

Localization and Solubilization of the Iron(III) Reductase of *Geobacter sulfurreducens*

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The iron(III) reductase activity of *Geobacter sulfurreducens* was determined with the electron donor NADH and the artificial electron donor horse heart cytochrome *c*. The highest reduction rates were obtained with Fe(III) complexed by nitrilotriacetic acid as an electron acceptor. Fractionation experiments indicated that no iron(III) reductase activity was present in the cytoplasm, that approximately one-third was found in the periplasmic fraction, and that two-thirds were associated with the membrane fraction. Sucrose gradient separation of the outer and cytoplasmic membranes showed that about 80% of the iron(III) reductase was present in the outer membrane. The iron(III) reductase could be solubilized from the membrane fraction with 0.5 M KCl showing that the iron(III) reductase was weakly bound to the membranes. In addition, solubilization of the iron(III) reductase from whole cells with 0.5 M KCl, without disruption of cells, indicated that the iron(III) reductase is a peripheral protein on the outside of the outer membrane. Redox difference spectra of KCl extracts showed the presence of *c*-type cytochromes which could be oxidized by ferrihydrite. Only one activity band was observed in native polyacrylamide gels stained for the iron(III) reductase activity. Excision of the active band from a preparative gel followed by extraction of the proteins and sodium dodecyl sulfate-polyacrylamide gel electrophoresis revealed the presence of high-molecular-mass, cytochrome-containing proteins in this iron(III) reductase activity band. From these experimental data it can be hypothesized that the iron(III) reductase of *G. sulfurreducens* is a peripheral outer membrane protein that might contain a *c*-type cytochrome.

Dissimilatory microbial iron(III) reduction has been proposed as the most important chemical change that takes place in the development of certain anaerobic soils and sediments (26). Iron(III) reduction may have been the first globally significant mechanism for the oxidation of organic matter to carbon dioxide (17). Several studies have demonstrated that iron(III) reduction accounts for the oxidation of up to 65% of organic matter in anaerobic sediments (7, 17). Microbial oxidation coupled with iron(III) reduction is an important natural mechanism for contaminant removal from groundwater or shallow aquifers polluted with landfill leachates (1). Furthermore, iron(III) reduction can influence the biogeochemical cycles of important nutrients, such as carbon, sulfur, and phosphorus (17, 28). Up to now, numerous iron(III)-reducing bacteria have been described in the literature (8, 16). These bacteria are versatile towards electron donors and acceptors which can be used for growth.

Although iron(III)-reducing bacteria have been studied for many years, little is known about the biochemistry and the mechanism of iron(III) reduction. Iron(III)-reducing bacteria are able to utilize Fe(III) either chelated or in insoluble inorganic minerals such as ferrihydrite, goethite, and others. The question arises whether Fe(III) is taken up and reduced in the periplasm or cytoplasm or whether electrons are transferred to Fe(III) minerals outside of the cells. Evidence has been presented that a direct contact between the cells and the metal oxide is necessary for the conservation of energy (3). Iron(III) reductase activity was found to be predominantly in the membranes of the iron(III)-reducing bacteria (23). In the case of *Shewanella putrefaciens*, 54 to 56% of this activity was localized

in the outer membrane (23). The involvement of *b*- and *c*-type cytochromes as electron carriers in the iron(III) respiration chain has been reported previously (23). In addition, it has been shown that the outer membranes of anaerobically grown cells of *S. putrefaciens* have a high cytochrome content (22). This suggests that those cytochromes localized in the outer membrane may play a key role in iron(III) reduction. In this study, we report investigations on the localization of the iron(III) reductase activity of *Geobacter sulfurreducens* with a new enzyme assay. Results obtained by isolating the outer membrane and by treating whole cells with a high KCl concentration to solubilize the iron(III) reductase indicated that the iron(III) reductase of *G. sulfurreducens* is a peripheral protein of the outer membrane. Evidence is presented that a *c*-type cytochrome is involved in iron(III) reduction.

MATERIALS AND METHODS

Growth conditions. *G. sulfurreducens* was cultivated anaerobically in 1-liter bottles filled with 0.7 liter of defined medium (5). One liter of basal medium contained the following: 2.5 g of NaHCO₃, 1.5 g of NH₄Cl, 0.68 g of NaH₂PO₄, 0.1 g of KCl, 6.8 g of CH₃COONa, 13.7 g of Fe(III)-citrate, 10 ml of vitamin solution, and 10 ml of trace element solution, whose composition was as described previously (19). The pH was adjusted to 6.8. The bottles were sealed with butyl rubber stoppers, and the headspace was replaced with N₂-CO₂ (80%-20% [vol/vol]). Cultures were incubated in the dark for 4 days without shaking at 30°C.

