

Extended X-ray Absorption Fine Structure and Nuclear Resonance Vibrational Spectroscopy Reveal that NifB-co, a FeMo-co Precursor, Comprises a 6Fe Core with an Interstitial Light Atom

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Abstract: NifB-co, an Fe–S cluster produced by the enzyme NifB, is an intermediate on the biosynthetic pathway to the iron molybdenum cofactor (FeMo-co) of nitrogenase. We have used Fe K-edge extended X-ray absorption fine structure (EXAFS) spectroscopy together with ⁵⁷Fe nuclear resonance vibrational spectroscopy (NRVS) to probe the structure of NifB-co while bound to the NifX protein from *Azotobacter vinelandii*. The spectra have been interpreted in part by comparison with data for the completed FeMo-co attached to the NafY carrier protein: the NafY:FeMo-co complex. EXAFS analysis of the NifX:NifB-co complex yields an average Fe–S distance of 2.26 Å and average Fe–Fe distances of 2.66 and 3.74 Å. Search profile analyses reveal the presence of a single Fe–X (X = C, N, or O) interaction at 2.04 Å, compared to a 2.00 Å Fe–X interaction found in the NafY:FeMo-co EXAFS. This suggests that the interstitial light atom (X) proposed to be present in FeMo-co has already inserted at the NifB-co stage of biosynthesis. The NRVS exhibits strong bands from Fe–S stretching modes peaking around 270, 315, 385, and 408 cm⁻¹. Additional intensity at ~185–200 cm⁻¹ is interpreted as a set of cluster “breathing” modes similar to those seen for the FeMo-cofactor. The strength and location of these modes also suggest that the FeMo-co interstitial light atom seen in the crystal structure is already in place in NifB-co. Both the EXAFS and NRVS data for NifX:NifB-co are best simulated using a Fe₆S₉X trigonal prism structure analogous to the 6Fe core of FeMo-co, although a 7Fe structure made by capping one trigonal 3S terminus with Fe cannot be ruled out. The results are consistent with the conclusion that the interstitial light atom is already present at an early stage in FeMo-co biosynthesis prior to the incorporation of Mo and R-homocitrate.

Introduction

All life depends on the input of the element nitrogen into the biosphere by biological nitrogen fixation performed only by diazotrophic microorganisms. These organisms achieve N₂ fixation using at least one member of the nitrogenase family of metalloenzymes, with the Mo-dependent enzymes being the most widely distributed and best studied, although V-dependent and Fe-only forms are also well-known. The active site of Mo-nitrogenase, which is contained within the so-called MoFe protein, is a remarkable [7Fe-9S-Mo-X-homocitrate] cluster, called the iron–molybdenum cofactor (FeMo-co). This cluster both binds the substrate N₂ and catalyzes the chemically difficult reduction of the N₂ triple bond to form NH₃.¹ The V-

nitrogenases and Fe-nitrogenases contain similar cofactors where the Mo is thought to be substituted by V or Fe, forming FeV-co and FeFe-co, respectively.²

The structure of FeMo-co (Figure 1) can be viewed as two Fe–S cubane clusters, each missing one bridging S, making [3FeMo-3S] and [4Fe-3S] components that are joined with three extra bridging S so that six of the seven Fe atoms of FeMo-co form a trigonal prism core. The cluster is anchored to the Mo-nitrogenase polypeptide by a cysteine at the Fe terminus and by a histidine at the Mo end. The Mo is also coordinated by R-homocitrate, a bidentate carboxylate ligand that is essential for N₂ reduction activity. Much interest has been generated by the most recent crystal structure for Mo-nitrogenase.³ It was proposed that, after correction for Fourier truncation artifacts, the electron density at the center of the FeMo-co could be assigned to a single light interstitial atom, namely C, N, or O.³

