Redox and Chemical Activities of the Hemes in the Sulfur Oxidation Pathway Enzyme SoxAX*

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Background: SoxAX enzymes initiate microbial oxidation of reduced inorganic sulfur compounds. Their catalytic mechanism is unknown.

Results: Cyanide displaces the CysS⁻ ligand to the active site heme following reduction by S₂O₄²⁻ but not Eu(II).

Conclusion: An active site heme ligand becomes labile on exposure to substrate analogs.

Significance: Elucidation of SoxAX mechanism is necessary to understand a widespread pathway for sulfur compound oxidation.

SoxAX enzymes couple disulfide bond formation to the reduction of cytochrome c in the first step of the phylogenetically widespread Sox microbial sulfur oxidation pathway. Rhodovulum sulfidophilum SoxAX contains three hemes. An electrochemical cell compatible with magnetic circular dichroism at near infrared wavelengths has been developed to resolve redox and chemical properties of the SoxAX enzymes. In combination with potentiometric titrations monitored by electronic absorbance and EPR, this method defines midpoint potentials (Eₘ) at pH 7.0 of approximately +210, −340, and −400 mV for the His/Met, His/His/Cys−, and active site His/Cys−-ligated heme, respectively. Exposing SoxAX to S₂O₄²⁻, a substrate analog with Eₘ ≈ −450 mV, but not Eu(II) complexed with diethylene triamine pentacetic acid (Eₘ ≈ −1140 mV), allows cyanide to displace the cysteine persulfide (CysS⁻) ligand to the active site heme. This provides the first evidence for the dissociation of CysS⁻ that has been proposed as a key event in SoxAX catalysis.

Microbial oxidation of reduced inorganic sulfur compounds forms an integral part of the biogeochemical sulfur cycle. It is exploited in biomining and is the defining metabolic pathway in both photosynthetic and nonphotosynthetic sulfur bacteria where it produces electrons that may be used for CO₂ fixation or fed into respiratory electron transfer chains (1, 2). The best studied and most widely distributed pathway for this process is the Sox² (sulfur oxidizing) pathway that converts thiosulfate and certain other sulfur species to sulfate (1). Intermediates in this pathway are covalently bound to a Cys residue in the SoxYZ protein. SoxYZ uses a swinging arm mechanism to deliver these intermediates to the active sites of the other Sox components (3, 4). SoxAX catalyzes the first step of the pathway in which thiosulfate is oxidatively coupled to the Cys of SoxYZ with the reduction of two molecules of cytochrome (cyt) c. SoxAX enzymes have been classified into three groups (5). Groups 1 and 2 are heterodimers containing three and two c-type hemes, respectively. Group 3 comprises diheme-containing heterotrimers with an additional SoxA subunit. Although details of the mechanism of thiosulfate and SoxY co-oxidation by SoxAX are unclear, it is likely that one of the roles of the hemes is to mediate electron transfer from substrates to cyt c (2).

The SoxAX enzymes of both Rhodovulum sulfidophilum and Paracoccus pantotrophus belong to group 1. In these enzymes, the SoxA subunit binds one heme with His/Met axial ligands, whereas both hemes in the larger SoxA subunit have the highly unusual His/thiolate axial ligand set (6–8). Crystal structures of both proteins reveal an almost linear constellation of hemes that have been labeled Heme-1, -2, and -3 in the convention of Bamford et al. (8). The heme of SoxA (Heme-3) and one heme of SoxA (Heme-2) are positioned close to the dimer interface such that their shortest edge-to-edge separation is of the order of 10 Å. By contrast, the second SoxA heme (Heme-1) is positioned ~24 Å from its neighbor, precluding physiologically relevant rates of intramolecular electron transfer to the other SoxA hemes. In Starkeya novella SoxAX, the most studied of the diheme group 2 and 3 enzymes, the arrangement and ligation of the hemes straddling the dimer interface are conserved (9). In these enzymes, Heme-1 is replaced with a disulfide bond (10), and the role of either group in enzyme activity is unclear.

