

Copper Ions Stimulate Polyphosphate Degradation and Phosphate Efflux in *Acidithiobacillus ferrooxidans*

Sergio Alvarez and Carlos A. Jerez*

Laboratory of Molecular Microbiology and Biotechnology, Millennium Institute for Advanced Studies in Cell Biology and Biotechnology, and Department of Biology, Faculty of Sciences, University of Chile, Santiago, Chile

For some bacteria and algae, it has been proposed that inorganic polyphosphates and transport of metal-phosphate complexes could participate in heavy metal tolerance. To test for this possibility in *Acidithiobacillus ferrooxidans*, a microorganism with a high level of resistance to heavy metals, the polyphosphate levels were determined when the bacterium was grown in or shifted to the presence of a high copper concentration (100 mM). Under these conditions, cells showed a rapid decrease in polyphosphate levels with a concomitant increase in exopolyphosphatase activity and a stimulation of phosphate efflux. Copper in the range of 1 to 2 μ M greatly stimulated exopolyphosphatase activity in cell extracts from *A. ferrooxidans*. The same was seen to a lesser extent with cadmium and zinc. Bioinformatic analysis of the available *A. ferrooxidans* ATCC 23270 genomic sequence did not show a putative *pit* gene for phosphate efflux but rather an open reading frame similar in primary and secondary structure to that of the *Saccharomyces cerevisiae* phosphate transporter that is functional at acidic pH (Pho84). Our results support a model for metal detoxification in which heavy metals stimulate polyphosphate hydrolysis and the metal-phosphate complexes formed are transported out of the cell as part of a possibly functional heavy metal tolerance mechanism in *A. ferrooxidans*.

Acidithiobacillus ferrooxidans (formerly *Thiobacillus ferrooxidans*) is a chemolithoautotrophic bacterium that obtains its energy from the oxidation of ferrous iron, elemental sulfur, or partially oxidized sulfur compounds (19, 24). This ability makes it of great industrial importance due to its application in bio-mining to recover metals such as copper, gold, and uranium (19, 23). These microorganisms are normally subjected to stress in their environment, such as temperature and pH changes and the presence of toxic heavy metals and nutrient starvation, which affect their physiological state (30).

Unlike most heterotrophic bacteria, *A. ferrooxidans* is capable of resisting high concentrations of heavy metals such as copper, zinc, arsenic, and uranium (9). The genetic basis for mercury and arsenic resistance has been studied in detail in this acidophile (6, 26). Copper is an essential trace element for all cells. However, it can cause serious cell damage through radical formation (10). Information regarding copper resistance in *A. ferrooxidans* is scarce. Although copper-tolerant strains have been obtained by growth and adaptation to increasingly higher concentrations of this metal (5, 8, 18), only a few genes were recently identified by RNA arbitrarily primed PCR as being induced or repressed in *A. ferrooxidans* subjected to copper (21). Nevertheless, the role of these genes in the mechanism of copper resistance is still unclear, and their expression may be related to indirect metabolic responses to stress (21).

Many heavy metal resistance systems involve either active efflux or detoxification of metal ions by different transformations (27). For copper, these include intracellular complexation, reduced accumulation, extracellular complexation, and

sequestration in the periplasm (13, 25). One of the proposed mechanisms for metal tolerance is the sequestration of metal cations with long polymers of inorganic polyphosphate (17). Polyphosphate is a linear polymer of hundreds of orthophosphate residues linked by phosphoanhydride bonds. Several physiological functions have been attributed to polyphosphate in addition to being a reservoir of phosphate, such as a substitute for ATP, chelator of metals, and adaptation to stress conditions in the cell (17). The main enzyme involved in the biosynthesis of polyphosphate is the polyphosphate kinase, which catalyzes the reversible conversion of the terminal phosphate of ATP into polyphosphate (17). An exopolyphosphatase (PPX), on the other hand, is known to hydrolyze polyphosphate, liberating inorganic phosphate (P_i) (17). These enzymes have been purified from *Escherichia coli*, and their genes have been identified in several bacteria, including *A. ferrooxidans* (30). These genes show a relatively high degree of sequence conservation (7, 28).