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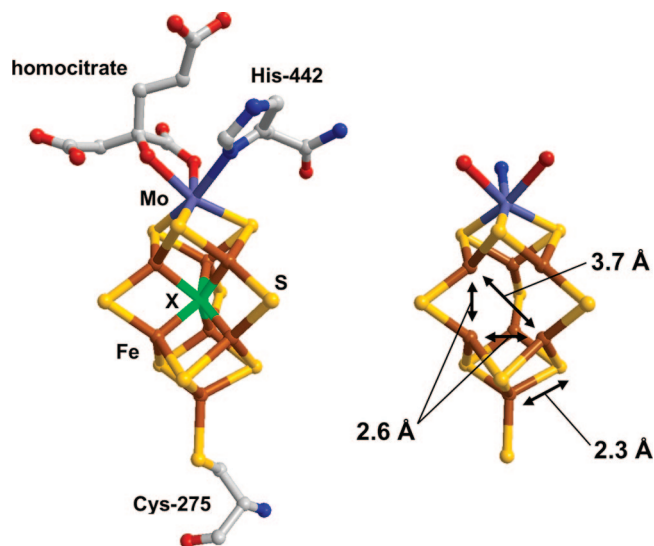


Figure 1. The structure of FeMo-co: (left) the structure as determined by X-ray crystallography on the Mo-nitrogenase enzyme (Protein Data Bank: 1MIN³); (right) the Mo-7Fe-9S core showing the 2.3 Å Fe–S, 2.6 Å short Fe–Fe, and 3.7 Å long Fe–Fe distances probed by EXAFS spectroscopy. Color code: Fe (brown), S (yellow), O (red), N (blue), interstitial atom X (green). The central atom has been omitted in the right-hand structure for clarity.

As the element of this atom is unknown it is commonly denoted as “X”. The presence of this atom has been supported by theoretical studies⁴ as well as recent analyses of nuclear resonance vibrational spectroscopy (NRVS) data.⁵

The genetic basis for the Mo-dependent nitrogenase is the *nif* gene cluster, which not only contains the structural genes encoding the nitrogenase component proteins, but importantly also codes for enzymes and chaperones which cooperate to biosynthesize FeMo-co and the other unique metallo-cofactors found in the nitrogenase systems. These Nif proteins make up these biosynthetic pathways by performing an intriguing and complex series of bioinorganic reactions using Fe and S together with Mo or V. Figure 2 illustrates our understanding of the Nif catalyzed biosynthetic pathways for Mo-nitrogenase. However, the detailed chemistry of these pathways is currently undetermined.⁶

A key metabolic intermediate in the biosynthesis of FeMo-co is NifB-co, a low-molecular weight Fe–S cluster that is the product of *S*-adenosyl methionine mediated radical (SAM-radical) chemistry catalyzed by the 108 kDa dimeric enzyme NifB.⁷ In addition to being a precursor for synthesis of FeMo-co, NifB-co is also hypothesized to be the precursor for the synthesis of FeV-co and FeFe-co, suggesting that NifB-co forms the core portion that is common among all three nitrogenase active site cofactors.⁶ The essential natures of NifB-co and the SAM-radical enzyme NifB are highlighted by the inability for diazotrophic growth upon deletion of the *nifB* gene in *N₂* fixing organisms such as *Azotobacter vinelandii* or *Klebsiella pneumoniae*. Genetic disruption of FeMo-co biosynthetic steps

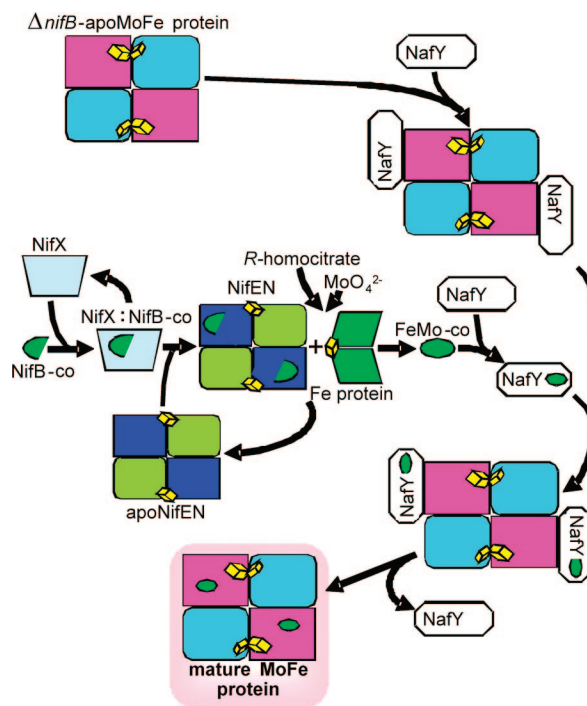


Figure 2. Proposed scheme for FeMo-co biosynthesis showing the proposed roles of NifB, NifX, NifEN, NifH, NafY, and the MoFe protein of Mo-nitrogenase.