Cell harvest and preparation of subcellular fractions. All steps were carried out in an anaerobic glove box (Coy Laboratory, Ann Arbor, Mich.). If not mentioned otherwise, N₂-saturated Tris buffer (100 mM, pH 8.1), containing 0.1 mM dithiothreitol, was used. *G. sulfurreducens* cultures in the late-growth phase were acidified to pH 5 to dissolve iron(II) precipitates, harvested by centrifugation (10,000 × g, 20 min, 4°C), and washed three times with Tris buffer. To prepare cellular fractions, a protocol adapted from a method described by Myers and Myers (22) was used. For the preparation of spheroplasts, cells (110 mg of protein) were resuspended in 30 ml of the Tris buffer containing 25% (wt/vol) sucrose. To accomplish cell wall lysis, a 1/10 volume of a lysozyme solution (6.4 mg · ml⁻¹) and a 1/10 volume of a Na₂-EDTA solution (20 mg · ml⁻¹) were added with constant stirring at 15-min intervals. Finally, MgCl₂ (0.1 M) was added to a final concentration of 12.8 mM. Separation of spheroplasts from the periplasmic fraction was obtained by centrifugation at 20,000 × g for 30 min.

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TABLE 1. Electron donors tested for iron(III) reductase activity^a

Electron donor	Concn (mM)	Chemical reduction of Fe(III)	Iron(III) reductase activity (mU/mg of protein)
NADH	1	No	35
NADPH	1	No	0
Benzyl viologen	0.5	Yes	ND
Methyl viologen	0.5	Yes	ND
Phenosafranin	0.5	Yes	ND
Safranin T	0.5	Yes	ND
Neutral red	0.5	Yes	ND
Bromophenol blue	0.5	No	0
Horse heart cytochrome <i>c</i>	0.004	No	30

^a Reactions were performed in 50 mM HEPES buffer (pH 7) containing 0.3 mM Fe(III)-NTA and 0.5 mM ferrozine. An activity of 1 mU corresponds to 1 nmol of Fe(II) formed per min. ND, not determined.

Spheroplasts (pellet) were resuspended in Tris buffer to a final protein concentration of 9 mg/ml.

To obtain the membrane and soluble fraction, a few crystals of DNase were added to an EDTA-lysozyme-treated cell suspension, and the cells were broken by ultrasonic treatment (10 times, 100 W, 30 s) with a sonifier (model 250; Branson Ultrasonics, Danbury, Conn.) in an ice-water bath. Cell debris was removed by centrifugation (4,000 × g, 10 min, 4°C). The crude extract (supernatant) was centrifuged at 200,000 × g (1 h, 4°C), yielding the soluble fraction containing the cytoplasmic and periplasmic fractions and the membrane fraction (pellet). The latter was resuspended in Tris buffer to a final concentration of 15 mg/ml. The protein concentration was determined by using bicinchoninic acid as described previously (30), with bovine serum albumin as a standard.

Separation of outer and cytoplasmic membranes. Separation of outer and cytoplasmic membranes was achieved according to a procedure adapted from Myers and Myers (22). An aliquot of 1 ml of the membrane fraction containing 1.585 mg of protein was layered onto 10.5 ml of a 50 to 80% (wt/wt) sucrose gradient and centrifuged for 17 h at 82,500 × g. Two colored discrete bands were observed, one between 5 and 8 cm from the bottom of the tube (low-density band) and one on the bottom of the tube (high-density band). The bands were retrieved from the tube by removing fractions of 0.5 to 1 ml with a syringe. The fractions were resuspended in and subsequently dialyzed for 5 h at 4°C against a 10 mM HEPES buffer containing 0.1 mM dithioerythritol (DTE). To identify the fraction containing the outer membrane, the content of KDO (2-keto-3-deoxyoctonate), a specific constituent of the lipopolysaccharide in the outer membrane, was determined (13).

Enzyme assay. Iron(III) reductase activity was assayed anaerobically in 1-cm cuvettes by photometric monitoring of the appearance of Fe(II) over time as adapted from the method of Lascelles and Burke (15). The reaction mixture contained either 1 mM NADH or 4 μM dithionite-reduced horse heart cytochrome *c* as the electron donor, 0.5 mM ferrozine as the Fe(II) chelating agent, and either 0.15 mM Fe(III)-nitriloacetic acid (NTA), Fe(III)-citrate, or Fe(III)-EDTA as electron acceptor in N₂-saturated 50 mM HEPES buffer (pH 7). Horse heart cytochrome *c* was added from a stock solution that was prepared by adding 50 μl of a 4.6 mM sodium dithionite solution to 1 ml of 50 mM HEPES buffer (pH 7) containing 2 mg of horse heart cytochrome *c*. The concentration of reduced horse heart cytochrome *c* in the sample cuvette was calculated by measuring the increase in A_{552} ($A_{552} = 29,500 \text{ M}^{-1} \cdot \text{cm}^{-1}$). The enzyme assay was started by adding whole cells or cell extracts of *G. sulfurreducens* (0.03 to 0.5 mg of protein) to both sample and reference cuvettes. The activity was monitored by measuring the increase in A_{562} (32) due to the formation of a Fe(II)-ferrozine complex ($A_{562} = 28,000 \text{ M}^{-1} \cdot \text{cm}^{-1}$) against a reference where no electron donor was added. An activity of 1 mU corresponded to 1 nmol of Fe(II) formed per min. In the absence of ferrozine, the enzyme activity could be also monitored by measuring the decrease in A_{552} due to the reoxidation of the reduced horse heart cytochrome *c* as previously described ($A_{552} = 29,500 \text{ M}^{-1} \cdot \text{cm}^{-1}$) (12).

