Two-Dimensional Magnetization Exchange Spectroscopy of *Anabaena* 7120 Ferredoxin. Nuclear Overhauser Effect and Electron Self-Exchange Cross Peaks from Amino Acid Residues Surrounding the 2Fe–2S* Cluster†,‡

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**ABSTRACT:** Hyperfine 1H NMR signals of the 2Fe–2S* vegetative ferredoxin from *Anabaena* 7120 have been studied by two-dimensional (2D) magnetization exchange spectroscopy. The rapid longitudinal relaxation rates of these signals required the use of very short nuclear Overhauser effect (NOE) mixing times (0.5–20 ms). The resulting pattern of NOE cross-relaxation peaks when combined with previous 1D NOE results [Dugad, L. B., La Mar, G. N., Banci, L., & Bertini, I. (1990) *Biochemistry* 29, 2263–2271] led to elucidation of the carbon-bound proton spin systems from each of the four cysteines ligated to the 2Fe–2S* cluster in the reduced ferredoxin. Additional NOE cross peaks were observed that provide information about other amino acid residues that interact with the iron-sulfur cluster. NOE cross peaks were assigned tentatively to Leu, Arg, and Ala on the basis of the X-ray coordinates of oxidized *Anabaena* 7120 ferredoxin [Rypniewski, W. R., Breiter, D. R., Benning, M. M., Wesenberg, G., Oh, B.-H., Markley, J. L., Rayment, I., & Holden, H. M. (1991) *Biochemistry* 30, 4126–4131]. Three chemical exchange cross peaks were detected in magnetization exchange spectra of half-reduced ferredoxin and assigned to the 1H* protons of Cys and Cys [both of whose sulfur atoms are ligated to Fe(III)] and Arg (whose amide nitrogen is hydrogen-bonded to one of the inorganic sulfurs of the 2Fe–2S* cluster). The chemical exchange cross peaks provide a means of extending assignments in the spectrum of reduced ferredoxin to assignments in the spectrum of the oxidized protein. Our results suggest that 2D magnetization exchange spectroscopy employing short mixing times will be useful for the assignment and characterization of hyperfine 1H peaks in a variety of paramagnetic proteins.

In macromolecules, magnetization exchange by chemical exchange and cross relaxation (the nuclear Overhauser effect or NOE) are formally equivalent (Noggle & Schirmer, 1971; Neuhaus & Williamson, 1989). The first magnetization exchange investigation of a paramagnetic protein was the classic pulsed NMR study of chemical self-exchange in a mixture of oxidized and reduced cytochrome c (Redfield & Gupta, 1971). Cross-relaxation (NOE) studies of paramagnetically shifted (hyperfine) resonances appeared later (Gordon & Wüthrich, 1978) and became a valuable tool for spectral assignments, particularly in heme proteins (Trewhella et al., 1979; Keller & Wüthrich, 1980; Lecomte et al., 1985).

Hyperfine proton signals of iron–sulfur proteins were first reported by Poe et al. (1970). The hyperfine signals of ferredoxins generally are characterized by their relative intensities (number of protons) and by the magnitudes and temperature dependencies of their chemical shifts [reviewed by Markley et al. (1986)]. Two new strategies have been used recently to assign ferredoxin hyperfine proton resonances: incorporation of deuterium-labeled cysteine into the protein (Cheng et al., 1990) and detection of one-dimensional (1D) nuclear Overhauser enhancements (NOE's) (Dugad et al., 1990). The latter approach led to preliminary sequence-specific assignments of several of the cysteine proton resonances in the ferredoxins from *Spirulina platensis* and *Prophyra umbilicalis* (Dugad et al., 1990).