A solvent channel leads from the protein surface to the distal face of Heme-2, suggesting that this forms part of the active site for thiosulfate oxidation (6, 8, 9). Distal ligation of this heme is provided by a cysteine persulfide (CysS⁻) formed by post-translational modification of Cys-222A (R. sulfidophilum numbering). This modification may arise from decomposition of a catalytic intermediate during enzyme purification.
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Expression and Purification of Recombinant R. sulfidophilum SoxAX—A plasmid was constructed to allow heterologous expression of R. sulfidophilum soxXA in Rhodobacter capsulatus under the control of the dimethyl sulfoxide (DMSO)-inducible dorA promoter. soxX was amplified from R. sulfidophilum chromosomal DNA using primers MKS001 (5'-AGACTGGA-TCCATGCGAGGTATGG-3') and MKS002 (5'-ACC-GGACTAGTCATTGCCGTACG-3'). The amplicon was digested with BamHI and ligated into BamHI- and Smal-digested pGEM-7Zf(+) (Promega) to produce plasmid pMKSO1. soxA, together with the soxX ribosome binding site, were amplified from R. sulfidophilum chromosomal DNA using primers MKS003 (5'-ACTAGTCGTAGGCGCGCTCCGTG-3') and MKS006 (5'-GCTAGATGTCATGCGGCACG-3'). The amplicon was digested with SpeI and XbaI and ligated into the same sites in pMKSO1 to give plasmid pMKSO8. The soXXA genes were excised from plasmid PMKS08 with BamHI and XbaI and cloned into the same sites in pD rex (17) to produce plasmid pMKSO10. The soXXA-containing HindIII-XbaI fragment from pMKSO10 was then cloned into the same sites in pRK415 (18) to give plasmid pMKSO12.

Plasmid pMKSO12 was introduced into R. capsulatus strain 37B4 ΔdorA (19) by conjugation from Escherichia coli strain S17-1 (20). Starter cultures were grown photoheterotrophically at 28 °C on malate-containing RCV medium (21) supplemented with 1 μg ml⁻¹ tetracycline. 80 ml of starter culture at an A₆₀₀nm of 0.3 was used to inoculate 5-liter bottles of RCV-malate medium supplemented with 1 μg ml⁻¹ tetracycline, 60 mM DMSO, and 1 mM sodium molybdate, and the resulting static cultures were incubated under illumination at 28 °C for 5 days. Cells were harvested, a periplasmic fraction was isolated, and the periplasmic fraction from R. capsulatus strain

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tions composed of reagents of AnalR quality or equivalent and water of resistivity >18 megaohms cm.

Potentiometric Titration by Electrode Equilibration—Optically transparent electrochemical cells of three designs were used to poise samples of SoxAX at various potentials for examination by (magneto-) optical spectroscopies. The cells employed three electrodes connected to an Autolab PGSTAT12 potentiostat under the control of the GPES software. The working electrode was selected to optimize the quality of data in terms of (a) spectral signal-to-noise and (b) the time needed for sample equilibration with the applied potential because this limited the number of potentials for which a spectrum could be collected over a practical time period.

Potentiometric titrations monitored by electronic absorbance were performed with a working electrode composed of mesoporous nanocrystalline SnO2 in a previously described cell configuration (23, 24). SoxAX was adsorbed onto SnO2 electrodes during 2 h of cyclic voltammetry (+60 mV to −540 mV, 20 mV s−1) in 35 μM SoxAX, 50 mM Heps, 100 mM NaCl, pH 7, 20 °C. The SoxAX loaded electrode was then rinsed with buffer-electrolyte to remove loosely bound material prior to placing it inside the spectroelectrochemical cell. Taking the extinction coefficient of adsorbed SoxAX to be the same as that of the protein in solution and the path length of the adsorbed sample to be the 4-μm thickness of the SnO2 layer, a surface coverage of 1.2 nmol of SoxAX cm−2 geometric electrode area was calculated. These electrodes typically have a surface area 300 × the geometric area (25). Thus, SoxAX coverage corresponded to 4 pmol cm−2, which for a protein with dimensions of ~34 × 36 × 75 Å indicates good penetration of protein into the mesoporous structure.

Spectropotentiometric titrations monitored by MCD in the visible region were performed in an optically transparent electrochemical cell (1-mm path length) with a working electrode composed of a gold mini-grid as described previously (26). The samples typically contained 53 μM SoxAX in 50 mM Heps, 100 mM NaCl, pH 7, and 40 μM each of the following mediators: methylene blue, 2,6-dichlorophenolindophenol, phenazine methosulfate, ferricyanide, diaminodurene, anthraquinone-2,6-sulfate, anthraquinone-2-sulfate, benzyl viologen, and methyl viologen.

At near infrared (nIR) wavelengths, it was found that gold mini-grid electrodes were depolarizing and that the SnO2 electrodes were not transparent when they were conducting. As a consequence, potentiometric titrations of mediated SoxAX solutions monitored by MCD at nIR wavelengths were performed with a modification of the cell used for MCD at visible wavelengths. The mini-grid working electrode was replaced by a ribbon of gold foil folded to an “S”-shape 2 mm wide and 4 mm high. The gold ribbon was 1 mm wide so that when the “S” was inserted into the cell, the working faces of the electrode spanned the path length of the cell. The front window of the cell was masked off to ensure that only the sample within ~0.5 mm of the extremities of the working electrode lay in the light path. In this way, equilibration with applied potential, that is, reliant on diffusion of protein and redox mediators to the electrode, could be achieved within ~30 min. Spectra collected during MCD-monitored potentiometry in both the UV-visible and the nIR regions are presented without base-line correction.