downstream of NifB also has deleterious effects on diazotrophic growth, but causes the cellular accumulation of NifB-co. The NifB-co Fe–S cluster can be isolated from a *nifN* mutant strain of *K. pneumoniae*. Isolated NifB-co can be used in vitro to complement a cellular extract from a $\Delta nifB$ strain, which is incapable of FeMo-co synthesis, and permits the synthesis of FeMo-co producing a functional nitrogenase. Isolated NifB-co does not contain Mo required for FeMo-co synthesis, but is thought to contain the majority of the Fe and S components of FeMo-co. NifB-co is subsequently processed and converted to FeMo-co by the combined activities of NifEN and NifH. The NifB-co cluster binds to NifEN⁸ and is subjected to further modifications that may include the addition of more Fe, the addition of more S, and the attachment of Mo and R-homocitrate.⁶

In addition to the key Nif proteins such as NifB, NifEN, and NifH, additional *nif* and non-*nif* encoded proteins enhance the efficiency of FeMo-co biosynthesis by binding and stabilizing the Fe–S cluster intermediates of the pathway or the protein components. One such protein, NafY, is a 26 kDa protein that tightly binds FeMo-co ($K_d \approx 60$ nM) to form the NafY:FeMo-co complex.⁹ Another, NifX, is a 17 kDa protein that is able to bind isolated NifB-co with high affinity ($K_d \approx 1$ μ M)⁸ and the resulting NifX:NifB-co complex can donate NifB-co to the form of NifEN that is “uncharged”, that is without precursor already bound.⁸ Interestingly, NifX can also extract a different FeMo-co precursor from “charged” NifEN. We have recently characterized this Fe–S cluster—which we have termed the “VK-cluster”—and shown that it is spectroscopically distinct from NifB-co, and that it can act as a precursor for in vitro FeMo-co synthesis.⁸ This chemistry led to the hypothesis that NifX serves

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as an intracellular carrier for NifB-co or other FeMo-co biosynthetic intermediates.

Little is known about the molecular composition and structure of NifB-co, how it is assembled by NifB, or how it is subsequently used as a substrate by NifEN. Clearly, structural information on the NifB-co Fe-S cluster is required to understand the FeMo-co biosynthetic pathway and the pertinent enzyme mechanisms. Recently, some structural information was obtained on a NifEN bound FeMo-co precursor using EXAFS difference spectroscopy of charged and uncharged NifEN¹⁰ and these authors identified structural elements of partially assembled FeMo-co. However it is not clear whether this study addressed bound NifB-co, the VK cluster, or some other FeMo-co precursor. To fully elucidate the bioinorganic chemistry underpinning diazotrophy, additional study of purified NifB-co samples seems necessary.

In this paper, we use Fe K-edge extended X-ray absorption fine structure (EXAFS) spectroscopy, a well-developed technique,¹¹ together with the more novel ⁵⁷Fe nuclear resonance vibrational spectroscopy (NRVS)^{12,13} to probe the structure of NifB-co bound to NifX from *A. vinelandii*. We use EXAFS and NRVS measurements on the NafY:FeMo-co complex as a control to help calibrate and interpret the NifX:NifB-co results. The two techniques are shown to be highly complementary. EXAFS spectroscopy yields the number, distance, and approximate atomic number (*Z*) of the atoms immediately surrounding the absorbing atom. By contrast, ⁵⁷Fe NRVS analysis provides the ⁵⁷Fe-specific partial vibrational density of states (PVDOS). This spectrum extends from approximately 10–500 cm⁻¹, and it is a direct measure of the motion of the resonant ⁵⁷Fe nucleus along the direction of the incident X-ray beam in particular normal modes.¹²

Experimental Section

Protein Purification and Sample Preparation. NifX was overexpressed in *E. coli* and purified using recently described procedures.⁸ NifB-co was produced in *K. pneumoniae* UN1217 and purified using a modified larger scale version of the procedure described by Shah and co-workers.¹⁴ All manipulations of the protein and sample preparation were performed under strict anaerobic conditions using crimp-sealed vials and gastight syringes or inside an anaerobic (<5 ppm O₂) glovebox (Coy) with a 95% N₂/5% H₂ environment.