Weak NOE's present in regions of overlapping peaks are difficult to detect in 1D difference spectra; they should be more easily resolved in 2D NOE spectra. We show here that hyperfine 2D magnetization exchange spectroscopy can be carried out with a ferredoxin provided that short mixing times and rapid recycle times are used in the data collection. Analysis of the patterns of NOE cross peaks from reduced *Anabaena* 7120 ferredoxin enabled us to characterize the proton spin systems of all four cysteine ligands to the 2Fe–2S* cluster plus those of three other residues located near the cluster. In addition, chemical exchange cross peaks found in 2D magnetization exchange spectra of half-reduced ferredoxin

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‡NMR data will be deposited in BioMagResBank (Ulrich et al., 1989) under ref 1D 889.

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EXPERIMENTAL PROCEDURES

Ferredoxin was isolated from the vegetative form of *Anabaena* 7120 as described previously (Oh & Markley, 1990a). Samples were reduced (totally or half) as described by Skjeldal et al. (1990). Protein samples used for NMR experiments were dissolved in 0.5 mL of a 50 mM phosphate buffer in \( \text{D}_2\text{O} \). The pH* was 8.2. The final protein concentration was 4 mM. The oxidation state of the protein sample was checked by analysis of 1D \(^1\text{H} \) NMR spectra obtained before and after each NOESY data set was collected (Skjeldal et al., 1990).

NMR data were collected on a Bruker AM 600-MHz spectrometer. \(^1\text{H} \) NOESY experiments (Macura & Ernst, 1980; Anil Kumar et al., 1980) were collected in the pure absorption mode. The TPPI method (Marion & Wüthrich, 1983) was used for quadrature detection in the second dimension. The receiver and transmitter were aligned to decrease baseline distortions (Marion & Bax, 1988). The NOESY mixing times were between 0.5 and 20 ms, as specified along with other details for individual experiments in the figure captions. \(^1\text{H} \) chemical shifts are referenced to internal 3-(trimethylsilyl)propionate (TSP), which was assigned as 0 ppm.

RESULTS AND DISCUSSION

One-dimensional \(^1\text{H} \) NMR data from reduced *Anabaena* 7120 vegetative ferredoxin are shown in Figure 1. Peaks a–e, j, and k show weak Curie-type temperature dependence (the peaks shift toward their diamagnetic positions as the temperature is raised) whereas peaks f–i show anti-Curie temperature dependence (the peaks shift away from their diamagnetic positions as the temperature is raised) (Chan & Markley, 1983; Skjeldal et al., 1990). Signals with Curie scalar shifts are assigned to protons on residues that are ligated to Fe(III), and signals with anti-Curie scalar shifts are assigned to protons on residues that are ligated to Fe(II) (Dunham et al., 1971; Bertini et al., 1984).
Limited radio frequency pulse power ($\gamma B_1/2\pi \approx 25$ kHz) prevented us from covering the full spectral window in the 2D magnetization exchange experiments. We decided not to include the four peaks at lowest field (a, b, c, and d, Figure 1B). Dugad et al. (1990) did not detect any 1D NOEs among these four peaks, but they found an NOE between peaks a and j and another NOE between peaks c and e. This evidence was used to assign peaks c and e to the $^1$H protons of these two cysteines (Dugad et al., 1990). Note that our exchange spectra were obtained at 27 °C, which is below the temperature at which peaks i and j cross in spectra of that ferredoxin. We reexamined this assignment by adding hydrogen to the 1H 2O solution, pH 8.2, and 27 °C.

Table I: Hyperfine Related Two-Dimensional NOESY Cross Peaks Observed in the Spectrum of Reduced Anabaena 7120 Vegetative Ferredoxin in $^1$H 2O Solution, pH 8.2, and 27 °C

