SYNTHETIC ANALOGS OF THE ACTIVE SITES OF OXIDIZED AND REDUCED RUBREDOXIN

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Abstract. — The bischelate monoanion \([Fe((SCH_2)\_2C\_6H_4)\_2]^\-\) contains an unconstrained, near-tetrahedral Fe(III)-S\(_4\) coordination unit (similar to the Fe-S\(_4\) coordination unit occurring in rubredoxin proteins) and exhibits the Fe(III)/Fe(II) redox couple. Mössbauer parameters at 77 K are isomer shift \(\delta = 0.13\) mm/s (relative to iron metal) and electric quadrupole splitting \(\Delta E_Q = 0.57\) mm/s. Application of an external magnetic field parallel to the direction of observation at 4.2 K induces a saturation magnetic hyperfine field of \(-380\) kOe. Upon reduction to the dianion \([Fe((SCH_2)\_2C\_6H_4)\_2]^2\-\) the isomer shift changes to \(\delta = 0.61\) mm/s and the quadrupole splitting at 4.2 K to \(\Delta E_Q = 3.34\) mm/s. From the temperature dependence of \(\Delta E_Q\) for 1.4 K < \(T\) < 295 K we infer the splitting of the ground orbital doublet, \(\epsilon(d_{z^2}, d_{x^2-y^2})\), to be \(d \approx 900\) cm\(^{-1}\). External magnetic fields up to 80 kOe reveal the sign of \(\Delta E_Q\) to be negative indicating that the ground orbital has \(d_{z^2}\) symmetry and induce a magnetic hyperfine interaction of negative sign. For \(H_0 = 60\) kOe, \(H_m = -128\) kOe. \(|H_{hf}|\) increases with increasing \(H_0\), showing that saturation has not been reached up to \(H_0 = 80\) kOe, and implying a zero field splitting of several cm\(^{-1}\) of the electronic ground state. These results as well as magnetic moment measurements for the unconstrained model compounds are comparable to measurements in oxidized and reduced rubredoxin and imply that an «entatic state» description of rubredoxin is not valid.
More significantly, X-ray diffraction studies of oxidized Rd from *Clostridium pasteurianum* by Jensen and coworkers [4] at the present 1.5 Å stage of resolution demonstrate considerable deviations of the Fe-S₄ unit from a perfect tetrahedron. One of the Fe-S bond lengths (2.05 Å ± 0.03) is abnormally short and indeed falls outside the range of iron-sulfur bond lengths observed in any synthetic Fe(II, III)-SR complex [5]. The other three Fe-S bond lengths are nearly equal and fall within the 2.24-2.35 Å range. This stereochemistry has been taken as evidence that the Fe-S₄ active site in Rdₐx exists in a constrained state in accordance with the *entatic state* hypothesis of Vallee and Williams [6]. This hypothesis postulates that the conformation of the polypeptide chain imposes structural constraints on the active site of metalloenzymes resulting in a state of *entasis* or tension of unique physical and chemical properties with implications for the catalytic activity, redox potentials, and electron transfer kinetics of the proteins.

Recent extended X-ray absorption fine structure (EXAFS) studies of Rd using the high intensity X-ray flux from synchrotron radiation show, however, that the four Fe-S bond lengths are probably closer to being equal than the X-ray diffraction crystal structure of Rdₐx indicates. Studies on Rdₐx from *Peptococcus aerogenes* by Shulman et al. [7] give an average Fe-S bond distance of 2.24 Å with a deviation of 0.1 Å. EXAFS measurements by Sayers et al. [8] on *C. pasteurianum* Rd indicate an average Fe-S bond length of 2.30 Å ± 0.04 Å and a rms deviation about this average of 0.06 Å ± 0.004 Å.

Elucidation of the conflicting results on the structure of the Fe-S₄ cluster in Rd and a critical examination of the validity of the description of the Fe-S₄ complex in Rd as an *entatic state* may be approached by comparative studies of the protein and inorganic synthetic iron-sulfur compounds for which the local site symmetry is essentially unconstrained, and whose physical and electronic properties sufficiently closely approach those of the Fe-S₄ cluster in Rd that they can be considered suitable synthetic analogues [5, 9, 10] of the protein itself. To this end we have carried out Mössbauer measurements on synthetic analogues of Rdₐx and Rdₐz and compare the results with those of Rdₐx and Rdₐz available in the literature.