Preparation of Chemically Poised Samples for MCD and EPR Spectroscopy—Samples of SoxAX (200 μM) in 50 mM Hepes, 100 mM NaCl, D2O, pH* 7 (where pH* is the apparent pH of the D2O solution determined using a standard glass pH electrode) were chemically poised at defined potentials in a sealed vessel housed in a nitrogen-filled chamber as described previously (27). Aliquots of chemical reductants/oxidants were titrated into the sample until equilibration at the desired potential was achieved and during which time the sample was stirred and maintained at 4 °C. Equilibrated samples were rapidly transferred to an anaerobic MCD cell for analysis by room temperature nIR-MCD or into EPR tubes and frozen. Samples of fully oxidized SoxAX were prepared by the addition of potassium ferricyanide, whereas samples poised at +45, −360, and −500 mV were prepared by reduction with ascorbate, Eu(II), and Eu(II) complexed by diethylene triamine pentaacetic acid (DTPA), respectively (28). To facilitate equilibration at a measurable potential, each experiment included 40 μM of each of the following mediators: trimethylhydroquinone, menadione, 5-hydroxynaphthoquinone, and methylene blue. For poised at −360 and −500 mV, the samples additionally contained 40 μM each of benzyl viologen, sulfonyl viologen, and methyl viologen.

Stock solutions of EuCl3 (20 mM) were prepared anaerobically and immediately prior to use by dissolution of the desired mass in 50 mM Hepes, 100 mM NaCl, D2O, pH* 7. A stock solution of DTPA was prepared in 0.2 M NaOH, D2O.

Preparation of Samples with Cyanide-ligated Heme—Samples for optical spectroscopy comprised either 40 μM (electronic absorbance) or 100 μM (nIR-MCD) SoxAX in 50 mM Tris/HCl, pH 7.8, or 50 mM Tris/HCl, pH* 7.8 D2O, respectively. Reduction of the samples was carried out in a nitrogen-filled chamber by the addition of sequential aliquots of either a 5 mM solution of sodium dithionite or a saturated (25 mM) solution of Eu(II)DTPA prepared as described above. Aliquots of KCN were titrated into the samples to produce the maximum change in the electronic absorbance spectra. Samples for EPR spectroscopy were prepared in a similar fashion with a protein concentration of 114 μM. The sample was transferred to an EPR tube and frozen prior to measurement. Between measurements, the sample was thawed, in a nitrogen-filled chamber if necessary, to allow the addition of dithione or KCN or air equilibration, the consequences of the chemical manipulations being monitored using electronic absorbance spectroscopy.

Spectroscopy—Electronic absorption spectra were recorded on either a Hitachi U4100 or a Hitachi U2900 spectrophotometer. MCD spectra were recorded using circular dichrographs, JASCO models J810 and J730 for the UV-visible and nIR regions, respectively. Either an Oxford Instruments SM1 split-coil superconducting solenoid with a 25-mm ambient temperature bore or an Oxford Instruments Special Spectromag 1000, a bespoke split-coil superconducting solenoid with a 50-mm ambient temperature bore, was used to generate a magnetic field of 6 T for the ambient temperature MCD measurements. EPR spectra were recorded on a spectrometer comprising an ER-200D electromagnet and microwave bridge interfaced to an EMX control system (Bruker, Spectrospin) and fitted with a
liquid helium flow cryostat (ESR-9, Oxford Instruments) and a dual mode X-band cavity (Bruker, type ER4116DM). EPR spectra were simulated using the Bruker WinEPR and SimFonia software and quantified by double integration of the simulated component spectra.

Global Analysis of Potentiometric Data—Ambient temperature methods allow the convenience of measuring the spectrum of a single sample across a wide range of electrochemical potential applied in situ. However, none of the spectroscopic techniques suitable for monitoring ambient temperature potentiometry were capable of resolving the different responses of the two thiolate-ligated hemes of SoxAX as was possible through the EPR of frozen samples. As a consequence, the midpoint potentials of the three SoxAX hemes were obtained from a global analysis of the potentiometric data collected from all of these approaches. Six data sets were included in this analysis. Five of these arose from optically monitored potentiometry: variation of signal intensity with potential from i) electronic absorbance at 552–565 nm (25 data points), ii) MCD at 548–553 (10 data points), iii) MCD at 1170 nm (20 data points), iv) MCD at 1800 nm (11 data points), and v) MCD at 1500 nm (11 data points). The sixth data set quantified the EPR intensity attributed to the LS1 and LS2 species as a function of potential (4 data points).