It has been proposed that polyphosphate sequesters the heavy metals, thereby reducing their intracellular concentration and, on the other hand, that the hydrolysis of polyphosphate detoxifies the metals (1, 14). Van Veen (29) has shown that the inorganic phosphate transport system (Pit) in *E. coli* and *Acinetobacter johnsonii* can reversibly transport metal phosphates. Later, Keasling and Hupf (15), using genetically engineered strains of *E. coli*, obtained results indicating that not only a large quantity of intracellular polyphosphate but also the ability to synthesize and degrade polyphosphate is important for tolerance to heavy metals. Based on these results and those mentioned above, Keasling (14) proposed a model in which the intracellular cation concentration in bacteria would regulate the activity of PPX, which would in turn degrade polyphosphate and the P_i generated accompanied by cation transport out of the cell through the Pit system.

* Corresponding author. Mailing address: Laboratorio de Microbiología Molecular y Biotecnología, Departamento de Biología, Facultad de Ciencias, Universidad de Chile, Santiago, Chile. Phone: (56 2) 678 73 76. Fax: (56 2) 271 29 83. E-mail: cjerez@uchile.cl.

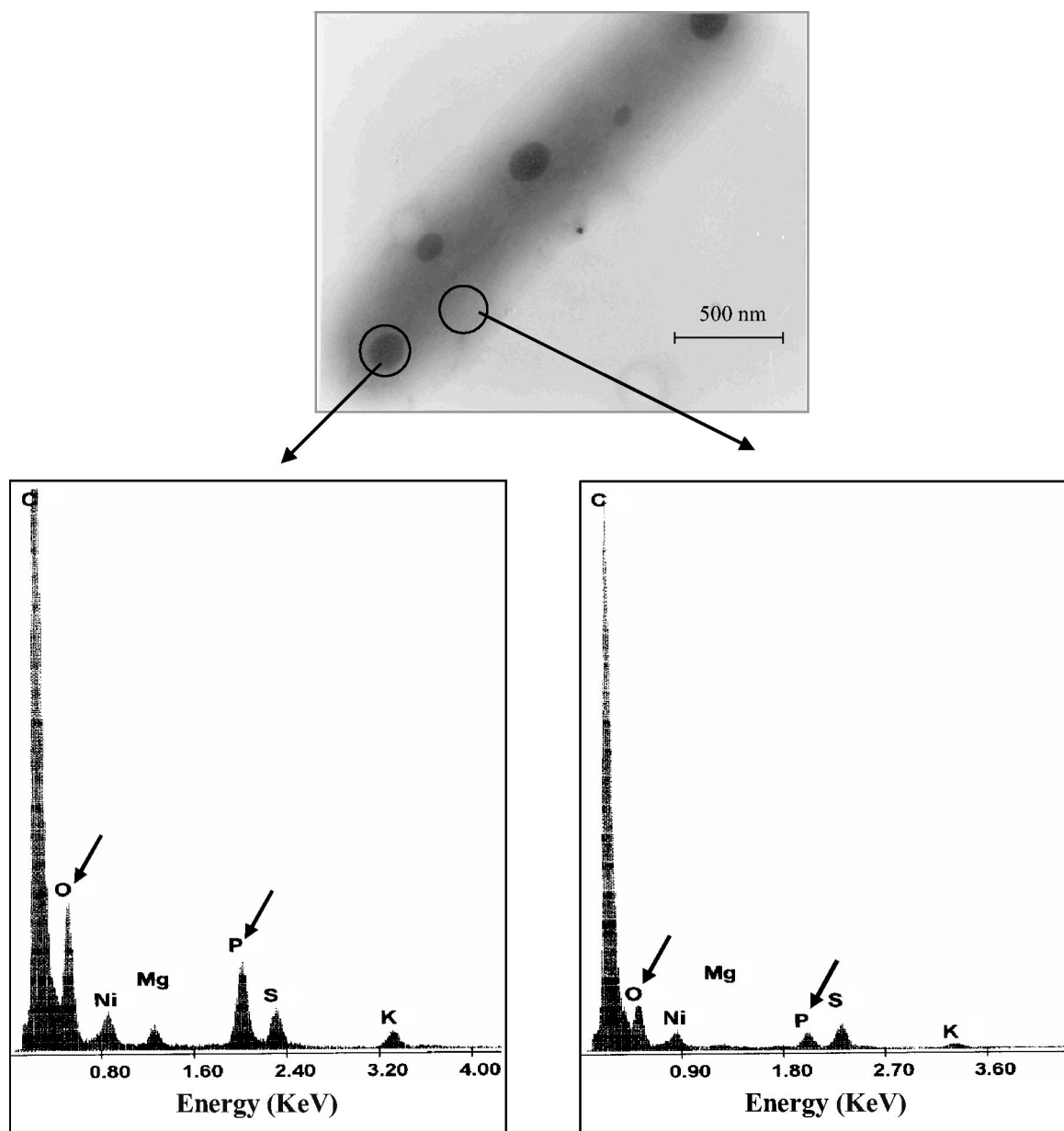


FIG. 1. Transmission electron microscopy and energy dispersive X-ray analysis of *A. ferrooxidans*. Unstained and unfixed cells taken from sulfur-containing medium were examined directly for the presence of electron-dense granules. The elemental composition of a granule (left spectrum) and a cytoplasmic area (right spectrum) was analyzed by energy dispersive X-ray analysis. Arrows indicate the signals corresponding to oxygen and phosphorus.