The bischelate monoanion \([\text{Fe}((\text{SCH}_2)_2C_6H_4)_2]^\text{1-}\) (bis(o-xylyl-α,α'-dithiolato) ferrate (III) monoanion) contains an unconstrained, near-tetrahedral Fe(III)-S₄ coordination unit and exhibits the Fe(III)/Fe(II) redox couple. Figure 1a shows the X-ray diffraction structure obtained by Ibers [9] and figure 1b gives the details of the structure around the iron atom. The X-ray structure [10] of the reduced compound \([\text{Fe}((\text{SCH}_2)_2C_6H_4)_2]^\text{2-}\) (bis(o-xylyl-α,α'-dithiolato) ferrate (II) dianion) is similar to that of the oxidized compound with an average increase of 0.1 Å in Fe-S bond distance upon reduction. These are the only synthetic analogues of Rd which have been isolated in both oxidation states with a common ligand structure. Thiolate sulfur coordination simulates cysteinate binding to iron in the native proteins.

2. **Experimental.** — Details on the synthesis, chemical and physical characterization and X-ray structure determination of the compounds are given in detail elsewhere [9, 10]. Mössbauer spectra were taken with a conventional constant acceleration spectrometer operated in the normalized mode. The spectrometer was equipped with a Nb-Sn superconducting magnet and cryogenic equipment for measurements in longitudinal magnetic fields up to 85 kOe and temperatures as low as 1.5 K. The source, Co⁵⁷ in Rh, and absorber were maintained at the same temperature. All Mössbauer spectra were taken on crystalline or frozen solution samples with an average thickness of 3-5 mg Fe/cm², resulting on the average in ~ 10% effects. Mössbauer parameters were obtained by least square fitting of the experimental data to Lorentzians, and all errors quoted are ± 1 standard derivation obtained from the computer fits.

3. **Results and discussion.** — \([\text{Fe}((\text{SCH}_2)_2C_6H_4)_2]^\text{1-}\): Mössbauer spectra of \([\text{Fe}((\text{SCH}_2)_2C_6H_4)_2]^\text{1-}\), the analog of Rdₐx obtained as the tetraethylammonium salt were taken at room temperature, 77 and 4.2 K. Samples in the crystalline phase and in dimethylformamide and dimethylformamide/dichloromethane (3:1 v/v) frozen solutions were studied. All spectra consist of a single quadrupole doublet with temperature-independent quadrupole splitting. Mössbauer parameters are summarized in table I together with results for Rdₐz for comparison. The results are consistent with a high spin Fe(III)-S₄ coordination unit distorted from T₄ symmetry. Mössbauer spectra at 4.2 K in the absence and presence of an applied
The line shape at 77 K is asymmetric with the low velocity component broader and less intense than the high velocity component. At 4.2 K the lines are more nearly symmetric.

Application of longitudinal external magnetic fields up to 80 kOe show that the sign of the principal component of the electric field gradient tensor, \( V_{zz} \), is positive and that a magnetic hyperfine field is induced parallel to the applied field but in the opposite direction. The variation of the induced field with the magnitude of the applied field indicates that at 80 kOe applied field saturation of the induced field has been reached with a saturation value \( H_{sat} = -380 \) kOe. This is in good agreement with \( H_{sat} = -370 \) kOe obtained for \( \text{Rd}_{\text{ox}} \) \[11\] and implies similar bond covalency effects in the analogue and \( \text{Rd}_{\text{ox}} \). These internal fields are reduced from the usual saturation value of about \(-500 \) kOe \[12\] found for Fe(III) with oxygen or nitrogen coordination, and presumably reflect the effects of spin depolarization and greater metal-ligand bond covalency. The absence of magnetic hyperfine structure at low temperatures in zero magnetic field and its presence in the \( \text{Rd}_{\text{ox}} \) spectra reflect different electronic relaxation times for the two cases.

The near identity of Mössbauer spectra for [Fe((SCH\(_2\))\(_2\)C\(_6\)H\(_4\))\(_2\)]\(^{3+}\) observed in the crystalline state with those found in frozen solutions, including the magnetically perturbed spectrum in dimethyl-formamide-dichloromethane, implies that the solid and solution structures are the same. This indicates that no crystallographic constraints are imposed on the Fe-S\(_2\) structure. EPR spectra of [Fe((SCH\(_2\))\(_2\)C\(_6\)H\(_4\))\(_2\)]\(^{3+}\)
show a sharp $g = 4.3$ signal [9] indicative of high-spin rhombic Fe(III) but overall the spectra are more complicated than those of $R_{d_{	ext{ox}}}$ and have not as yet been interpreted. This might be a result of exchange interactions between Fe-S$_4$ clusters which are absent in the protein due to their greater separation.