Two different NifX:NifB-co samples were prepared. The first contained ⁵⁶Fe (natural abundance) and was studied by EXAFS spectroscopy only. This was prepared by binding NifB-co (1.2 μmol total Fe) to phenyl sepharose equilibrated in 50 mM MOPS, pH 7.2, 2 mM Na₂S₂O₄, washed free of detergent, and then eluted by applying 13 mg of purified NifX. The NifB-co-eluted from the phenyl sepharose resin bound to NifX as a complex and was concentrated by ultrafiltration to ~1 mM in protein and loaded into Lucite cells (dimensions: 3 × 20 × 2 mm³). The sample was prepared without use of a glassing agent such as 10% glycerol as it was feared that this could disrupt the cluster. The second sample contained the ⁵⁷Fe necessary for NRVS studies and this was studied by both NRVS and EXAFS. Incorporation of ⁵⁷Fe was achieved by replacing the regular source of Fe with 11 μM ⁵⁷Fe (98%, Isotec) in the *K. pneumoniae* growth medium. A sarkosyl membrane extract

containing ⁵⁷Fe NifB-co was prepared from 190 g of ⁵⁷Fe-enriched cells and was applied to a column of Sephacryl 200 (S200), which NifB-co binds. The column was washed free of all proteins and then washed with detergent-free buffer. Pure and metal-free NifX (4 mg) was applied to the NifB-co bound S200 column and the NifB-co was eluted as the NifX:(⁵⁷Fe) NifB-co complex. The NifX:(⁵⁷Fe)NifB-co samples were concentrated to ~5 mM in protein by ultrafiltration and loaded into 3 × 7 × 1 mm³ (interior dimensions) Lucite cuvettes. The final Fe concentration, as estimated from the Fe K-emission and NRVS resonance peak count rates was approximately 30 mM, consistent with the sample preparation and a stoichiometry of 6–7 Fe per NifX molecule.

The NafY:FeMo-co complex was prepared using a modification of previously described procedures.⁹ FeMo-co was prepared from *A. vinelandii* MoFe protein¹⁵ and *A. vinelandii* NafY was purified⁹ as previously described. To prepare the sample of NafY:FeMo-co for EXAFS, 1.4 μmol of NafY was diluted with 50 mM MOPS, pH 7.4, and 2 mM Na₂S₂O₄ in a stirred ultrafiltration cell fitted with a YM10 membrane (Millipore). FeMo-co (0.7 μmol total), dissolved in NMF solution, was added to the diluted NafY in several portions to limit the NMF concentration (<3%), with repeated concentration and dilution with every addition of FeMo-co. The NMF concentration was decreased by an additional round of dilution and concentration to a volume of 550 μL. The solution of NafY:FeMo-co was then passed through a Sephadex-10 size exclusion column (1 × 15 cm²) to remove residual NMF, concentrated to 60 μL, and frozen in a Lucite cell (dimensions: 3 × 20 × 2 mm³). The sample for NRVS was prepared using essentially the same method using ⁵⁷Fe MoFe protein and 3 × 7 × 1 mm³ Lucite cuvettes

X-ray Absorption Spectroscopy. Fe K-edge X-ray absorption data were measured at Stanford Synchrotron Radiation Laboratory beamline 9-3, with a Si 220 double-crystal monochromator and two rhodium-coated mirrors: one flat premonochromator mirror for harmonic rejection and vertical collimation and one toroidal postmonochromator mirror for focusing. Fluorescent X-rays were measured using a 30-element Ge fluorescence detector (Canberra Industries), fitted with Soller slits and a Mn filter to minimize the relative contribution of scattered radiation. An Oxford Instruments CF1208 liquid helium cooled sample cryostat was used to maintain the sample temperature at 9 K. The X-ray energy was calibrated using the first and major inflection point of a standard Fe foil set as 7112 eV. This was measured at the same time as the sample spectrum using two ion chambers positioned downstream of the cryostat. To minimize the effect of X-ray photoreduction, the sample was moved after each scan so that the beam irradiated a different spot on the sample for each scan; typically a total of 3–4 spots were available for each sample, and 5–11 scans were collected requiring reuse of each spot. The Fe K-edge structure and position were monitored to ensure no photochemistry.