In the present work, it was found that *A. ferrooxidans* normally accumulates high amounts of polyphosphate granules and that the levels of intracellular polyphosphate are greatly reduced when the bacterium is grown in or shifted to 100 mM Cu^{2+} ions. In the presence of this metal, PPX activity and P_i efflux increased greatly. Our results support a model for metal tolerance mediated through polyphosphate in *A. ferrooxidans*.

MATERIALS AND METHODS

Bacterial strains and growth conditions. *A. ferrooxidans* ATCC 19859 was grown in spherical prills, 0.15 to 0.20 cm in diameter, of elementary sulfur (2) in the presence of 1.75 mM P_i (phosphate-sufficient conditions), unless otherwise indicated. Growth was monitored by measuring cell numbers or the optical

density at 600 nm (OD_{600}). Experiments were conducted when cells reached the late exponential phase ($\text{OD}_{600} = 0.2$). Cultures were adapted to grow in the presence of CuSO_4 by continuous subculturing. During the first culture in the presence of Cu^{2+} (10 mM), a 4- to 5-day lag period was observed. Bacteria were completely adapted to grow in the presence of Cu^{2+} (10 to 100 mM) in the second subculture. It is a known phenomenon that *A. ferrooxidans* cells adapted to grow under these conditions lose this tolerance when they are cultured again in medium without cupric ions (18).

Electron microscopy and X-ray microanalyses. Unstained and unfixed cells were examined for the presence of electron-dense granules by transmission electron microscopy (12). A suspension (10 μl) of *A. ferrooxidans* cells ($\text{OD}_{600} = 0.25$) was placed on a Formvar-coated grid and left for 2 min to allow the cells to sediment. Excess liquid was removed with a piece of filter paper, and the grids were air dried. For analysis, a transmission electron microscope (Philips Tecnai 12) operating at 80 kV was used. Energy-dispersive X-ray microanalysis (EDAX)