For each data set, the variation in intensity with potential was assumed to arise from the sum of the intensities arising from three hemes acting as independent sites undergoing single electron oxidation/reduction in a manner described by the Nernst equation

\[ y_{\text{calc}} = \frac{\sum A_i(1 + F/RT\exp(x_i - E_m(n)))}{y_{\text{obs}}} \]  
(Eq. 1)

where \( y_{\text{calc}} \) is the calculated normalized intensity of point \( i \), \( A_i \) is the fractional intensity attributed to heme \( n \), \( F \) is the Faraday constant, \( R \) is the ideal gas constant, \( T \) is the temperature, \( x_i \) is the potential of point \( i \), and \( E_m(n) \) is the midpoint potential of heme \( n \). The midpoint potential of each heme was constrained to be identical for each data set examined. The fractional change in intensity contributed by each heme to each data set was parameterized in the model but held constant during refinement when the value could be determined a priori by consideration of the system. The latter was the case for (a) the nIR-MCD intensity at 1500 and 1800 nm that must arise solely from the His/Met-ligated heme and (b) the EPR intensity that must arise solely from the His/Cys(S)-ligated heme, respectively. A broader feature of lower intensity was parameterized in the model but differing in axial ligation, as postulated for R. sulfidophilum.

The best fit to the experimental data was obtained using the parameters described in Table 3, yielding a \( \chi^2 \) of 7. The major features of all data sets are well described by this model, the main difference between calculated and observed behavior being due to residual intensity in the 1500 nm MCD response between 0 and +100 mV. Accounting for this feature in a model assuming Nernstian behavior would require the introduction of a fourth oxidation/reduction event not evident in any of the other data sets. Constraining the fractional intensities at the selected wavelengths at \( \sim 550 \) nm in the electronic absorbance and MCD data to be equal for both low potential hemes led to a 60% increase in \( \chi^2 \). Constraining the heme with the lowest \( E_m \) to give the lowest amplitude responses in these data sets led to a doubling of \( \chi^2 \). The significantly lower quality of the fits in these cases, indicated by the higher \( \chi^2 \) values, was caused by the model predicting that 550 nm absorbance increased at more positive potentials than observed.

RESULTS

Overview of the Properties of Recombinant SoxAX—A system for the heterologous expression of R. sulfidophilum SoxAX in R. capsulatus was developed to provide sufficient SoxAX protein for detailed spectroelectrochemical analysis. The recombinant R. sulfidophilum SoxAX protein crystallized in space group P2₁. The asymmetric unit had dimensions similar to those found in the 2.55 Å structure of the native protein (8) and contained four copies of SoxAX. The structure of the recombinant protein is essentially identical to that of the native protein (supplemental Fig. 1) with an overall average root mean square deviation of 0.26 Å (supplemental Table 1). Ligation of all three SoxAX hemes was the same as in the native structure. All four monomers of the asymmetric unit had CysS⁻ ligation of the active site Heme-2, although the recombinant SoxAX protein had been produced in the absence of thiosulfate or other Sox pathway components (supplemental Fig. 2). Equivalent turn-over-independent modification of the active site heme ligand has also been reported for another recombinant SoxAX protein (9).

The nIR-MCD spectrum of fully oxidized recombinant protein at ambient temperature (Fig. 1) was indistinguishable from that of the wild-type enzyme (7). Prominent features were present at 1170 and 1800 nm that have been assigned to the LMCT bands from the His/Cys(S)⁻ (29, 30) and His/Met (16) ligated ferric hemes, respectively. A broader feature of lower intensity spans the range 1300–1600 nm and arises from the vibrational sidebands of the peak at 1800 nm. The spectral features of the oxidized recombinant protein when viewed by X-band EPR (Fig. 2) and MCD in the UV-visible region (Fig. 3A) were also indistinguishable from those of wild-type enzyme (7). The recombinant enzyme, henceforth simply referred to as SoxAX, was used in all studies presented below.

Potentiometric Titrations—In contrast to the prominent nIR-MCD arising from low-spin ferric hemes (Fig. 1), the corresponding LMCT transitions of their ferrous counterparts are symmetry-forbidden (31). Consequently, potentiometric titration monitored by nIR-MCD affords the opportunity to resolve the redox properties of hemes closely matched in midpoint potential but differing in axial ligation, as postulated for R. sul-
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FIGURE 1. Potentiometric titration of SoxAX monitored by nIR-MCD spectroscopy. A, spectra of 160 \( \mu \)M SoxAX electrocally poised at increasingly negative potentials of +350 (solid black), +200 (broken black), −300 (solid gray), and −440 (broken gray) mV. B, normalized MCD signal intensity at 1800 (■), 1500 (○), and 1170 nm (△) as a function of equilibration potential; data were collected for oxidative and reductive titrations. Solid lines indicate the intensity predicted for Nernstian behavior of three independent n = 1 centers described by the parameters listed in Table 3. Open data points represent chemically poised samples. C, spectra of 200 \( \mu \)M SoxAX chemically poised at +400, +45, −360, and −510 mV using air, ascorbate, Eu(II), and Eu(II)DTPA, respectively. The greater noise in the spectra of SoxAX (8), because the energy of the LMCT for the ferric hemes will differ. To exploit this possibility, a thin layer electrochemical cell optically transparent to circularly polarized light of nIR wavelengths was developed as described under “Experimental Procedures.” The effect of electrochemical potential on the nIR-MCD of SoxAX is summarized in Fig. 1, A and B. When poised at potentials above +300 mV, SoxAX yields spectra that are typical of the fully oxidized enzyme. Lowering the sample potential to 0 mV removed all MCD intensity at wavelengths above 1400 nm and produced a small (15%) drop in the intensity of the feature at 1170 nm. Between 0 and −300 mV, the spectra were indistinguishable. Equilibrating at lower potentials produced a decrease in the intensity of the 1170 nm feature such that at −440 mV, it had 20% of that displayed by fully oxidized enzyme. Posing the sample at increasingly positive potentials reversed the spectral changes seen on reduction and demonstrated the reversibility of the events being monitored.