EXAFS Data Analysis. EXAFS data were analyzed using the EXAFSPAK software suite.¹⁶ As the samples contained no glassing agent, it was first necessary to rigorously screen the scans from the individual detector elements to eliminate any diffraction artifacts. Curve fitting used the EXAFSPAK program OPT, with single-scattering phase and amplitude functions calculated using FEFF 7.0.¹¹ The NafY:FeMo-co complex was used as a reference sample to provide experimentally determined estimates of Debye–Waller factors.

NRVS Measurements. NRVS is a synchrotron radiation technique where a highly monochromatic ($\Delta E \approx 1$ meV) X-ray beam is scanned through the nuclear resonance.^{12,17} In addition to the familiar “zero phonon” (recoil-free) Mössbauer resonance, there

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are additional transitions corresponding to nuclear excitation combined with excitation (Stokes) or de-excitation (anti-Stokes) of vibrational modes. The only requirement for NRVS intensity is motion of the resonant nucleus (in this case ^{57}Fe) along the direction of the incident X-ray beam in the given normal mode. The details of NRVS theory have been detailed elsewhere.^{12,18} Essentially, the single phonon NRVS excitation probability $S_1(\bar{\nu})$ is directly proportional to the Fe partial vibrational density of states (PVDOS) $D_{\text{Fe}}(\bar{\nu})$.¹⁹ The PVDOS function $D_{\text{Fe}}(\bar{\nu})$ arises from convolution of a line shape $L(\bar{\nu} - \bar{\nu}_\alpha)$ with the Fe mode composition factor, e_{Fe}^2 , which represents the fraction of kinetic energy due to Fe for normal modes α at resonant energies $\bar{\nu}_\alpha$.^{18,20}

^{57}Fe NRVS spectra were recorded using published procedures¹² at beamline BL09XU at SPring-8, Japan.²¹ The flux was $\sim 3 \times 10^9$ in a 1.1 meV bandwidth (1 meV $\equiv 8.0655 \text{ cm}^{-1}$). During NRVS measurements, the samples were maintained at $\sim 20 \text{ K}$ using a liquid He cryostat. Exact temperatures for individual spectra were calculated using the ratio of anti-Stokes to Stokes intensity according to: $S(-E) = S(E) \exp(-E/kT)$. Spectra were recorded between -20 and 80 meV in 0.25 meV steps. Nuclear fluorescence and Fe K fluorescence (from internal conversion) were recorded using an APD array.²² Each scan took about 40 min, and all scans were added and normalized to the intensity of the incident beam. The reported NRVS spectrum represents the average of 16 scans.

Normal Mode Calculations. Normal mode calculations for NifB-co candidate structures were conducted with a Urey–Bradley force field using a modification of program “Vibrat”.²³ A QR algorithm was used for finding eigenvectors.²⁴ The structures used the distance constraints from the EXAFS analysis. The mode composition factor $e_{j\alpha}^2$ for mode α and atom j was calculated by^{13,18,25}

$$e_{j\alpha}^2 = \frac{m_j r_{j\alpha}^2}{\sum_j m_j r_{j\alpha}^2}$$

where m_j and $r_{j\alpha}$ are the mass of atom j and its eigenvector in mode α , respectively. For comparison with the experimental spectra, Gaussian functions were calculated centered around each ω_α , the frequency of mode α , using a 7 cm^{-1} full width at half-maximum and an amplitude proportional to $e_{j\alpha}^2$. The partial vibrational density of states for atom j was then obtained by summing these Gaussian functions over all modes.

Results

We have measured EXAFS and NRVS spectra of the NifX:NifB-co complex. We have also measured analogous data for NafY:FeMo-co. As this protein complex contains the completed FeMo-co center it provides an excellent control sample for the

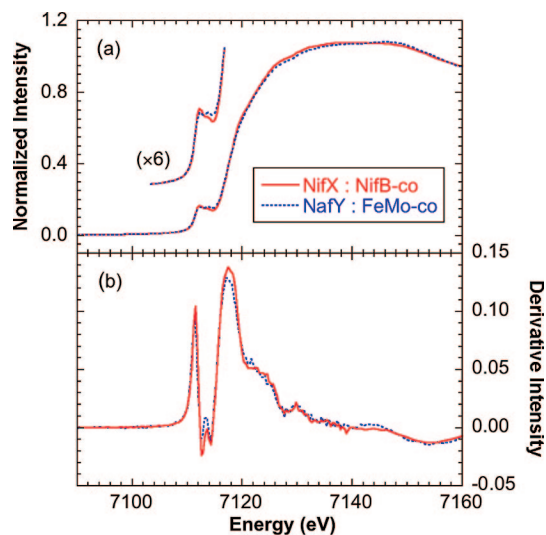


Figure 3. Fe K-edge near-edge spectra of NifX:NifB-co (solid line) compared with NafY:FeMo-co (broken line): (a) absorption spectra and (b) first derivative spectra.