<table>
<thead>
<tr>
<th>cross peak designation</th>
<th>chemical shifts (ppm)</th>
<th>diagonal peaks</th>
<th>assignments</th>
<th>distance (Å)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1a</td>
<td>-0.56, 24.72</td>
<td>-f</td>
<td>Ala$^4$H$_3$, Cys$^1$H$_3$</td>
<td>3.73 (3.56)</td>
</tr>
<tr>
<td>1b</td>
<td>3.20, 24.72</td>
<td>-/g</td>
<td>Cys$^6$H$_4$, Cys$^6$H$_4$</td>
<td>2.55 (2.32)</td>
</tr>
<tr>
<td>1c</td>
<td>16.38, 24.72</td>
<td>i, g</td>
<td>Cys$^6$H$_3$, Cys$^6$H$_3$</td>
<td>2.98</td>
</tr>
<tr>
<td>2a</td>
<td>5.00, 19.97</td>
<td>-h</td>
<td>Cys$^6$H$_4$, Cys$^6$H$_4$</td>
<td>2.29</td>
</tr>
<tr>
<td>2b</td>
<td>16.33, 19.97</td>
<td>i, h</td>
<td>Cys$^6$H$_2$, Cys$^6$H$_2$</td>
<td>1.76</td>
</tr>
<tr>
<td>3a</td>
<td>0.29, 18.04</td>
<td>-j</td>
<td>Leu$^{27}$H$<em>{311}$, Cys$^7$H$</em>{311}$</td>
<td>5.09 (5.04)</td>
</tr>
<tr>
<td>3b</td>
<td>1.08, 18.04</td>
<td>-j</td>
<td>Leu$^{27}$H$<em>{311}$, Cys$^7$H$</em>{311}$</td>
<td>3.16 (3.07)</td>
</tr>
<tr>
<td>3c</td>
<td>6.84, 18.04</td>
<td>-j</td>
<td>Leu$^{27}$H$<em>{311}$, Cys$^7$H$</em>{311}$</td>
<td>2.53</td>
</tr>
<tr>
<td>3d</td>
<td>8.39, 18.04</td>
<td>-j</td>
<td>Leu$^{27}$H$<em>{311}$, Cys$^7$H$</em>{311}$</td>
<td>2.09</td>
</tr>
<tr>
<td>4a</td>
<td>5.05, 16.89</td>
<td>i, h</td>
<td>Cys$^4$H$_3$, Cys$^4$H$_3$</td>
<td>2.56</td>
</tr>
<tr>
<td>4b</td>
<td>19.45, 16.89</td>
<td>h, i</td>
<td>Cys$^4$H$_3$, Cys$^4$H$_3$</td>
<td>1.76</td>
</tr>
<tr>
<td>4c</td>
<td>24.16, 16.89</td>
<td>g, i</td>
<td>Cys$^4$H$_3$, Cys$^4$H$_3$</td>
<td>2.98</td>
</tr>
<tr>
<td>5a</td>
<td>0.055, 12.57</td>
<td>-k</td>
<td>Arg$^{42}$H$_2$, Arg$^{42}$H$_2$</td>
<td>2.96</td>
</tr>
<tr>
<td>5b</td>
<td>1.77, 12.57</td>
<td>-k</td>
<td>Arg$^{42}$H$_2$, Arg$^{42}$H$_2$</td>
<td>2.64</td>
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<tr>
<td>5c</td>
<td>4.20, 12.57</td>
<td>-k</td>
<td>Arg$^{42}$H$_2$, Arg$^{42}$H$_2$</td>
<td>2.28</td>
</tr>
<tr>
<td>5d</td>
<td>5.47, 12.57</td>
<td>-k</td>
<td>Arg$^{42}$H$_2$, Arg$^{42}$H$_2$</td>
<td>2.75</td>
</tr>
</tbody>
</table>

*See Figure 2. Peaks f and g overlap at 27 °C. 2D NOE connectivities were observed between these two peaks at 11 °C where they do not overlap (spectrum not shown); they are not listed here. NOE connectivities are assigned as i, j, where the dipolar transfer is from i to j. Hydrogens were added onto the 1.5-Å refinement (manuscript in preparation) of the X-ray structure of oxidized Anabaena 7120 vegetative ferredoxin (Rypniewski et al., 1991) by means of the MOLEDT routine of the INSIGHT II graphics molecular modeling package (Biosym Technologies, Inc.). 1H−H distances were measured interactively by using the INSIGHT II software. Average distances were calculated for protons on methyl groups assuming free methyl group rotation. Two distances are given: that outside parentheses is for a third-power average $\langle(r^3)^{1/3}\rangle$, and that within parentheses is for a sixth-power average $\langle(r^6)^{1/6}\rangle$. A dash (−) indicates that the diagonal peak was not observed.