Attempts to equilibrate samples below −440 mV and generate fully reduced SoxAX, as evidenced by the absence of intensity in the nIR-MCD, were unsuccessful as current continued to flow from the samples even after the desired potential had been applied for several hours. Consequently, SoxAX was chemically poised at four potentials selected to confirm and extend the description of its redox chemistry (Fig. 1C). The spectrum of SoxAX poised at +400 mV was again characteristic of the fully oxidized enzyme. Posing the sample at +45 mV was sufficient to remove all intensity at wavelengths above 800 nm, and the sample poised at −360 mV showed a 40% reduction in the intensity at 1170 nm when compared with the fully oxidized sample.

FIGURE 2. Perpendicular mode X-band EPR spectra of SoxAX. A, spectra of SoxAX chemically poised at +400, +45, −360, and −490 mV with air, ascorbate Eu(II), and Eu(II)DTPA, respectively, and for the sample poised at −490 mV following reoxidation in air. Sample conditions are as in Fig. 1. Spectrometer parameters: microwave frequency 9.68 GHz, microwave power 2 milliwatts, modulation amplitude 10 G, and temperature 20 K. B, the spectrum of SoxAX poised at +400 mV together with its simulation by contributions from LS1a, LS1b, LS2a, and LS2b with the parameters and proportions indicated in Tables 1 and 2. The sum of the simulated contributions is indicated by the broken line in the upper trace.
\[ \Delta \varepsilon_{1170 \text{ nm}} \] values to Heme-1 and -2 by comparison with the nIR-MCD response of other hemoproteins having His/Cys \(^{-}\) ligation or that of the His/CysS \(^{-}\) -ligated Heme-2 of the \textit{S. novella} enzyme, respectively. The loss in 1170 nm intensity may reflect redox transformation of one of the SoxA hemes or simply a change in ligand orientation at one or both of these hemes in response to the reduction of Heme-3. Alternatively, the spectroscopic response of Heme-1 and -2 may be invariant at potentials greater than \(-300 \text{ mV}\) with the change in 1170 nm intensity at high potential reflecting the loss of the "tail" of the feature at 1800 nm.

For independent assessment of the redox properties of the His/Cys(S) \(^{-}\) -ligated Heme-1 and -2, they were probed further by the EPR of frozen SoxAX solutions chemically poised at potentials guided by the optical data collected at ambient temperatures (Fig. 2A). The X-band EPR of oxidized SoxAX is dominated by multiple rhombic trios between 260 and 380 mT characteristic of His/thiolate-ligated ferric hemes, whereas Heme-3 gives rise to a much smaller feature at \(g_0 = 3.48\). Each of these hemes is EPR-silent in the ferrous state. Poising of samples at \(+45 \text{ mV}\) removes the feature at \(g = 3.48\), consistent with reduction of the His/Met-ligated Heme-3 (Fig. 2A). The sample poised at \(-490 \text{ mV}\) was essentially EPR-silent, indicating reduction of all three SoxAX hemes.

Quantification of the effect of intermediate potentials on the EPR of SoxAX required spectral simulation. Although Heme-3 gives rise to only one detectable feature, the resonances from Heme-1 and -2 can only be satisfactorily simulated using four rhombic components (Fig. 2B). The latter are grouped into two sets (Table 1). As described previously (7), these are termed LS1 and LS2, each of which is composed of two components, a and b, reflecting minor heterogeneities. The total EPR intensity arising from ferric Heme-1 and -2 in the sample poised at \(+45 \text{ mV}\) is within error of that from fully oxidized SoxAX (Table 2).

Thus, neither of these hemes is redox-active between \(+300\) and \(0 \text{ mV}\). The redistribution of the LS1 and LS2 intensity between the four contributing species in these samples is indicative of changes in ligand orientation at Heme-1 and/or Heme-2 in response to reduction of Heme-3. This accounts, at least in part, for the change in MCD intensity at 1170 nm across this potential range. Significantly, the sample poised at \(-360 \text{ mV}\) displayed 50% of the LS2 intensity and 100% of the LS1 intensity displayed by fully oxidized SoxAX (Table 2). Therefore, the species giving rise to LS1 EPR signals must have a lower \(E_m\) value than those giving rise to the LS2 features.