interpretation of the NifX:NifB-co data. We present first the EXAFS and near-edge spectra, together with their analysis, followed by the NRVS results.

EXAFS and Near-Edge Spectra. The two samples, NifX: (^{56}Fe) NifB-co and NifX: (^{57}Fe) NifB-co, gave very similar X-ray absorption spectra. The data presented here were obtained from the ^{57}Fe sample that was also used for NRVS measurements. Figure 3 shows the Fe K-edge near-edge spectrum and first derivative from NifX:NifB-co and compares these with those from NafY:FeMo-co known to be in the resting $S = 3/2$ $\text{Na}_2\text{S}_2\text{O}_4$ reduced state. Both spectra exhibit the low-intensity broad rounded edge typical of iron–sulfur clusters together with small pre-edge features at $\sim 7112.2 \text{ eV}$ that arise from $1s \rightarrow 3d$ transitions. The data sets from the NifX:NifB-co and NafY:FeMo-co are almost superimposable, with only subtle differences in both the pre-edge and edge regions. This indicates that the oxidation states and ligand environments of the Fe atoms in NifX:NifB-co and NafY:FeMo-co are very similar.

We also searched the Mo K-edge region for evidence of cluster bound Mo in NifX:NifB-co. These measurements (not shown) conclusively demonstrated the absence of any measurable Mo in NifX:NifB-co, corroborating previous chemical analyses that indicated NifB-co does not contain Mo.¹⁴

Figure 4 presents the EXAFS spectrum and Fourier transform for NifX:NifB-co together with a representative fit using the 6Fe model and compares these data to those from NafY:FeMo-co fitted to the crystallographically determined FeMo-co structure in Figure 1. Initial inspection of the EXAFS for NifX:NifB-co suggests a strong similarity to the well-examined EXAFS of FeMo-co,²⁶ and comparisons between these spectra are key

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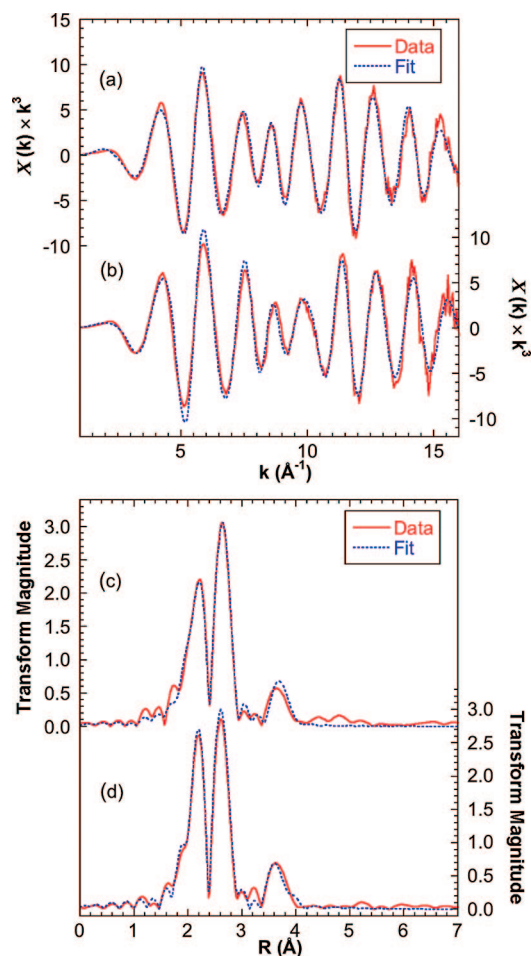


Figure 4. Fe K-edge EXAFS of NifX:NifB-co with fits taken from the 6Fe model compared with data and fit of NafY:FeMo-co: (a) unsmoothed k^3 -weighted EXAFS of NifX:NifB-co; (b) unsmoothed k^3 -weighted EXAFS of NafY:FeMo-co; (c) NifX:NifB-co Fourier transforms; (d) NafY:FeMo-co Fourier transforms. In each plot data is given as a solid line while the fit is a broken line. Fitting parameters are given in table 1. A deconvolution of the NifX:NifB-co 6Fe model fit is shown in Figure S1 and Figure S2 of the Supporting Information.