UV-visible spectroscopy. UV-visible spectra of KCl-extracted proteins were recorded on a U-2000 spectrophotometer (Hitachi, Tokyo, Japan) in a 1-cm quartz cuvette containing 0.1 to 0.3 mg of protein/ml. To record the redox difference spectra of KCl extracts, the protein solution was reduced with sodium dithionite and subsequently reoxidized by adding increasing amounts of an anaerobic solution of ferrihydrite to the reaction mixture in order to obtain final concentrations ranging from 5.4 to 13.7 mM.

Extraction of iron(III) reductase. Membrane fractions, crude extracts or whole cells were stirred for 1 h at 4°C in a high-ionic-strength salt buffer (100 mM Tris, 0.1 mM DTE, 0.5 M KCl [pH 7.6]) in order to release the iron(III) reductase.

Native polyacrylamide gel electrophoresis (PAGE) stained for the iron(III) reductase. For detection of enzymatic reduction of Fe(III) in the presence of ferrozine in native polyacrylamide gels, a procedure adapted from Moody and Dailey (21) was used. An aliquot of 40 μl of the KCl-extracted proteins contain-

ing 950 μg of protein after dialysis and concentration (Amicon ultrafiltration cell with PM-10 filter) was applied to a 1-mm, 12% Ready gel (Bio-Rad). The gel was run for 3 h at 10 mA. The gel was removed and washed for 1 min in 200 ml of 0.1 mM Tris buffer (pH 7.6). After the washing buffer was removed, the gel was placed in 170 ml of the same buffer containing 30 mg of NADH, 2 mg of reduced horse heart cytochrome *c*, and 107 mg of ferrozine. The reaction was started by adding 150 μl of a 300 mM Fe(III)-NTA solution. A purple band was observed where iron(III) reduction took place after 30 min of incubation at ambient temperature. Photographs could then be taken.

To isolate the protein of the activity band, a preparative gel was performed according to the following procedure. The dialyzed and concentrated KCl-extracted proteins were applied on the 8 lanes of the 1-mm, 12% Ready gel. To avoid disturbance due to iron for further processing, one lane was separated from the others by cutting the gel and then was stained for the iron(III) reductase activity. With this procedure, the position of the active band of the other lanes could be estimated and the bands could be excised. These bands were suspended in 5 ml of 50 mM HEPES (pH 7) containing 0.1 mM DTE and homogenized on ice with a Potter. Sodium dodecyl sulfate (SDS) was added to a final concentration of 1%. The solution was incubated for 30 min at 50°C and then for 90 min at 37°C. The sample was centrifuged for 5 min at 12,000 × g. The proteins of the supernatant were precipitated by adding 11 ml of ethanol to 4 ml of supernatant followed by incubation at -20°C for 2 days. Precipitated proteins were collected by centrifugation at 12,000 × g for 10 min and then resuspended in 100 μl of H₂O.

Denaturing SDS-PAGE analysis of KCl-extracted proteins. SDS-PAGE was carried out as described by Laemmli (14) after the samples were desalted by dialysis. Proteins were visualized by a silver-staining procedure (27), and the cytochromes were stained by a method with *o*-dianisidine and hydrogen peroxide (10).

Chemicals. Ferrozine, DNase I, horse heart cytochrome *c*, NADPH, safranin T, phenosafranin, DTE, and *o*-dianisidine-HCl tablets were purchased from Fluka (Buchs, Switzerland); NADH, neutral red, and bromophenol blue were obtained from Merck (Darmstadt, Germany); and benzyl viologen, methyl viologen, rotenone, and 2-*n*-heptyl-4-hydroxyquinoline *N*-oxide (HOQNO) were from Sigma (Munich, Germany).

RESULTS

Iron(III) reductase activity. The iron(III) reductase activity of *G. sulfurreducens* could be measured with NADH but not with NADPH (Table 1). If Fe(III) NTA was used as an electron acceptor, the highest Fe(III)-reducing activities were measured (10.6 mU/mg of protein), whereas with Fe(III)-citrate and Fe(III)-EDTA, lower (2.1 mU/mg of protein) and no activities were detected, respectively. Cell extracts that were heated at 100°C for 5 min showed no iron(III) reductase activity. Exposure to air for 2 h decreased the activity by 80%, indicating an oxygen lability of the iron(III) reductase activity. The