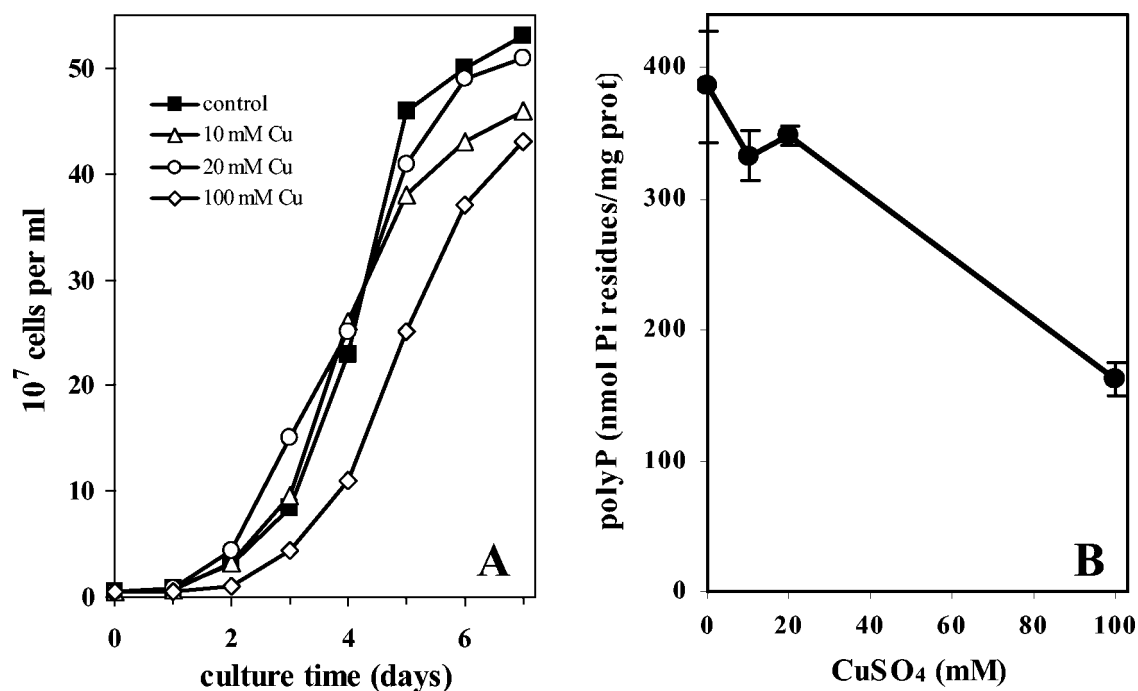


FIG. 2. Growth and polyphosphate levels of *A. ferrooxidans* in the presence of copper ions. *A. ferrooxidans* cultures were inoculated in sulfur medium with 1.75 mM P_i in the presence of the indicated concentration of $CuSO_4$, and cells were counted daily (A). To determine polyphosphate levels (B), the cells in A were harvested in the early stationary phase, and polyphosphate was extracted and quantified by the nonradioactive enzymatic method. Two independent determinations were performed. The error bars represent the standard deviations.

was performed with an EDAX-PV 9800 energy-dispersive microanalyzer at an accelerating voltage of 120 kV (11).

Polyphosphate quantification. Polyphosphate was quantified with a two-step conversion of polyphosphate into ATP by polyphosphate kinase and quantification of the ATP formed by using luciferase to generate light (3). First, polyphosphate was extracted from cell extracts with Glassmilk and then assayed by the reverse reaction of *E. coli* polyphosphate kinase in ADP excess. Finally, the ATP content was assayed with the firefly luciferase ATP assay, and luminescence was measured with a luminometer (BioScan Lumi/96). The concentration of polyphosphate is given in terms of P_i residues.

Preparation of crude cell extracts from *A. ferrooxidans*. Cultures (200 ml) grown to the late exponential phase were harvested by centrifugation ($10,000 \times g$ for 20 min), and the cell pellets were resuspended in 500 μ l of a buffer containing 50 mM Tris-HCl (pH 7) and 10% sucrose and lysed by four cycles of freezing ($-80^\circ C$) and thawing by sonication. The lysate was centrifuged ($5,000 \times g$ for 5 min) to eliminate cellular debris, and the supernatant (crude cell extract) was used to measure PPX activity. These cell extracts were also obtained from nonadapted cells shifted to copper.

Assay for PPX activity. PPX activity was determined as previously reported (16). The 20- μ l reaction mixture contained 50 mM Tris-HCl (pH 6.5), 1 mM $MgCl_2$, 100 mM KCl, 250 μ M [^{32}P]poly P_{750} (polyphosphate with an average of 750 P_i residues). After incubation of the mixture at $30^\circ C$ for 60 min, the reaction was stopped by loading the mixture in polyethyleneimine-cellulose plates for thin-layer chromatography (Aldrich) and development in 0.75 M KH_2PO_4 , pH 3.5. Radioactive spots corresponding to the P_i liberated by the hydrolysis of polyphosphate were visualized and quantified with a phosphor imager (Molecular Imager FX; Bio-Rad). [^{32}P]poly P_{750} was synthesized in vitro by the method of Ault-Riché et al. (3) as described in Cardona et al. (7). One unit of enzyme was defined as the amount releasing 1 pmol of P_i from polyphosphate per min.