The stereochemical designation used here is the RS system of Cahn et al. (1966), which has been endorsed by the IUPAC–IUB Joint Commission on Biochemical Nomenclature (1984).
weaker NOE cross peak (3a) to the longer distance. No NOE was observed from the 'H of Leu52, which is 4.66 Å distant from the Cys79 H*.

All the assigned NOE's correspond to distances less than 3.0 Å with the exception of peak 3a at 4.58 Å assigned to interaction between the Cys101 H* and the Leu52 H51. We reexamined the electron density to see if there was any evidence for partial occupancy that might support the existence of rotation about the C49—C50 bond but found none. This inconsistency requires further study. If the assignment is correct, the results would indicate either a difference in the structure of the protein in solution and in the crystal or a difference in the structures of the oxidized and reduced proteins.

The combined 1D and 2D NOE data provide the first complete observation of the carbon-bound cysteine proton resonances of a 2Fe-2S* ferredoxin (the cysteine 'H's are excluded because the data were collected in H2O). The only ambiguity remaining in the assignments is whether peak b corresponds to cysteine C (Cys49) and peak d to cysteine D (Cys79) or vice versa (Dugad et al., 1990). Since all the cysteine protons are included in the above assignments, peak k (12.57 ppm, Figure 2), which shows a Curie-type contact shift, must correspond to some other residue, most probably one that is hydrogen-bonded to the iron-sulfur cluster. The four NOE cross peaks (3a–d) arise most likely from nearby protons on the same residue. Of the residues that show potential H-bonds to the iron-sulfur cluster in the X-ray structure of Anabaena 7120 ferredoxins (Ser46, Arg42, Ala43, Gly44, Ala45, Thr48, Ser47, Thr49, Val10), all but Arg42 and Val10 can be ruled out as candidates because their spin systems contain too few protons. Since both Arg42 and Val10 are potentially hydrogen-bonded to the iron-sulfur cluster by their backbone amide group, we must consider the possibility that the amide proton is protected and may not have exchanged with deuterium in the solvent. Thus we need to consider three possibilities for the assignment of peak k: the IH* of Arg42, the IH* of Arg42, or the IH* of Val10. Upon examining the distances in the Anabaena 7120 X-ray structure, the best fit is provided by assigning peak k to the IH* of Arg42. With this assignment, all four observed NOE's correspond to distances less than 3.0 Å (Table I); with either of the other two assignment possibilities, one or more of the observed NOE's would have to correspond to a distance in the X-ray structure of over 4.0 Å.

Electron self-exchange is slow on the NMR time scale when plant-type ferredoxins are reduced with dithionite but fast on the NMR time scale when methyl viologen is added (Skjeldal et al., 1990). The 1D spectrum of Anabaena 7120 ferredoxin half-reduced with dithionite appears as the superposition of spectra of the oxidized and reduced forms (Skjeldal et al., 1990). The 2D exchange spectrum obtained with a mixing time of 5 ms (Figure 3) shows six additional cross peaks (enclosed by boxes) that are not present in similar spectra of fully reduced or oxidized protein (Figure 2). These peaks arise from chemical exchange and correlate with chemical shifts of protons in the two redox states. One cross peak links peak e (43.3 ppm) to a resonance at 14.8 ppm. A second cross peak links peak j (18.0 ppm) to a resonance at 9.0 ppm. Peaks e and j are assigned to IH* atoms in the two cysteines that ligate Fe(II) in reduced ferredoxin. The 2D exchange results show that the signals from the IH* atoms of the cysteines ligated to Fe(III) shift upfield when the other iron is oxidized (Table III).

