Bp increased only 1.4-fold to 29% of the steady-state level. This slow recovery corresponds to the observation described above that strain 2Bp suffers more from CS$_2$ stress than the other four strains do. At 24 h of recovery, the culture density of all five strains was higher than during steady state. This indicates that the cultures had started growing and/or had S$^0$ present.

**DISCUSSION**

Our approach to the enrichment of extremely acidophilic CS$_2$-converting microorganisms from volcanic regions and CS$_2$-converting trickling filters proved successful. Conversion of CS$_2$ was observed within days after inoculation. Furthermore, the enrichments and isolated *Acidithiobacillus* strains could grow at very high H$_2$SO$_4$ concentrations. The only acidophilic CS$_2$-converting *Acidithiobacillus* strain described to date, TJ330, was isolated from an acidic (pH 6.1 to 1.4) CS$_2$- and H$_2$S-treating peat biofilter (19). The neutrophilic CS$_2$-converting species *Paracoccus denitrificans* was isolated from oak leaves and soil beneath the leaf canopy (7, 27). Our attempt to enrich acidophilic microorganisms from soil samples taken under an oak tree and from compost was unsuccessful. Of the *Acidithiobacillus* strains from culture collections screened for CS$_2$ conversion (at micromolar concentrations), only *A. caldus* DSM 8584 was positive. This trait was not previously reported for this species. Our screening confirmed the observation of Smith and Kelly (28), although they used a potentially toxic liquid CS$_2$ concentration (2 mM).

The new *Acidithiobacillus* isolates even showed slow growth in 6% H$_2$SO$_4$. The constant supply of CS$_2$ to the cultures was the key to experimentally creating a CS$_2$ stress.
factor in the maintenance of cultures at these very high H₂SO₄ concentrations. Culture activity decreased dramatically upon storage without a substrate at or above 4% H₂SO₄. Acidophiles require a continuous and high supply of maintenance energy to be able to actively pump out protons that leak from the acidic environment into the nearly neutral pH cell cytoplasm (29). Operation of biofilters under highly acidic conditions therefore puts additional stress on the microorganisms during periods of fluctuating CS₂ concentration and factory shutdown.

Although growth was very slow at 6% H₂SO₄, dense growth was already observed at 4% for all new Acidithiobacillus strains. Similar H₂SO₄ tolerance was reported for Acidithiobacillus sp. strain AZ11, which is able to respire on elemental sulfur in the presence of 4.2% H₂SO₄ (30). The pH optima determined for 5 of the 16 new CS₂-converting strains were between 1 and 2. This is slightly lower than the optimal-pH range of 1.8 to 2.5 described for other Acidithiobacillus strains (29, 31).

Growth at around pH 0, as observed for 15 of our A. thiooxidans isolates, is among the lowest pHs reported in the literature for any organism. *Ferromonas acidarmanus* (32), *Picrophilus oshimae*, and *Picrophilus torridus* could grow at pH 0 (optimum pH, 0.7) (33). *Picrophilus torridus* adapted to growth at pH 0.1 even showed significant growth at 1.2 M H₂SO₄ (pH −0.06) (34). The eukaryotic red alga *Cyanidium caldarium* was cultured at 0.5 M H₂SO₄ (35), and the green alga *Dunaliella acidophila* is able to survive pH 0.2 (36). Also, some fungal species were reported to grow at pH 0, i.e., *Acanthidium clyatum* (37), *Cephalosporium sp.*, and *Trichosporon cutaneum* (38). We also observed fungal growth at 6% H₂SO₄ (pH −0.05) in our enrichment cultures but did not further investigate the species present in these cultures. With growth at pH as low as −0.05 (6% H₂SO₄), the new Acidithiobacillus isolates obtained in this research exceed the previously reported pH limit of 0.5 for microbial CS₂ conversion for Acidithio- bacillus TJ330 (9).

The newly isolated CS₂-converting *A. thiooxidans* strains differed in colony morphology. The genus Acidithiobacillus comprises a physiologically and genetically heterogeneous group of microorganisms (39, 40), despite the often low sequence diversity in the 16S rRNA gene. For that reason, the 16S−23S ISR is used to discriminate at the intra species level (41, 42).

Two main colony types on Gelrite plates were distinguished, compact, creamy, shiny colonies and dry, white, spreading colonies (40). Reversible variation in colony morphology of several *A. ferrooxidans* strains has been described, resulting in the same large, white spreading colonies, as opposed to compact colonies, as observed here for some of the newly isolated *A. thiooxidans* strains (43, 44). *A. thiooxidans* strains may be motile via a polar flagellum (40). *A. ferrooxidans* ATCC 19859 spreading variants displayed increased motility and chemotaxis toward thiosulfate, which may be a selective advantage over biofilm growth during periods of low substrate concentrations. These variants arose through the rearrangement of insertion sequences in the genome, potentially acting as a genetic switch (44). The *A. ferrooxidans* group of strains has recently been reclassified into four separate species (45). Of these, only *A. ferrivorans* and some *A. ferrioxidans* strains were shown to be motile (45). The type strain *A. ferrooxidans* ATCC 23270 lacks flagellum and chemotaxis genes (46). The draft genome of *A. thiooxidans* ATCC 19377 (47) and the draft genomes of both *A. thiooxidans* strains Sts 4-3 and G8 (Daan Speth, personal communication) do contain the operons for flagellum biosynthesis and chemotaxis. However, we did not observe swimming motility in liquid cultures of our strains when we examined them microscopically. Therefore, the mechanism and role of the spreading colony phenotype in these strains are not clear.