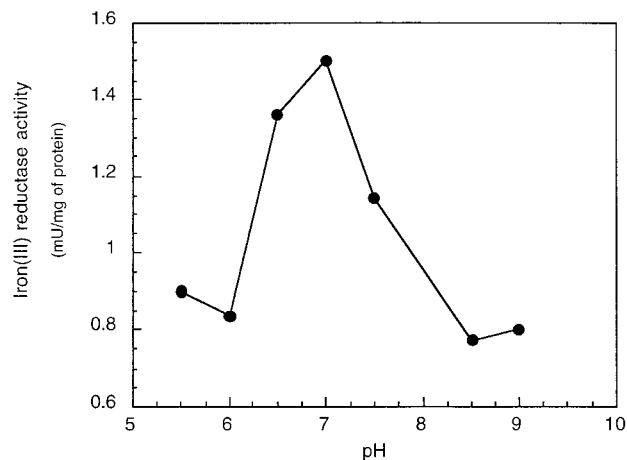


FIG. 1. Iron(III) reductase activity as a function of pH. The assay was done in a solution containing HEPES, MES (morpholineethane sulfonic acid), PIPES [piperazine-*N,N'*-bis(2-ethanesulfonic acid)], TAPS [*N*-[Tris-(hydroxymethyl)-methyl]-3-amino-propane sulfonic acid], and KH₂PO₄ buffer, each constituent at a concentration of 20 mM. In addition, the reaction mixture contained 0.3 mM Fe(III)-NTA, 0.5 mM ferrozine, and 1 mM NADH.

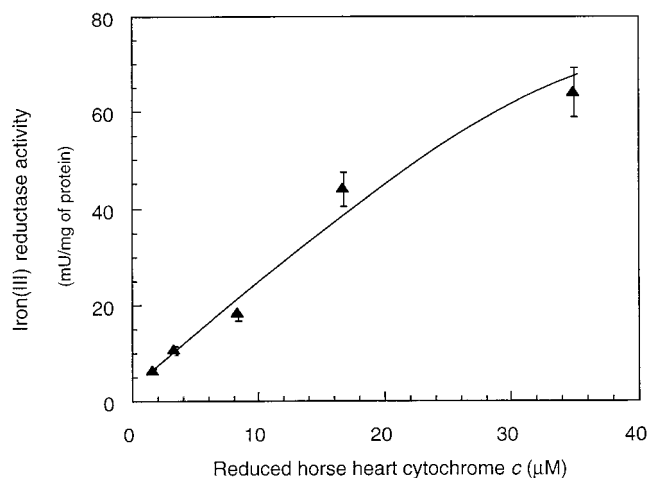


FIG. 2. Iron(III) reductase activity as a function of reduced horse heart cytochrome *c* concentration. The assay was done in 50 mM HEPES buffer (pH 7) containing 0.3 mM Fe(III)-NTA and 0.5 mM ferrozine.

activity was highest at pH 7 (Fig. 1). In order to develop an NADH-independent assay, different artificial electron donors were tested (Table 1). The major difficulty was the rapid chemical reduction of Fe(III) by these artificial electron donors, which did not allow us to distinguish between an enzymatic and chemical reduction. Also, reducing agents such as cysteine, dithionite, and Ti(III) citrate could not be used to reduce the iron(III) reductase since they also reduced Fe(III) chemically. Only horse heart cytochrome *c* reduced by dithionite could be used as an alternative electron donor (Table 1). The iron(III) reductase activity increased with increasing concentrations of reduced horse heart cytochrome *c* (Fig. 2). However, at concentrations above 35 μM , activity measurements with ferrozine were not possible due to a background absorption by reduced horse heart cytochrome *c* at 562 nm that was too high. The iron(III) reductase activity could alternatively be determined by measuring the oxidation of reduced horse heart cytochrome *c* at 552 nm. In this case, due to the high absorbance of the reduced horse heart cytochrome *c*, it was not possible to use concentrations above 55 μM , and the highest iron(III) reductase activity measured was of 75 mU/mg of protein.

Iron(III) reductase activity was not affected by 50 μM

HOQNO and was partially inhibited (50 to 80%) by cyanide (KCN; 200 μM) with both NADH and horse heart cytochrome *c* as the electron donor. Rotenone completely inhibited the iron(III) reductase activity only with NADH.

Localization of the iron(III) reductase. Breakage of the cells of *G. sulfurreducens* by EDTA-lysozyme treatment followed by sonication led to a 1.5-fold increase of iron(III) reductase activity in comparison with unbroken cells when NADH was used as an electron donor (Table 2). No significant increase caused by breaking the cells was observed when horse heart cytochrome *c* was used as the electron donor.

The spheroplast fraction contained 77 and 68% of iron(III) reductase activity, respectively, when horse heart cytochrome *c* and NADH were used as electron donors, whereas the periplasmic fractions contained 23 and 32% of the iron(III) reductase activity, respectively (Table 2). The soluble fraction of crude extracts accounted for approximately the same percentage of iron(III) reductase activity as the periplasmic fraction of the spheroplast preparation (17 and 33%, respectively). The membrane fraction contained 83 and 67% of the iron(III) reductase activity, respectively.

Separation of the outer and cytoplasmic membranes was achieved by using a highly concentrated sucrose gradient of 50 to 80%. The outer membrane, identified by its high KDO content of 77 ± 5 ng/mg of protein, contained 75 and 79% of the iron(III) reductase measured with horse heart cytochrome *c* and NADH, respectively (Table 3). The cytoplasmic fraction had a low KDO content of 2.2 ± 0.2 ng/mg of protein and contained 25 and 21% of the iron(III) reductase activity, respectively (Table 3).