Magnetization measurements of

\[ [\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^{1-} \]

at 4.2 K are shown in figure 3. The solid lines are theoretical assuming a Hamiltonian of the form

\[ H = D[S^2 - \frac{1}{3} S(S+1)] + E(S^2 - S_z^2) + 2\beta H_0 \cdot S \tag{1} \]

with $E/D \approx \frac{1}{8}$ as indicated by the EPR results. At high field, the experimental values are reproduced parallel exchange coupling between iron-centers. This observation is consistent with the complexities of the EPR spectrum mentioned above and also the rapid electronic relaxation observed in the Mössbauer spectrum.

\[ [\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^{1-} : \text{Mössbauer spectra for crystalline samples of } [\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^{2-} \text{obtained as sodium (tetraphenylarsonium) salt were taken at 295, 77, 4.2 K and temperatures between 4.2 and 1.4 K. All spectra consist of a single symmetric quadrupole doublet with a quadrupole splitting ($\Delta E_Q$) of the order of 3 mm/s and isomer shift ($\delta$) of $\sim 0.6$ mm/s indicative of high spin Fe(II). Spectral parameters are listed in table I and spectra at $H_0 = 0$ and various temperatures are shown in figure 4.

\[ \text{FIG. 3.} \quad \text{Mössbauer spectra of } \text{Na(Ph}_4\text{As)} [\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2] \at various temperatures. The solid lines give least-squares fits of the experimental points to Lorentzian lines. The small broad feature centered about 0 mm/s is presumably due to an Fe(III) impurity. \]

In a ligand field of $T_d$ symmetry the free ion $^5D$ term of Fe(II) is split into a lower $^5E(c^3 t^2_2)$ and upper $^3T_2(c^2 t^2_0)$ state separated by $\sim 5000 - 6000$ cm$^{-1}$ in energy [10]. The large quadrupole splitting observed indicates that the ground state orbital degeneracy is lifted via a rhombic distortion of the tetrahedral point symmetry. From the small temperature dependence of $\Delta E_Q$ observed over the temperature range 1.4 - 295 K (Table II), an estimate of the energy separation between the ground orbital to the next excited orbital state can be obtained by fitting the experimental data to an equation of the form $\Delta E_Q(T) = \Delta E_Q(0) \tanh (A/2kT)$ expected for the thermal population of the two isolated orbital levels, $d_{z^2}$ and $d_{x^2-y^2}$, split by an energy $A$. $\Delta E_Q(0)$ is the quadrupole splitting at zero temperature when only the ground orbital state is populated and $\Delta E_Q(T)$ is
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its value at temperature \( T \). The above equation is correct within the approximation that any splittings due to the spin-orbit interaction are much smaller than \( A \) and the direct lattice contribution to the EFG is negligible. By using \( \Delta E_{Q}(0) = 3.32 \text{ mm/s} \), the value observed at 1.4 K, an estimate of \( A \approx 900 \text{ cm}^{-1} \) is obtained. The temperature dependence observed by Rao et al. [11] for \( \text{Rd}_{\text{red}} \) is somewhat smaller than that of \( [\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^{2-} \). The latter, however, is very similar to that of lyophilized reduced Rd observed by Phillips et al. [14] for which Eaton and Lovenberg [1] estimate \( A \approx 850 \text{ cm}^{-1} \). Due to the small temperature variation observed and the experimental error associated with the values of \( \Delta E_{Q} \), the above estimates of \( A \) are probably correct only within \( \pm 100 \text{ cm}^{-1} \).

To characterize further the ground state of the \( \text{Fe}^{2+} \) ion, Mössbauer spectra in longitudinal magnetic fields were obtained. As discussed by Collins [15] and Gabriel and Ruby [16] when the line width is taken into account, the superposition of the absorption lines arising from transitions between the magnetically perturbed nuclear levels in the presence of the randomly oriented EFG tensor in a powder absorber results in the splitting of the \( | \frac{3}{2}, \pm \frac{3}{2} \rangle \rightarrow | \frac{3}{2}, \pm \frac{3}{2} \rangle \) transition line into an apparent doublet and the \( | \frac{1}{2}, \pm \frac{1}{2} \rangle \rightarrow | \frac{1}{2}, \pm \frac{1}{2} \rangle \) line into an apparent triplet. The spectrum of \( [\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^{2-} \) in the presence of a magnetic field of 80 kOe at 4.2 K is shown in figure 5.