SEM studies of frozen colonies indicated that the difference in colony appearance may be caused by the absence of a clear EPS layer on the dry white colonies, as has been observed for colony morphology mutants of *Mycobacterium smegmatis* (22). In support, *A. thiooxidans* strains Sts 4-3 and some other white-dry colonies, which had a considerably higher cell surface hydrophobicity than strains 2Bp and Sts 4-3, which produced compact shiny colonies, suggesting the presence of relatively more hydrophilic compounds on the cell surface of the latter strains. Acidithiobacillus strains produce EPS containing both neutral sugars and fatty acids (48, 49), the proportion varying depending on the substrate the cells are grown on (50, 51). The EPS of *A. thiooxidans* grown on S⁰ consists of 40% sugars and 60% fatty acids (mainly eicosanoic acid) (50). The fatty acids are released from the outer membrane by blebbing (52) and cause the “wetting” of sulfur particles described in 1961 by Jones and Starkey (53), making S⁰ available as an energy source. In addition, EPS is essential for successful attachment and bioleaching of *A. ferrooxidans* to pyrite (50). Differences in EPS production and potential motility observed in our strains may have implications for the degree of colonization and clogging of biofilters.

Efficient biofiltration of CS₂-contaminated air streams requires microorganisms with a higher affinity for CS₂ than the concentration present in the air stream, to ensure its removal to concentrations complying with increasingly stringent regulations, as well as resistance to changes in operating conditions, low biomass production, low S⁰ accumulation, and rapid S⁰ removal to prevent clogging of the biofilters. The affinities of the five strains tested for CS₂ were similar (around 100 μM) and correspond to that reported for the purified CS₂ hydrolase of *Acidianus A1-3* (10). Comparison of the *Vₘₐₓ* values revealed strain Sts 1p to have a consistently higher *Vₘₐₓ* under these conditions than the other four strains tested, possibly because it has a relatively larger amount of CS₂ hydrolase present in the cells than the other strains (15). The *Vₘₐₓ* is only reached at substrate concentrations of 200 μM (nmol ml⁻¹) or higher. Bioreactors treating CS₂-contaminated air from the viscose industry are operational at much lower concentrations of around 4 to 20 nmol ml⁻¹ (100 to 500 ppm) (2), indicating that although strain Sts 1p converts CS₂ with a higher *Vₘₐₓ* than the other strains, it would not usually reach its full potential reaction rate in a bioreactor. Indeed, a higher *Vₘₐₓ* may even be deleterious, as it might cause H₂S to accumulate to toxic levels in the cell more rapidly upon CS₂ peaks.

Comparison of the resistance to changes in operating conditions in the form of an interruption in the CS₂ supply revealed that all five strains were affected by 24 h of CS₂ starvation in terms of the ability the reprise after a CS₂ pulse during starvation, but they all recovered within 24 h after reconnection of the CS₂ supply. However, during recovery, all five strains will also produce S⁰, as we found in all of the strains tested that CS₂ pulses during starvation resulted in the transient accumulation of 80% of the total S added as S⁰, independently of the amount of CS₂ added. This implies that it will be difficult to avoid S⁰ accumulation in biofiltration systems with uneven CS₂ loading. However, differences in S⁰ removal were observed, with strains Sts 4-3 having a much lower but constant S⁰ removal rate than the other strains and strain Sts 4-3
having a long period in which the S0 respiration rate slowly declined. Whatever the cause, the strains that remove S0 more slowly will cause more clogging problems in bioreactors because of S0 accumulation.

In summary, we have successfully isolated extremely acidophilic, CS2-converting *A. thiooxidans* strains from both environmental and industrial ecosystems that grow optimally around pH 1 to 2 and can grow at sulfuric acid concentrations of up to 6% (vol/vol). Currently bioreactors are operated at pH 0.5 to 1 (2). Use of the new strains would reduce water use and improve the prospect for reuse of the produced sulfuric acid in the rayon/viscose industry. The isolated strains displayed different growth and colony morphology characteristics, which may be due to differences in motility and/or the presence or absence of an EPS layer surrounding the cells. To circumvent the bottlenecks in the biofiltration of CS2, cocultures of extremely acidophilic *A. thiooxidans* strains to combine the best acid tolerance, affinity for CS2, S0-removing potential, and stability during periods of fluctuating CS2 loads are most promising for inoculation of industrial biofilters. This application is expected to result in reduced sulfur accumulation, increased CS2 removal rates, reduced water consumption, a more stable operation, and recycling of the sulfuric acid produced.

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