Solubilization of the iron(III) reductase. The activity present in the soluble fractions might have been due to iron(III) reductase that was only loosely associated with the membranes. This hypothesis was substantiated by the finding that the iron(III) reductase activity can be solubilized by incubation of the membrane fraction with KCl 0.5 M (Table 4). When horse heart cytochrome *c* was used as the electron donor, an iron(III) reductase activity of 45.2 ± 5 mU was found in the soluble fraction that corresponds to 188% activity compared to the membrane fraction before treatment (Table 4). With NADH as the electron donor, only 14% of the iron(III) reductase activity was present in the soluble fraction. The results obtained after solubilization with KCl also showed that with NADH as electron donor 59% less iron(III) reductase activity

TABLE 2. Localization of iron(III) reductase in different cellular fractions of *G. sulfurreducens*

Fraction	Protein (mg)	Iron(III) reductase activity ^a with:					
		NADH			Horse heart cytochrome <i>c</i>		
		Sp act (mU/mg of protein)	Total activity (mU)	%	Sp act (mU/mg of protein)	Total activity (mU)	%
Whole cells vs. crude extract							
Whole cells	110 \pm 7	4.8 \pm 0.2	528 \pm 20	100	7.0 \pm 0.7	770 \pm 70	100
Crude extract	114 \pm 10	7.7 \pm 0.5	878 \pm 50	166	7.4 \pm 0.5	844 \pm 50	110
Periplasm vs. spheroplasts							
Periplasm	1.7 \pm 0.2	9 \pm 1	15.3 \pm 2	23	9.5 \pm 1	16.2 \pm 2	32
Spheroplasts	8.7 \pm 1.0	6.0 \pm 1	52.0 \pm 9	77	4.0 \pm 0.5	35.0 \pm 4	68
Soluble fraction vs. membrane fraction							
Soluble fraction	7.5 \pm 1	9.7 \pm 0.5	72.7 \pm 5	17	11.0 \pm 1	82 \pm 7.0	33
Membrane fraction	60.3 \pm 0.5	6.0 \pm 1	362 \pm 6	83	2.7 \pm 0.5	163 \pm 3	67

^a Reactions were performed in 50 mM HEPES buffer (pH 7) containing 0.3 mM Fe(III)-NTA, 0.5 mM ferrozine, and either 1 mM NADH or 4 μM horse heart cytochrome *c*. An activity of 1 mU corresponds to 1 nmol of Fe(II) formed per min.

TABLE 3. Localization of the iron(III) reductase in the outer membrane fraction of *G. sulfurreducens*

Fraction	Protein (mg)	KDO content		Iron(III) reductase activity ^a with:			
		Total (mg)	Specific (ng/mg of protein)	NADH		Horse heart cytochrome <i>c</i>	
				Sp act (mU/mg of protein)	Total activity (mU)	Sp act (mU/mg of protein)	Total activity (mU)
Low-density band	0.85 ± 0.04	2.60 ± 0.04	77 ± 5	112 ± 12	95.2 ± 10	122 ± 15	103 ± 12
High-density band	1.02 ± 0.06	0.20 ± 0.02	2.2 ± 0.2	25 ± 4	25.5 ± 4	33 ± 3	33.6 ± 3

^a Reactions were performed in 50 mM HEPES buffer (pH 7) containing 0.3 mM Fe(III)-NTA, 0.5 mM ferrozine, and either 1 mM NADH or 4 μM horse heart cytochrome *c*. An activity of 1 mU corresponds to 1 nmol of Fe(II) formed per min.

was recovered compared with untreated membrane fraction, whereas with horse heart cytochrome *c* an increase of total activity by a factor of 2.6 was found.

The incubation of whole cells in Tris buffer containing 0.5 M KCl did not disrupt the cells, as verified by microscopy, and led to a solubilization of the iron(III) reductase activity. The soluble fraction obtained with the KCl treatment contained 54% of the iron(III) reductase activity (measured with horse heart cytochrome *c*), whereas no iron(III) reductase activity was present in the supernatant in the absence of KCl (Table 4). With NADH as electron donor, 34% of the iron(III) reductase activity was solubilized by KCl. The chelating agent EDTA, known to destabilize the outer membrane by complexing Ca²⁺ and Mg²⁺ ions present in the outer membrane, caused in combination with KCl a solubilization of up to 81% of the iron(III) reductase activity (Table 5).