In vivo labeling of *A. ferrooxidans* with $^{32}P_i$. Cells were grown in sulfur medium to the late exponential phase in P_i -sufficient conditions (1.75 mM). Cells were collected by centrifugation and resuspended at a higher cell density (10^{10} cells/ml) in medium with reduced P_i (0.18 mM P_i). To label the cells, $H_3^{32}PO_4$ (100 μ Ci/ml) was added, and the microorganisms were further incubated for 17 h, after which the radioactively labeled cells were harvested by centrifugation.

P_i efflux measurements. The $^{32}P_i$ -labeled cells were exhaustively washed by resuspension and centrifugation with fresh medium containing sufficient P_i (1.75 mM) to eliminate the nonincorporated radioactive label and finally resuspended

in the same medium to an OD_{600} of 0.26 (10^9 cells/ml) in the presence or absence of $CuSO_4$. To determine the amount of $^{32}P_i$ released into the medium, samples (1.5 ml) were taken periodically, and the radioactivity in the supernatants obtained by centrifugation at $12,000 \times g$ for 10 min (1.0 ml) was determined by scintillation counting.

Genome sequence analysis. Preliminary sequence data for *A. ferrooxidans* strain 23270 was obtained from the Institute for Genomic Research website at <http://www.tigr.org>. Identity and similarity searches in the databases were done with the tBlastn program from NCBI (<http://www.ncbi.nlm.nih.gov>). The finished available *A. ferrooxidans* ATCC 23270 genomic sequence (<http://www.tigr.org>), which is not yet annotated, was used. The amino acid sequences of the PitA and PitB transporters from *E. coli* and the Pho84 transporter from *Saccharomyces cerevisiae* were used as probes. The possible presence of transmembrane domains in the open reading frames analyzed was studied with the Top-Pred program (<http://bioweb.pasteur.fr/seqanal/interfaces/toppred.html>).

RESULTS AND DISCUSSION

Accumulation of polyphosphate in *A. ferrooxidans*. Previously, it was demonstrated that *A. ferrooxidans* possesses a polyphosphate kinase activity and the corresponding gene, suggesting the existence of polyphosphate in this bacterium (30). Typically, accumulation of polyphosphates in the form of electron-dense granules has been reported in many bacteria (12). To detect the presence of such granules in *A. ferrooxidans*, cells grown in sulfur medium in a P_i -sufficient condition were analyzed by transmission electron microscopy. As Fig. 1 shows, abundant spherical electron-dense granules were observed. All cells grown under these conditions presented at least two granules, and over 90% of the cells presented three to four granules. These granules disappeared when the cells were subjected to phosphate starvation (results not shown).

To confirm the chemical nature of the electron-dense granules, an elemental analysis with EDAX in the transmission