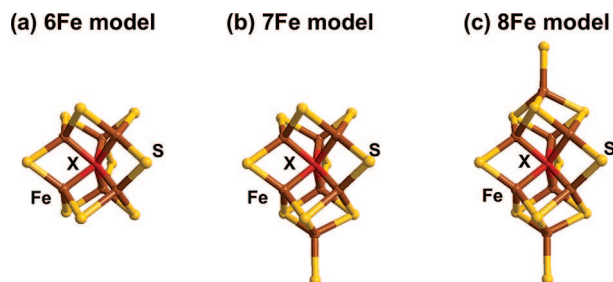


Figure 5. Proposed models for NifB-co as described in the text: (a) 6Fe, (b) 7Fe, (c) 8Fe. Color code: Fe (brown), S (yellow), interstitial atom X (red).

to interpreting and understanding the differences between FeMo-co and its biosynthetic precursor, NifB-co. Curve fitting analysis reveals that the two main features in the Fourier transform in Figure 4c correspond to a set of S ligands at ~ 2.3 Å and a set of Fe next nearest neighbors at ~ 2.6 Å. Importantly, a further set of Fe–Fe interactions at 3.7 Å is also clear. These interactions are very similar to those well-known from FeMo-co (Figure 4d) and their origin in this cofactor is illustrated in Figure 1. The crystallographically determined structure of FeMo-

co in MoFe protein can be viewed as a trigonal prism of 6 Fe atoms, with each of the nine sides bridged by S, and each triangular S-bridged face capped with either Fe or Mo. The 2.3 Å Fe–S interaction is normal for Fe–S bonds. The “short” Fe–Fe interactions at ~ 2.6 Å are the sides of the trigonal prism as well as the distance of the capping Fe from its neighboring core Fe. The “long” 3.7 Å Fe–Fe distance arises from interactions across the square faces of the [6Fe-9S] trigonal prism. Such an interaction has so far only been observed in Fe–S centers for FeMo-co, and it can therefore be viewed as a strong indicator for the presence of the [6Fe-9S] core in NifB-co.

To rationalize these data, we constructed three working models, containing 6Fe, 7Fe, and 8Fe atoms, all based on the FeMo-co structure (Figure 5). The 6Fe model is simply a D_{3h} symmetry [6Fe-9S] trigonal prismatic core similar to that of FeMo-co. The 7Fe and 8Fe models were formed by capping one or both of the trigonal 3S faces of the [6Fe-9S] core. With coordination numbers constrained by these models, the short Fe–S, short Fe–Fe, and long Fe–Fe distances and Debye–Waller factors were then optimized to fit the experimental data. The resulting values for each model are summarized in Table 1 and the quality of the fits is illustrated in Figure 4, which includes the 6Fe model fit. All three models give reasonable fits with fitted short Fe–Fe and long Fe–Fe distances about 0.02 Å larger than those of NafY:FeMo-co, but the 6Fe D_{3h} model gives the best fit to the experimentally observed data. As expected, the fitting process compensates for increasing the iron content of the cluster by changing the highly correlated Debye–Waller factor. All the Debye–Waller factors for each fit are within the sensible range for their bond types; that for the short Fe–Fe interaction for the 6Fe cluster at 0.0032 Å² is exactly the minimum reasonable value.³⁰ This low value suggests a highly symmetric cluster with little variation between the 9 short Fe–Fe distances and supports the 6Fe D_{3h} model for NifB-co. We note that the 6Fe and 7Fe models are more consistent with the values from the completed FeMo-co compared to the 8Fe model as well as giving the better fit.