CONCLUSION

Hyperfine NMR peaks, which arise from what usually are the most interesting parts of the paramagnetic protein, have been difficult to assign, largely because it appeared that they could not be studied by conventional 2D NMR methods (Oh & Markley, 1990). The present results indicate that two-dimensional magnetization exchange spectroscopy can be adapted to hyperfine resonances in proteins by the use of short
mixing times and recycle times. The choice of the mixing time
depends on the relaxation rates of the protons undergoing
magnetization exchange; a theoretical analysis is provided by
Cheng (1991). Interproton distances of about 3 Å or less can
be sampled by this approach (Table I). Spin diffusion effects
should be negligible at the mixing times used. We expect that
the optimized NOESY methods used here will be applicable
to hyperfine resonances of other paramagnetic proteins. The
hyperfine signals of heme proteins have longer relaxation times
and can be studied with longer NOESY mixing times (Wu
et al., 1991).

The 2D experiments complement and extend 1D NOE re-
sults obtained previously on two other plant-type ferredoxins
(Dugad et al., 1990). One limitation of the 2D approach stems
from the wide chemical shift range that needs to be covered.
In the present studies, the best results were obtained when the
spectral window was limited to about 36 kHz in each di-

dimension (60 ppm at 600 MHz). We relied on the 1D results
for information about the spectral region not covered. For
investigations of the more crowded regions, 2D exchange
spectroscopy has important advantages over 1D spectroscopy.
Only one of the connectivities between the hyperfine and
diamagnetic spectral regions found by 2D NOESY (Table I)
had been observed in the 1D NMR study: the short distance
between a Cys$^\text{H1}$ (peak f) and a methyl peak (peak m)
(Dugad et al., 1990).

There is a strong theoretical basis (Dunham et al., 1971;
Banci et al., 1990) for the division of hyperfine resonances from
reduced plant-type ferredoxins into two classes on the basis of
their temperature dependence: Curie-type signals from groups
that interact with Fe(III) and anti-Curie-type signals from
groups that interact with Fe(II). The more detailed assignments
of reduced Anabaena 7120 vegetative ferredoxin (Tables II and III)
were derived here from comparison of the NMR data with the X-ray
structure (Figure 4) of oxidized Anabaena ferredoxin (Rypniewski et al., 1991). The observed pattern of NOE's shows that the iron ligated to Cys$^\text{46}$ and Cys$^\text{49}$ (Fe1 of the X-ray structure) is predominantly Fe(II)
and that the iron ligated to Cys$^\text{46}$ and Cys$^\text{79}$ (Fe2) is predominantly Fe(III). This result is consistent with interpretations of the X-ray data on oxidized ferredoxins, which show Fe1 to be more exposed and involved in more potential hydrogen-bonding interactions than Fe2 (Tsukihara et al., 1981; Rypniewski et al., 1991).

The NMR assignment method used here relies heavily on
the ferredoxin X-ray structure. Since the NMR assignments
are of reduced ferredoxin in solution and the X-ray structure
is of oxidized ferredoxin in a crystal, implicit assumptions are
that the residues near the iron–sulfur cluster do not change
their positions significantly on reduction of the protein and
that the solution and crystal structures are very similar. Since
the diamagnetic resonances of Anabaena 7120 ferredoxin do
not undergo large chemical shift changes when the redox state
is changed (unpublished results from this laboratory), the
protein appears not to undergo a large conformational change
upon reduction. The earlier assignments (Dugad et al., 1990)
were made by reference to an unrefined structure of a closely
related ferredoxin. They reported an NOE peak similar to
1a (Figure 2). If they had used the more highly refined X-ray
structure used here (Rypniewski et al., 1991), they would have
obtained the same assignments of the Cys$^\text{41}$ and Cys$^\text{46}$ spin
systems reported here rather than the reverse.

Our future aims are to assign the NMR signals by methods
that are independent of a crystal structure and to compare the
structures of the ferredoxin in the oxidized states by X-ray
and NMR methods. It should be possible to test the NMR
assignments by selective isotopic labeling; in preparation for
such studies, we are optimizing the overexpression of the
Anabaena 7120 vegetative ferredoxin gene (Alam et al., 1986)
in Escherichia coli.