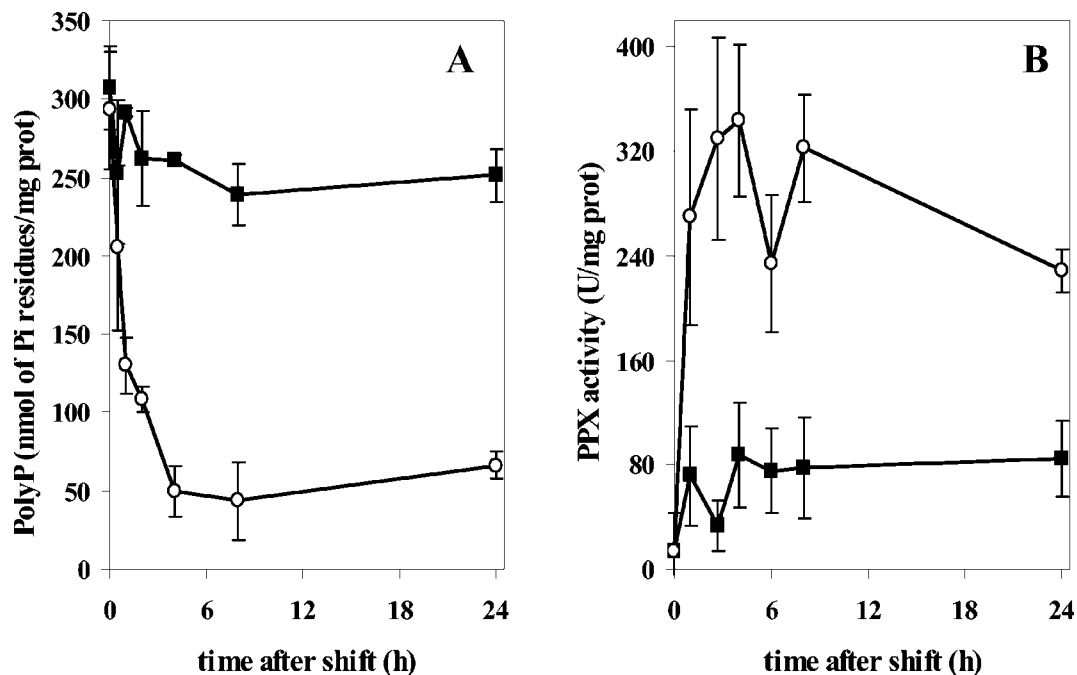


FIG. 3. (A) Reduction in polyphosphate content during exposure to copper ions. *A. ferrooxidans* cells grown in sulfur medium in the absence of copper to the early stationary phase were divided into two portions. CuSO_4 (20 mM final concentration) was added to one sample (○), and an equal volume of fresh medium was added to the control sample (■). Both were then incubated at 30°C. Aliquots were taken at the indicated times, and polyphosphate was quantified. (B) PPX activity in cells of *A. ferrooxidans* shifted to copper. *A. ferrooxidans* was cultured and transferred to medium with copper as in A. Cell extracts were then prepared at each of the indicated postshift times from control cells (■) and cells exposed to 20 mM copper (○), and the PPX activity was determined. The error bars represent the standard deviations.

electron microscopy mode was carried out. As can be seen in Fig. 1, the EDAX spectra showed that the electron-dense bodies present in *A. ferrooxidans* were mainly composed of phosphorus and oxygen (Fig. 1, left spectrum), in contrast to the elemental composition of a cytoplasmic area (Fig. 1, right spectrum), indicating that *A. ferrooxidans* synthesizes and accumulates electron-dense granules containing phosphate, which is most likely polyphosphate. Cells grown under control conditions (P_i sufficient) showed a very high level of polyphosphate, determined enzymatically, of approximately 400 nmol of P_i residues/mg of protein (Fig. 2). This high level of the polymer was in agreement with the presence of abundant electron-dense granules in the cells (Fig. 1). *A. ferrooxidans* can therefore be considered a polyphosphate-accumulating microorganism, like *Acinetobacter johnsonii* (29).

Effect of CuSO_4 on the growth of *A. ferrooxidans* and its polyphosphate levels. To evaluate the effect of Cu^{2+} on polyphosphate levels, cells were grown in the presence of different concentrations of Cu^{2+} and the cellular polyphosphate content was determined. The bacterial cells used in these experiments were previously adapted to grow at different concentrations of copper. There was a small decrease in the bacterial growth rates, and the curves reached the plateaus at slightly lower cell densities when cells were grown in the presence of increasing Cu^{2+} concentrations compared with the control culture in the absence of the metal (Fig. 2A). On the other hand, polyphosphate levels showed a clear drop (more than 50%) only when the Cu^{2+} concentration was raised to 100 mM (Fig. 2B). These results indicate a possible relationship between the polyphos-

phate level and the adaptation of *A. ferrooxidans* to growth in the presence of Cu^{2+} .

Polyphosphate levels in cells shifted to medium containing copper ions. To determine the effect of Cu^{2+} in unadapted *A. ferrooxidans*, cells grown under P_i -sufficient conditions to accumulate a large amount of polyphosphate (Fig. 2B) were shifted to the presence of 20 mM Cu^{2+} , and the polyphosphate levels were determined at different times postshift (Fig. 3A). A great decrease in polyphosphate levels was seen after 1 h, reaching the lowest level by 4 h, with polyphosphate rapidly dropping to about 20% of the level at time zero, and this level was maintained for the next 24 h. These results indicate that the presence of Cu^{2+} ions affects the polyphosphate content in *A. ferrooxidans*, probably by stimulating the degradation of this polymer, as has been suggested in other systems (1, 14).