Cytochromes and iron(III) reductase activity in KCl extracts. The UV-visible spectrum of the KCl extract obtained from whole cells exhibited a strong peak at 410 nm and shoulders centered at 520 and 580 nm (data not shown). These features are characteristic of hemoproteins. Redox difference spectra of this sample exhibited a Soret band at 420 nm and β and α bands at 522 and 552 nm, respectively, all of which are characteristic of *c*-type cytochromes. The addition of ferrihydrite to the cuvette containing the dithionite-reduced cytochrome *c* led to the disappearance of this spectrum (Fig. 3). This oxidation suggested that *c*-type cytochromes are involved in iron(III) reduction. Silver- and heme-stained SDS gels of the KCl fraction containing the iron(III) reductase revealed the presence of numerous proteins and several cytochromes (Fig. 4). Native PAGE of the KCl extract stained for the iron(III) reductase activity exhibited one purple band in the upper part of the gel, indicating that the iron(III) reductase could be a high-molecular-mass protein (data not shown). Analyzing the proteins obtained from a native preparative gel by SDS-PAGE revealed the presence of three cytochrome-containing proteins with molecular masses ranging from 67 to 97 kDa (Fig. 4).

DISCUSSION

The iron(III) reductase activity assay with ferrozine and Fe(III) complexed with NTA and citrate, respectively, showed that the reduction of Fe(III)-NTA occurred five times faster than the reduction of Fe(III)-citrate. This result is in agreement with previous findings obtained with *S. putrefaciens* (2). The Fe(II) species formed with citrate may be kinetically less labile than those formed with NTA due to steric effects (9). Additional evidence for the effect of kinetic lability was obtained with Fe(III) complexed with the hexadentate ligand EDTA. In this case, no iron(III) reductase activity could be measured in the assay, which was probably caused by the inability of ferrozine to unbind Fe(II) from the Fe(II)-EDTA complex. A similar finding has been described by Dobbin et al. (9). However, Fe(III)-EDTA can act as an electron acceptor for the iron(III)-reducing bacteria as shown by the stimulation of benzene degradation by iron(III) reducers (20).

The iron(III) reductase activity of *G. sulfurreducens* could be measured with NADH as a physiological electron donor but not with NADPH. The same was also reported for *S. putrefaciens* (23). Since iron(III) reduction by *G. sulfurreducens* is a respiratory process, electrons derived from NADH oxidation are probably not directly transferred to Fe(III) but rather are transferred via a respiratory chain. This assumption is supported by the inhibition of NADH-dependent iron(III) reduction by rotenone that was also found for *Geobacter metallireducens* (11). In addition, the increase of activity after the cells were broken that gave NADH better access to the NADH dehydrogenase and the loss of activity upon solubilization of the iron(III) reductase by KCl that disrupted the respiratory chain also provided evidence for this assumption.

Here, for the first time, an artificial iron(III) reductase activity assay could be set up by using reduced horse heart cytochrome *c* as the electron donor. The electrons of the latter probably enter the respiratory chain to the iron(III) reductase on a different site than the electrons of NADH for two reasons. First, in contrast to NADH, there was no increase in activity

TABLE 4. Solubilization of iron(III) reductase from membrane fractions of *G. sulfurreducens*

Fraction	Protein (mg)	Iron(III) reductase activity ^a with:					
		NADH			Horse heart cytochrome <i>c</i>		
		Sp act (mU/mg of protein)	Total activity (mU)	%	Sp act (mU/mg of protein)	Total activity (mU)	%
Membrane fraction before extraction	6.0 ± 1.0	6.0 ± 1.0	36.0 ± 6.0	100	4.0 ± 0.5	24.0 ± 3.0	100
Soluble fraction after extraction	3.6 ± 0.2	1.43 ± 0.2	5.1 ± 0.6	14	12.5 ± 1.0	45.2 ± 5.0	188
Membrane fraction after extraction	2.1 ± 0.3	4.5 ± 0.5	9.5 ± 1.0	27	7.9 ± 1.0	17.0 ± 2.0	71

^a Reactions were performed in 50 mM HEPES buffer (pH 7) containing 0.3 mM Fe(III)-NTA, 0.5 mM ferrozine, and either 1 mM NADH or 4 μM horse heart cytochrome *c*. An activity of 1 mU corresponds to 1 nmol of Fe(II) formed per min.

TABLE 5. Solubilization of iron(III) reductase from whole cells of *G. sulfurreducens*^a

Addition(s) ^b	Fraction	Protein (mg)	Iron(III) reductase activity ^c with:					
			NADH			Horse heart cytochrome <i>c</i>		
			Sp act (mU/mg of protein)	Total activity (mU)	Soluble fraction/cell ratio (%)	Sp act (mU/mg of protein)	Total activity (mU)	Soluble fraction/cell ratio (%)
None	Soluble fraction	2.0 ± 0.2	0	0	0	0	0	0
	Cells	2.2 ± 0.1	16.6 ± 1.0	36.5 ± 2.0	100	17.5 ± 3.0	38.5 ± 5.0	100
KCl	Soluble fraction	3.6 ± 0.2	2.1 ± 0.1	7.6 ± 0.2	34	4.4 ± 0.4	15.8 ± 1.0	54
	Cells	2.6 ± 0.1	5.7 ± 0.5	14.8 ± 2.0	66	5.1 ± 0.5	14.0 ± 2.0	46
EDTA	Soluble fraction	3.4 ± 0	0	0	0	0	0	0
	Cells	2.8 ± 0.4	11.7 ± 1.5	32.8 ± 4.0	100	9.5 ± 0.5	26.6 ± 1.0	100
KCl + EDTA	Soluble fraction	3.3 ± 0.2	6.4 ± 0.5	21.1 ± 1.0	66	11.7 ± 2.0	38.6 ± 4.0	81
	Cells	2.6 ± 0.1	4.2 ± 0.2	10.9 ± 0.5	34	3.4 ± 0.4	8.8 ± 1.0	19

^a For solubilization experiments, cells were dissolved in 100 mM Tris buffer (pH 7.8).