Determining the two \(E_m\) values describing redox transformation of the thiolate-ligated hemes requires spectral resolution with high-fidelity across the potential domain. Consequently, two further ambient temperature spectroscopies allowing for precisely defined and \textit{in situ} potentiometric control were employed, namely MCD and electronic absorption in the visible region. MCD in the wavelength range 300–700 nm contains features from both ferric and ferrous hemes that additionally inform on their spin state (32). Spectra recorded for potentials ranging from \(+450\) to \(-500 \text{ mV}\) showed no indication of positive intensity at \(-430 \text{ nm}\) or negative intensity at \(-630 \text{ nm}\) that would indicate high-spin ferrous and ferric heme, respectively (Fig. 3A). Instead, the spectral changes were characteristic of the interconversion of low-spin ferric and low-spin ferrous heme. These effects were quantified using the potential dependence of the peak-to-trough intensity at \(-550 \text{ nm}\) arising

![FIGURE 3. Potentiometric titration of \textit{R. sulfidophilum} SoxAX monitored by electronic absorbance and MCD spectroscopy in the visible region. A, room temperature MCD spectra of 53 \(\mu\)M SoxAX poised at \(+450\) (black), \(-200\) (broken gray), and \(-500 \text{ mV}\) (solid gray) by equilibration with a gold mini-grid electrode. Inset shows the intensity at 548 nm minus 553 nm as a function of equilibration potential. B, electronic absorbance spectra of SoxAX adsorbed on a nanocrystalline mesoporous SnO\(_2\) electrode poised at \(+230\) (black), \(-270\) (broken gray), and \(-540 \text{ mV}\) (solid gray). The features arising from SoxAX are superimposed on the scattering background arising from the nanocrystalline mesoporous SnO\(_2\) electrode. Inset shows the intensity at 552 nm minus that at the 565 nm isosbestic point as a function of potential. Solid lines in both insets indicate the intensity predicted for Nernstian behavior of three independent \(n = 1\) centers described by the parameters listed in Table 3.](image-url)

### TABLE 1

<table>
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<th>Species</th>
<th>(g_x)</th>
<th>(g_y)</th>
<th>(g_z)</th>
<th>(W_x)</th>
<th>(W_y)</th>
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from low-spin ferrous heme (Fig. 3A, inset). More rapid equilibration of SoxAX across a wide potential window and thus greater spectral resolution across the electrochemical potential domain was achieved through its adsorption on an optically transparent SnO2 electrode. Equilibration with the applied electrode potential was almost instantaneous as monitored by electronic absorption in the visible region such that the reversibility of SoxAX redox chemistry could be assessed over multiple cycles (Fig. 3B). Interconversion of features typical of low-spin ferric \( \epsilon_{412 \text{ nm}} 350 \text{ mm}^{-1} \text{ cm}^{-1}, \epsilon_{553 \text{ nm}} 45 \text{ mm}^{-1} \text{ cm}^{-1} \) and low-spin ferrous enzyme \( \epsilon_{420 \text{ nm}} 550 \text{ mm}^{-1} \text{ cm}^{-1}, \epsilon_{555 \text{ nm}} 100 \text{ mm}^{-1} \text{ cm}^{-1} \) was observed and quantified by the variation in the difference between the absorbance at 552 nm and that of the isosbestic point at 565 nm with potential (Fig. 3B, inset).

The optically monitored titrations confirmed that the redox chemistry of SoxAX is confined to the interconversion of low-spin ferric/ferrous heme and the two windows of potential identified by nIR-MCD-monitored potentiometry. As a consequence, a global analysis was employed to produce a model describing all of the potentiometric data through the sum of three independent Nernstian responses for single electron centers as described under “Experimental Procedures.” All data sets, i.e. the plots of intensity versus potential, were modeled satisfactorily using the same set of \( E_m \) values: +210 mV for the His/Met-ligated Heme-3 and −340 and −400 mV for the thiolate-ligated hemes (Table 3 and solid traces in Fig. 1B and the insets of Fig. 3). This validates the results obtained using the novel nIR-MCD compatible electrochemical cell.

Although both thiolate hemes are found to contribute equally to the intensity change monitored by nIR-MCD, it is striking that the intensity change monitored by both electronic absorption and MCD spectroscopies in the visible region is striking that the intensity change monitored by both electronic absorption and MCD spectroscopies in the visible region is.

**TABLE 2**

Relative contributions of the rhombic features to EPR spectra of SoxAX

<table>
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<th>Total spin integration</th>
<th>% of one heme equivalent*</th>
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<th>LS1b</th>
<th>LS2a</th>
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<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

* Relative intensity of the thiolate envelope to that at +400 mV.