The present results demonstrate that the non-cysteine proton that
gives rise to peak k in reduced ferredoxin experiences
appreciable unpaired spin density. The weak Curie temper-

ature dependence of peak k resonance (Figure 1C) suggests
that it is influenced slightly more by the spin on Fe(III) than
that on Fe(II). Peak k has been assigned tentatively to the
$^1\text{H}$ of Arg$^{42}$ on the basis of comparisons of NOE cross peaks
in the spectrum of reduced ferredoxin with the crystal structure
of oxidized ferredoxin. The mechanism of electron delocali-

zation presumably is a hydrogen bond between the Arg$^{42}$ $^1\text{H}$
and the cluster. The X-ray structure of the oxidized ferredoxin
shows the Arg$^{42}$ N within hydrogen-bonding distance (3.14 Å)
to one of the inorganic sulfur atoms (S1) (Rypniewski et al., 1991).
Assuming that the assignment of peak k to the $^1\text{H}$ of Arg$^{42}$ is correct, then its hyperfine shift in oxidized ferre-
doxin is 1.4 ppm in oxidized ferredoxin (5.8 ppm observed shift
minus 4.4 ppm for the diamagnetic chemical shift of an argi-
ine $^1\text{H}$) and 8.0 ppm in reduced ferredoxin (12.4 ppm
observed shift minus 4.4 ppm for the diamagnetic chemical
shift of an arginine $^1\text{H}$). The magnitude of the hyperfine shift

\begin{figure}
\centering
\includegraphics[width=\textwidth]{figure4}
\caption{Stereochemical view of the iron–sulfur cluster and selected surrounding residues of oxidized Anabaena 7120 vegetative ferredoxin. From the 1.5-Å refinement (manuscript in preparation) of the X-ray structure of this ferredoxin (Rypniewski et al., 1991). Fe1 is the iron ligated to Cys$^\text{41}$ and Cys$^\text{46}$; Fe2 is the iron ligated to Cys$^\text{46}$ and Cys$^\text{49}$.}
\end{figure}
of peak $k$ in reduced ferredoxin is intermediate between those of the $^{1}H$'s of the cysteines ligated to Fe(II) and Fe(III).

Additional evidence for electron delocalization onto at least one nitrogen comes from observation of hyperfine-shifted $^{14}$N signals in spectra of *Anabaena* 7120 ferredoxin uniformly labeled with $^{15}$N (Oh & Markley, 1990b). Also, a nitrogen component of the ENDOR signal of *Anabaena* 7120 vegetative ferredoxin has been identified by comparing the spectrum of [U-$^{15}$N]ferredoxin with that of the protein at natural isotopic abundance (A. L. P. Houseman, B.-H. Oh, M. C. Kennedy, L. Fan, M. M. Wester, R. Gurbiel, H. Beinert, J. L. Markley, and B. M. Hoffman, manuscript in preparation). It will be interesting to probe these effects (and to test the NMR assignment of peak $k$) by selective incorporation of $^{15}$Narginine into the ferredoxin.

It may be significant that Arg$^{42}$ is conserved in 36 of 38 sequences of low potential (plant-type) ferredoxins (Matsubara & Hase, 1983); the two exceptions are ferredoxins II from two species of *Equisetum* (horsetail) in which residue 42 is glutamic acid. By contrast, residue 42 is histidine in *Anabaena* 7120 heterocyst ferredoxin (Böhme & Haselkorn, 1983), which participates in electron transfer in the nitrogen-fixation system but not in the photosynthetic electron transport system (Böhme & Schrautemeier, 1987). The homologous residue in all sequenced vertebrate ferredoxins (adenodoxins) is glutamic acid (Mittal et al., 1988). The electronic properties of the latter two proteins differ from those of the photosynthetic ferredoxin. Site-directed mutagenesis can be used to determine whether residue 42 plays a role in these differences.

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REFERENCES