Effect of copper ions on PPX activity. A possible mechanism to explain the observed decrease in polyphosphate levels when cells were exposed to copper ions is an increase in PPX activity. Previously, a putative *ppx* gene was found in *A. ferrooxidans* which appears to be part of a Pho regulon in this microorganism (30). To investigate the effect of copper ions on PPX activity, nonadapted cells were shifted to the presence of 20 mM CuSO_4 . Cells were collected at different time intervals, and PPX activity was measured in the cell extracts. A rapid increase in PPX activity was seen when cells were exposed to copper (Fig. 3B). This increase corresponded in time to the decrease in polyphosphate levels seen in Fig. 3A, strongly suggesting that the decrease in polyphosphate levels observed

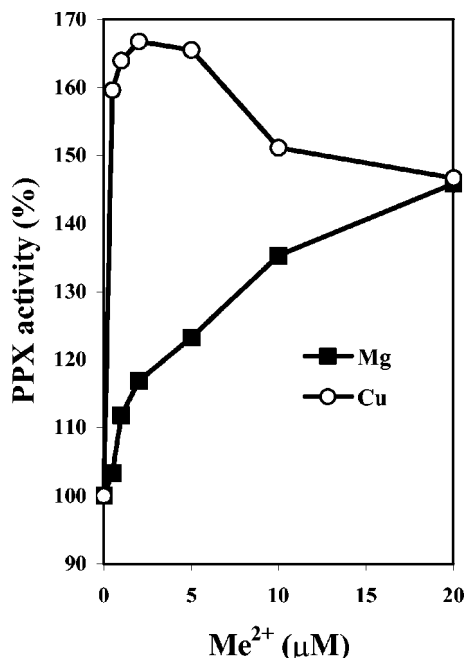


FIG. 4. PPX response to divalent cations *in vitro*. PPX activity was determined in the standard assay with cell extracts from *A. ferrooxidans* grown in the absence of copper. The indicated amounts of MgSO₄ or CuSO₄ were added. The enzyme activity in the absence of added metal was set at 100%.

when cells were shifted to the presence of copper could be due to an increase in this exopolyphosphatase activity.

To further analyze the previous phenomenon, the *in vitro* effect of copper and other metals on the PPX activity present in cell extracts of nonadapted *A. ferrooxidans* was determined, as shown in Fig. 4. It is clear that copper greatly stimulated PPX activity at very low concentrations. The same maximal PPX activity was reached with both copper and magnesium. However, the metal concentration required for this maximal activity was 1 to 2 µM for copper (Fig. 4) and 1,000 µM for magnesium (results not shown). At concentrations of copper higher than 5 µM, there was an inhibition of the activity, whereas magnesium continued to stimulate PPX. Cadmium and zinc also stimulated PPX activity at concentrations of 1 to 2 µM. However, this effect was only half that seen with copper (not shown). These results suggest an effect of the heavy metals on PPX activity and polyphosphate hydrolysis.

Effect of copper ions on efflux of P_i from *A. ferrooxidans* cells.

The decrease in polyphosphate levels due to increased PPX activity in *A. ferrooxidans* cells subjected to Cu²⁺ should generate free P_i. To evaluate if this P_i was transported out of the cells, *A. ferrooxidans* was grown to the late exponential phase and labeled *in vivo* with ³²Pi as indicated in Materials and Methods. After exhaustive washing of the radioactively labeled cells to eliminate nonincorporated label, they were resuspended in sulfur medium with or without Cu²⁺ ions, and P_i efflux was determined, as shown in Fig. 5. There was a continuous basal release of label from the control cells. However, an increase in the P_i efflux over this basal level was observed when cells were exposed to Cu²⁺ ions. The amount of ³²Pi released into the medium was higher in cells exposed to 50 mM Cu²⁺.

By analyzing the radioactivity released into the medium by thin-layer chromatography on polyethyleneimine-cellulose, it was found that it consisted mainly of P_i (not shown). No other radioactively labeled cellular metabolites appeared on the thin-layer chromatograph, indicating that the label released into the medium was not the product of cellular lysis during treatment with copper. These results strongly suggest that at least part of the P_i generated from polyphosphate hydrolysis in the presence of Cu²⁺ ions was transported out of the cells.