**TABLE 3**

Predicted midpoint potentials and optical properties of the hemes of SoxAX

<table>
<thead>
<tr>
<th>( E_m )</th>
<th>( \epsilon_{552} - \epsilon_{555} )</th>
<th>( \Delta \epsilon_{548} - \Delta \epsilon_{553} )</th>
<th>( \Delta \epsilon_{1170} )</th>
<th>( \Delta \epsilon_{1800} )</th>
<th>Ligand assignment</th>
</tr>
</thead>
<tbody>
<tr>
<td>+210 mV</td>
<td>31,000</td>
<td>680</td>
<td>0.14</td>
<td>1.0</td>
<td>His/Met</td>
</tr>
<tr>
<td>−340 mV</td>
<td>3,400</td>
<td>70</td>
<td>0.40</td>
<td>0.0</td>
<td>His/thiolate</td>
</tr>
<tr>
<td>−400 mV</td>
<td>22,000</td>
<td>150</td>
<td>0.46</td>
<td>0.0</td>
<td>His/thiolate</td>
</tr>
</tbody>
</table>

* Difference between the electronic absorbance at 552 nm and the 555 nm isosbestic point.

**FIG. 3**

B.

1. The isosbestic point at 565 nm with potential (Fig. 3B).
2. The absorbance at 552 nm and that of the isosbestic point at 565 nm.
3. The intensity change monitored by both electronic absorption and MCD spectroscopies in the visible region is.

**FIG. 4**

A.

The reactivity of SoxAX toward the potential exogenous ligand CN− was screened using electronic absorbance spectroscopy (Fig. 4A). Enzyme incubated with excess CN− in aerobic buffer had a spectrum similar to that of the enzyme as purified. The same was true for a sample reduced anaerobically with Eu(II)DTPA, equilibrated with excess CN−, and then re-equilibrated with air. EPR spectroscopy confirmed the small changes to the absorbance spectrum to arise from displacement of the methionine ligand of Heme-3 by CN− (not shown). By contrast, when the latter experiment was repeated using dithionite as the reductant, there was a 10% increase in the absorbance at 413 nm when compared with that of the other samples. The increase in electronic absorbance suggested that exposure to dithionite and cyanide produces a change of SoxAX heme ligation distinct to that produced by exposure to cyanide alone. This was confirmed and quantified by EPR and MCD spectroscopy (Fig. 4, B and C). Loss of the rhombic EPR features associated with LS1 species indicates displacement of thiolate as a ligand to Heme-2. As a consequence, the 50% reduction in 1170 nm MCD intensity shows Heme-2 to have \( \Delta \epsilon \) of \( -0.5 \text{ cm}^{-1} \text{ T}^{-1} \), which compares favorably with the value of 0.46 \( \text{ cm}^{-1} \text{ T}^{-1} \) predicted from global analysis of the potentiometric data (Table 3). The loss of MCD intensity at 1170 nm is accompanied by loss of all intensity at 1800 nm, the emergence of a new LMCT peak at 1650 nm, and increased EPR intensity at \( g \approx 3.4 \). Together these data indicate that after reaction of the enzyme with dithionite and CN−, Heme-2 in addition to Heme-3 has His/CN− ligation (16), whereas Heme-1 retains His/thiolate ligation.

Our attempts to use X-ray crystallography to resolve the structure of CN− -bound SoxAX were unsuccessful. We failed to find conditions where CN− -bound SoxAX would crystallize. We also failed to resolve any changes in the structure of SoxAX when crystals of the as prepared enzyme were soaked with dithionite and cyanide. Both results are consistent with SoxAX
undergoing conformational change on dissociation of the heme ligands, and further experiments are underway to identify the nature of these changes. As a consequence, further insight into the reactivity of SoxAX was obtained from spectroscopic studies. SoxAX reduced anerobically with dithionite, reoxidized by equilibration with air, and then exposed to cyanide displayed spectroscopic changes identical to those when cyanide was introduced prior to reoxidation of the enzyme (not shown). In contrast, very little change is observed in the EPR of SoxAX that has been reduced with dithionite and reoxidized by equilibration with air but not exposed to KCN.

**DISCUSSION**

We find no evidence to support the previous report that Heme-2 and -3 form a redox-linked pair in *R. sulfidophilum* SoxAX (8). Instead, the active site His/Cys$^-$-ligated Heme-2 has an $E_{m}$ that is some 600 mV lower than that of the His/Met- ligated Heme-3 (+210 mV). The $E_{m}$ values are consistent with those reported for the hemes located at the interface between SoxA and SoxX in the tri-heme *P. pantotrophus* and diheme *S. novella* enzymes. Similar behavior is also likely for the group 3 SoxAX of *Chlorobium tepidum* (5). As a consequence, a single catalytic mechanism is likely to operate in SoxAX enzymes. Kappler et al. (36) have proposed an essential role for Cu(II) in SoxAX catalysis. The work presented here corroborates previous studies in finding no indication of a site for binding Cu(II) within the structure of the enzyme resolved by x-ray crystallography (36). Thus, we propose that the activity of SoxAX originates from its heme cofactors and/or their ligands.