^b KCl was added to a final concentration of 0.5 M; EDTA was added to a final concentration of 1 mM.

^c Reactions were performed in 50 mM HEPES buffer (pH 7) containing 0.3 mM Fe(III)-NTA, 0.5 mM ferrozine, and either 1 mM NADH or 4 μM horse heart cytochrome *c*. An activity of 1 mU corresponds to 1 nmol of Fe(II) formed per min.

with horse heart cytochrome *c* after the cells were broken. Second, disruption of the respiratory chain by solubilizing the iron(III) reductase did not result in a loss of activity as was found with NADH but rather an increased iron(III) reductase activity. No indications for a different site of electron transfer were obtained by the inhibitor experiments. The reactions with either NADH or horse heart cytochrome *c* showed the same inhibition pattern. Interestingly, the inhibition pattern of *G. sulfurreducens* was different from the patterns described for either *S. putrefaciens* (9, 23) or *G. metallireducens* (11). HOQNO had no effect on the iron(III) reductase activity of the two *Geobacter* species, but it did inhibit the iron(III) reduction by *S. putrefaciens*. Cyanide, on the other hand, inhibited the iron(III) reduction by *S. putrefaciens* and *G. sulfurreducens*, whereas no effect was observed for *G. metallireducens*.

Similar to other iron(III) reducers, such as *S. putrefaciens* and *G. metallireducens*, where the total iron(III) reductase activity was found exclusively in the membrane fraction (9, 11, 23), the major part of the iron(III) reductase activity of *G. sulfurreducens* was also found in the membrane fraction. The presence of iron(III) reductase activity in the outer membrane fraction has also been found for *S. putrefaciens*. For *G. sulfurreducens*, 75 to 79% of the iron(III) reductase activity of the membrane fraction was present in the outer membrane, whereas for *S. putrefaciens* it was 54 to 56%. The iron(III) reductase activity found in the cytoplasmic membrane fraction of *G. sulfurreducens* could be an artifact due to the sonication treatment since cell breakage by sonication can cause an extensive redistribution of membrane proteins between membranes (31). The minor part of the iron(III) reductase activity recovered in the soluble fractions after ultracentrifugation or preparation of spheroplasts is probably due to the release of the loosely membrane-associated iron(III) reductase during treatment. The solubilization of iron(III) reductase by KCl actually showed that this enzyme activity is loosely membrane associated since a high salt concentration treatment is a well-known method for detaching peripheral membrane proteins (25).

Because the major part of Fe(III) exists in the environment as insoluble Fe(III) oxides, a localization of the terminal reductase in the outer membrane is certainly conceivable. It has been proposed that direct contact of the cell surface with the iron oxides is needed to enable the bacteria to reduce Fe(III) (3, 4, 23). This would require an outer membrane iron(III)

reductase facing the outside of the cells. Indications for such a localization of the iron(III) reductase in *G. sulfurreducens* were obtained with the solubilization of iron(III) reductase activity from whole cells by the KCl treatment. This treatment probably only detached peripheral outer membrane proteins located on the outside of the cells and did not detach proteins facing the periplasm. A recent study, however, showed that cell contact is not a requisite for reduction of solid Fe(III) oxides (6). In this study, an adhesion-deficient *Shewanella alga* strain reduced amorphous Fe(III) oxide at the same rates as a strain that strongly adhered to amorphous Fe(III) oxide. If the electrons are not directly transferred from the iron(III) reducers to the Fe(III) oxides, the electron transfer could be accomplished by diffusion of electron mediators or of free or complexed Fe(III). It has been shown that *G. sulfurreducens* excretes *c*-type cytochromes in the medium and that this cytochrome has iron(III) reductase activity (29). This suggested that this cytochrome could act as an electron mediator between the cells and the iron oxides. Indications for the production of a cell surface protein that efficiently chelate Fe(III) have been obtained for *S. alga* (6).

The presence of heme proteins in the protein fraction iso-

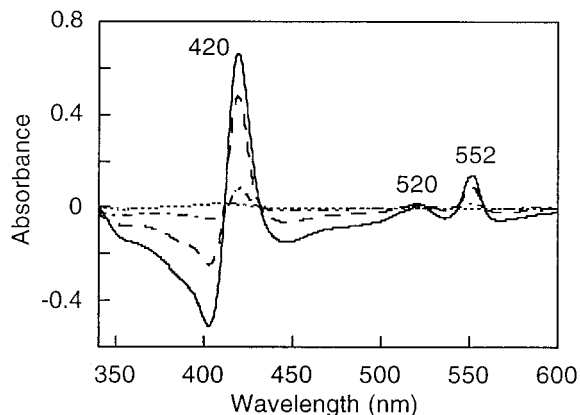


FIG. 3. UV-visible spectra of KCl extracts of whole cells. —, redox difference spectrum, after addition of ferrihydrite; ---, 8.3 mM; - · - ·, 10.8 mM; · · · ·, 13.7 mM.