![Mössbauer spectrum of Na(Ph$_4$As) $[\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^{2-}$ at 4.2 K in an external field of 80 kOe applied parallel to the γ-ray direction. The solid line is theoretical for a powder sample with $\Delta E_{Q} = 3.32 \text{ mm/s}$ and an effective magnetic field of 63 kOe with longitudinal configuration. The differences between the experimental data and calculated curve are due to anisotropy of the hyperfine interaction in the sample.](image)

The low energy absorption line of the quadrupole doublet observed at \( H_0 = 0 \) has been split into a doublet and the high energy absorption line into a triplet, indicating that the sign of the principal component of the EFG, \( V_{zz} \), is negative. Similar spectra were obtained at lower values of the applied field as well as at higher absorber temperatures. The negative sign of \( V_{zz} \) demonstrates that the ground state orbital of the \( \text{Fe}^{2+} \) ion has $d_{3z^2}$ symmetry, as has previously been shown for \( \text{Rd}_{\text{red}} \) [1]. The solid line in figure 5 is theoretical, assuming the field at the nucleus \( H_n = 63 \text{ kOe} \). The shape of the observed spectrum reflects anisotropy of the hyperfine interaction.

Figure 6 gives the energy level diagram within the $d^6$ electronic configuration for \( [\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^{2-} \).

![d-orbital energy level diagram for \( \text{Fe}^{2+} \) in \( [\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^{2-} \) in distorted \( T_d \) symmetry. Orbital energies for \( \text{Rd}_{\text{red}} \) are taken from Ref. [1].](image)

The ground state symmetry and rhombic splitting \( A \) were determined by Mössbauer spectroscopy as described above, while the splittings of the upper state \( t_2 \) were inferred from electronic spectral measurements [10]. Here orbital separations are approximated with spectroscopic energy levels since all quintet states derived from the same term $5D$ have the same mutual electron repulsion parameters. In the same diagram the corresponding energy levels for \( \text{Rd}_{\text{red}} \) [1] are also included for comparison. Although, the larger energy splittings of \( t_2 \) for the case of \( \text{Rd}_{\text{red}} \) indicates a probable greater distortion of the Fe-S$_2$ symmetry in Rd the overall cubic splitting \( A_x \) is of the same order for the two cases.

Comparisons of the observed Mössbauer spectra in the presence of external magnetic fields with computer generated spectra (Fig. 5) for an iron nucleus in the presence of a magnetic field in a randomly oriented EFG reveal that the observed field at the nucleus \( H_n \) differs from the applied magnetic field \( H_0 \), indicating the presence of a magnetic hyperfine interaction induced by the applied field. For \( H_0 = 60 \text{ kOe} \), \( | H_n | = 68.5 \text{ kOe} \); for \( H_0 = 80 \text{ kOe} \), \( | H_n | = 63 \text{ kOe} \). Since \( H_n \) decreases as \( H_0 \) increases from 60 to 80 kOe, we infer that the sign of \( H_n \) is negative. The 60 and 80 kOe spectral parameters yield \( H_n = -128 \) and \( -143 \text{ kOe} \), respectively. The parameters obtained at 1.4 K yield the same value of \( H_n \) as those obtained at 4.2 K for \( H_0 = 60 \) and 80 kOe. The fact that \( | H_n | \) continues to increase as \( H_0 \) increases from 60 to 80 kOe implies a zero field splitting of the
electronic ground state of the order of several degrees. The full saturation value of the magnetic hyperfine interaction constant would not be observed until \( \mu H_0 \) is large compared to the zero field splitting.

The magnetic hyperfine interactions induced by the applied field in the present case compare with those observed by Rao et al. [11] in Rd_{\text{red}} as follows. For \( H_0 = 60 \) kOe, they obtained \( H_m = -193 \) kOe at temperatures between 4.2 and 1.3 K. However, as discussed above, if there is a zero field splitting, this does not represent the full saturation value. If we assume comparable zero field splittings in the ground state electronic structures, and Fe-S bond covalency properties, this similarity doubtless extends to other Rd proteins such as, e.g., that from Ch. ethylica [11]. These results are also in agreement with the Mössbauer parameters of several tetrahedral Fe(II)-S_4 complexes recently measured by Kostikas et al. [18] such as \([\text{Fe(SPh)}_4]^2-\), which is also included in table I. Their values at 77 K are entirely similar to those of \([\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^2-\), whose isomer shift and quadrupole splitting in turn closely match those of \( C. \text{pasteurianum} \) Rd_{\text{red}} [11].

Although the metal ion site symmetry in Rd proteins may be slightly different than is the case in the present analogues, the close degree of agreement observed suggests that the actual structural disparity between the unconstrained structural analogue 

\[
[\text{Fe}((\text{SCH}_2)_2\text{C}_6\text{H}_4)_2]^2-
\]

and Rd_{\text{ox}} may not be as large as that suggested by a comparison of their X-ray diffraction crystal structures. Thus, the present results do not lend support to an entatic state description of the Fe-S_4 site in Rd_{\text{ox}} or Rd_{\text{red}}. As mentioned in the introduction, this is in accord with the results of EXAFS studies.

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**References**