The mechanism of SoxAX must account for the oxidative coupling of thiosulfate to the Cys$^-$ of SoxY accompanied by the reduction of two molecules of cyt $c$. Bamford et al. (8) proposed a mechanism in which the thiolate ligand of Heme-2 dissociates to form a disulfide bond with thiosulfate in a process accompanied by the reduction of Heme-2 and -3 (Fig. 5A). In this mechanism, the oxidized thiosulfate is subsequently fused to the Cys$^-$ of SoxY in a sulfur transferase step and the SoxAX is returned to the all-ferric state by the reduction of 2 equivalents of cyt $c$. The reactivity of dithionite treated SoxAX toward cyanide reported here provides the first evidence that the dissociation of Cys$^-$ from Heme-2, as required for this mechanism, can indeed occur.

![FIGURE 4. Binding of CN$^-$ to redox-cycled SoxAX. A, electronic absorption spectra of ferric SoxAX exposed to a 250-fold excess of KCN (gray) and reduction with dithionite (black) or Eu(III)DTPA (gray crosses) followed by exposure to a 250-fold excess of KCN and equilibration with air. B, X-band EPR of 114 nm SoxAX in the ferric state as purified, following reduction with sodium dithionite, following reduction with sodium dithionite and exposure to a 250-fold excess of KCN, and following reduction with sodium dithionite, exposure to a 250-fold excess of KCN, and reoxidation in air. The broken trace shows the spectrum of enzyme that has been reduced with sodium dithionite and reoxidized in air but not exposed to KCN. Spectrometer parameters: microwave frequency 9.68 GHz, microwave power 2 milliwatts, modulation amplitude 10 G, and temperature 20 K. C, nIR-MCD of SoxAX as purified (broken black) and following reduction with sodium dithionite, exposure to a 250-fold excess of KCN, and reoxidation in air (solid black). The lower intensity of the feature at 1800 nm relative to that at 1170 nm when compared with Fig. 1 is consistent with the partial reduction of Heme-3 in the enzyme as purified.](image)

![FIGURE 5. Proposed mechanisms of SoxAX activity. Reaction schemes for the SoxAX catalyzed attachment of thiosulfate to SoxY are shown. Scheme A, Cys-222A dissociates from Heme-2 and forms a disulfide bond with thiosulfate, liberating two electrons that reduce Heme-2 and -3 to the ferrous state. Subsequently, thiosulfate is transferred to SoxY, and SoxAX is returned to the resting state by the reduction of 2 equivalents of cyt $c$. Scheme B, thiosulfate binds directly to SoxY, liberating two electrons that reduce Heme-2 and -3 of SoxAX. Subsequent steps are as above. Cys-222A remains ligated to Heme-2 throughout.](image)
Heme Reactivity in SoxAX

Eu(II) reduces the SoxAX hemes as effectively as dithionite, but it does not act as a precursor for the displacement of the CysS⁻ ligand to Heme-2 by cyanide. The dissociation of CysS⁻ may then arise from the ability of dithionite and/or its oxidation products to mimic chemistry that can be induced by thiosulfate, the substrate of SoxAX. The details of this chemistry are unclear at present. The EPR of SoxAX following reduction with dithionite and equilibration with air demonstrates that Heme-2 retains a thiolate (RS⁻, where R is any alkyl ligand) ligand in the absence of CN⁻. This is consistent with the reported structure of dithionite-reduced SoxAX where the CysS⁻ ligand remains in place (8). However, it should be noted that the spectroscopic response of hemes with His/RS⁻ ligation is insensitive to the nature of R. It may be that Heme-2 is ligated by HS⁻ or Cys⁻ rather than CysS⁻ in solutions of dithionite exposed SoxAX.

An alternative to the mechanism of Bamford et al. (8) can be proposed for SoxAX activity (Fig. 5B). In this mechanism, SoxAX mediates the direct coupling of thiosulfate to SoxY without dissociation of any heme ligands. To our knowledge, SoxAX mediates the direct coupling of thiosulfate to SoxY (9). In this mechanism, proposed for SoxAX activity (Fig. 5B), Cys-Cys disulfide bonds and for which an oxidation state of 2 would be consistent with the described reaction, necessary to activate the thiosulfate oxidizing-monoxygenase enzyme. This mechanism must include ferrous Heme-2 without CysS⁻ or Cys⁻ formation, as demonstrated here, and is key to thiosulfate oxidation.

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REFERENCES