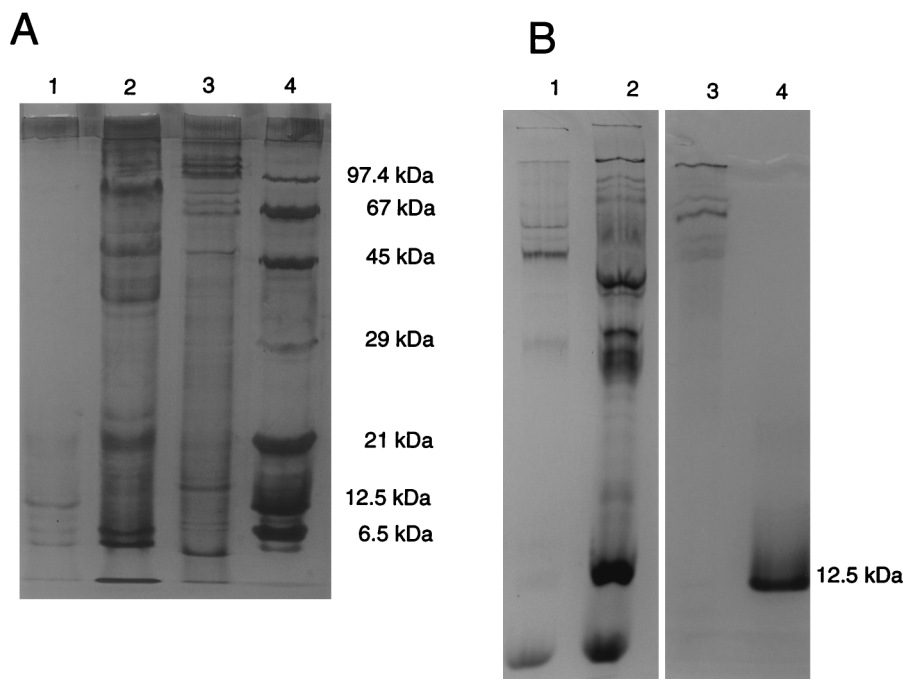


FIG. 4. SDS-PAGE of different fractions of *G. sulfurreducens* extracts. The gels were stained for protein with silver (A) and heme (B). Lanes 1, whole cells after KCl extraction; lanes 2, KCl extract from whole cells; lanes 3, isolated band from a preparative iron(III) reductase activity gel; lanes 4, molecular mass standards including phosphorylase *b* (97.4 kDa), bovine serum albumin (67 kDa), egg albumin (45 kDa), carbonic hydrase (29 kDa), soybean trypsin inhibitor (21 kDa), horse heart cytochrome *c* (12.5 kDa), and trypsin inhibitor (6.5 kDa).

lated from the iron(III) reductase activity band of a native gel, the solubilization of several *c*-type cytochromes by KCl from whole cells of *G. sulfurreducens*, and the possible involvement of these cytochromes in iron(III) reduction added further evidence for the hypothesized key role that periplasmic and outer membrane *c*-type cytochromes play in iron(III) reduction (18, 22, 24, 29). For *S. putrefaciens* it has been shown that the outer membrane of anaerobically grown cells contained, in addition to 54 to 56% iron(III) reductase activity (23), 80% of the membrane-bound cytochromes (22). Recently, the gene of a tetraheme cytochrome *c* of *S. putrefaciens* has been cloned and sequenced (24). This *c*-type cytochrome has been found in the cytoplasmic membrane as well as in the soluble fraction. It has been proposed that the tetraheme cytochrome *c* is involved in the transfer of electrons from the cytoplasmic membrane to acceptor proteins such as fumarate reductase located in the periplasm and iron(III) reductase located in the outer membrane (24). Finally, a *c*-type cytochrome that has been excreted into the medium by *G. sulfurreducens* and that has been purified is similar to the cytochrome *c*₃ of other bacteria (29). This *c*-type cytochrome was also present in the membrane and soluble fraction and was able to reduce different electron acceptors including iron oxides. In contrast to the heme proteins present in the iron(III) reductase activity band that had a high molecular mass of 70 to 100 kDa, the excreted *c*-type cytochrome had a molecular mass of only 9.6 kDa (29). High-molecular-mass cytochromes were present in the outer membrane of *Desulfovibrio vulgaris* (Hildenborough) when cultivated in a medium with a high Fe²⁺ concentration (100 ppm) (33). A role in anaerobic biocorrosion has been postulated for these high-molecular-mass cytochromes. Whether the iron(III) reductase of *G. sulfurreducens* is indeed a high-molecular-mass cytochrome will be investigated in the near future.

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