In silico search for P_i transporters in *A. ferrooxidans*. The proposed model for metal ion detoxification based on the hydrolysis of polyphosphate involves the transport of metal-phosphate complexes out of the cell. It has been proposed that the inorganic phosphate transport (Pit) system is a candidate for this purpose because it can reversibly transport metal-phosphate complexes (14, 29). P_i transporters have not been described in *A. ferrooxidans*, although the bacterium possesses a putative Pho regulon (30). A Pit-like phosphate transport system was searched for in the available genome of *A. ferrooxidans* ATCC 23270. A Pit-like transporter was not found in this bacterium, but instead we found an open reading frame coding for a protein similar to the Pho84 P_i transporter from *S. cerevisiae*. An alignment of the predicted amino acid sequences of the putative Pho84-like protein of *A. ferrooxidans* with Pho84 from *S. cerevisiae* showed 26% identity and 43% similarity, with the same number of highly conserved transmembrane segments (results not shown). On the other hand, experimental evidence indicates that yeast Pho84, like Pit, transports metal-phosphate complexes (22).

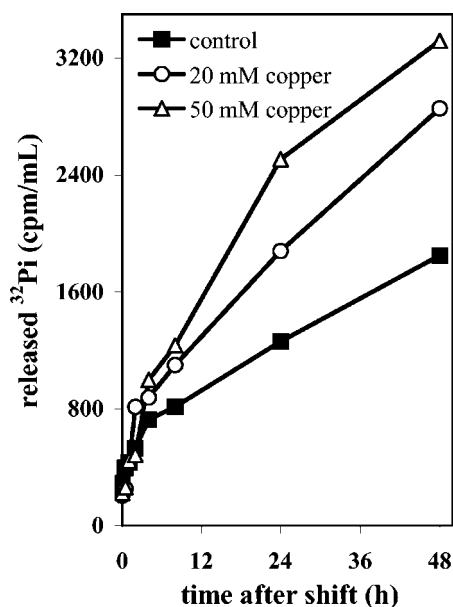


FIG. 5. Effect of copper ions on the efflux of P_i from *A. ferrooxidans* cells. *A. ferrooxidans* was grown in sulfur medium with 1.75 mM P_i to the exponential phase. These cells were then labeled *in vivo* with H₃³²PO₄ (100 µCi/ml) for 17 h in the presence of 0.18 mM P_i, as indicated in Materials and Methods. After the cells were exhaustively washed with unlabeled standard medium, they were shifted to the same fresh medium containing the indicated concentrations of CuSO₄. At the times indicated, the cells were removed by centrifugation, and the radioactive P_i released into the supernatants was determined.

Pho84 and Pho89 are the major P_i transporters in *S. cerevisiae*. Pho84, like Pit, belongs to the family of $P_i:H^+$ symporters and is a member of the major facilitator superfamily (20). The Pho84 transporter is functional only in acidic environmental conditions (22). Although currently there is no experimental evidence for a Pho84-like transporter in *A. ferrooxidans*, it is remarkable that this microorganism, an acidophilic bacterium, possesses a putative P_i transporter of this kind. In this regard, genes encoding proteins similar to Pho84 but not to Pit (or Pho89) were also found in the genomes of other acidophilic microorganisms, such as *Sulfolobus tokodaii*, *Sulfolobus solfataricus*, *Thermoplasma acidophilum*, *Thermoplasma volcanicum*, and *Ferroplasma acidarmanus* (results not shown).

Finally, the *A. ferrooxidans* ATCC 23270 genome sequence also shows the presence of a putative CopA uptake Cu^+ P-type ATPase and the CopB efflux Cu^+ P-type ATPase present in other bacteria (4). This suggests that *A. ferrooxidans* might have a copper homeostasis mechanism similar to that of other microorganisms, but no experimental evidence supporting this proposal is available for this bacterium. Irrespective of the existence of such metal cation uptake and efflux mechanisms, it is plausible that a polyphosphate-mediated metal tolerance mechanism such as the one described here is also of great functional survival value for this extremophilic microorganism.

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