We investigated whether introducing a small additional longer Fe–S or Fe–Fe interaction could significantly improve the quality of the fits. For Fe–Fe, we rationalized this search by postulating various movements of a single Fe away from the center of the trigonal prism. We found that we could improve the fit by including a small additional Fe–Fe interaction at ~ 2.9 Å and more significantly a split long Fe–Fe with a contribution at almost 4.0 Å which could be rationalized in terms of the distortion of a single Fe atom. However, at these distances the EXAFS from both these increased Fe–Fe interactions are almost exactly out of phase with the corresponding shorter distances, and hence there are significant risks of false minima with these distances in the curve fitting. In addition, in these fits, the Fe–Fe Debye–Waller factors are close to or below the reasonable minimum. Hence, at this time we believe the structure is best modeled by the fits in Table 1.

(30) Debye–Waller factors are frequently used (as they are here) as floating parameters to optimize the best fit to EXAFS data. They arise from the variation of distances for the given scatterer–backscatterer pair and hence have contributions from both molecular vibrations and unresolved structural variation. Clearly the smallest possible Debye–Waller factor for a given interaction is that arising from the vibrational component alone. For Fe–Fe interactions in a Fe–S cluster, this can be estimated from a protein sample containing a single Fe–Fe interaction, such as a [2Fe-2S] containing protein. Such measurements indicate a Debye–Waller of 0.0034 Å², so for this work we assign a value 0.0032 Å² as the “reasonable minimum”.

Table 1. EXAFS Curve Fitting Parameters for the NifX:NifB-co Spectrum Compared to Those Measured for NafY:FeMo-co^a

	NafY:FeMo-co			6Fe Model			7Fe Model			8Fe Model		
	<i>N</i>	<i>R</i> (Å)	σ^2 (Å ²)	<i>N</i>	<i>R</i> (Å)	σ^2 (Å ²)	<i>N</i>	<i>R</i> (Å)	σ^2 (Å ²)	<i>N</i>	<i>R</i> (Å)	σ^2 (Å ²)
Fe–S	3.10	2.251	0.00328	3.00	2.263	0.00424	3.00	2.264	0.00404	3.00	2.263	0.00383
Fe–Fe (short)	3.40	2.627	0.00456	3.00	2.655	0.00323	3.43	2.655	0.00385	3.75	2.654	0.00431
Fe–Fe (long)	1.70	3.726	0.00562	2.00	3.740	0.00646	1.71	3.742	0.00533	1.50	3.742	0.00436
Fe–X(N)	0.85	2.000	0.00199	1.00	2.046	0.00272	1.00	2.043	0.00240	1.00	2.037	0.00200
Fe–Mo	0.40	2.717	0.00350									
ΔE_0 (eV)		–12.66			–11.99			–11.81			–12.00	
<i>F</i>		0.242			0.214			0.237			0.256	

^a *N* = number of backscattering atoms; *R* = distance; σ^2 = Debye–Waller factor; ΔE_0 = threshold energy; *F* = fit quality defined as $\sqrt{[\sum(\chi_o - \chi_c)^2 k^6 / \sum \chi_o^2 k^6]}$ (χ_o = observed EXAFS; χ_c = calculated EXAFS). The backscatterer atom numbers (*N*) were fixed to the values expected for the models while *R*, σ^2 , and ΔE_0 were floated. Threshold energies (ΔE_0) were constrained to be the same for all components. The estimated uncertainties in *R*, σ^2 , and ΔE_0 were less than ± 0.01 Å, ± 0.0001 Å², and ± 0.4 eV, respectively. Phase and amplitude functions were calculated using FEFF 7.0. The scale factor used was 0.85. The Fe–X interaction was simulated assuming X = N; assuming X = C or O gave similar results but with different Debye–Waller factors.³⁰

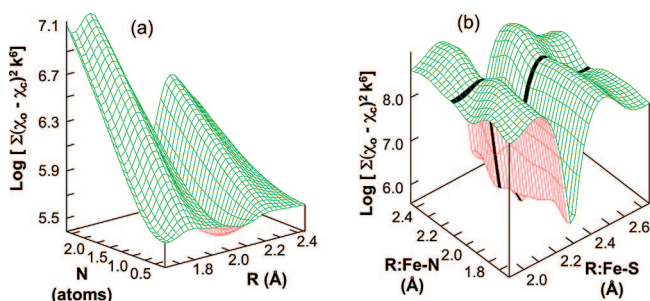


Figure 6. Search profiles from fitting the Fe K-edge EXAFS of NifX:NifB-co 6Fe model. (a) 3D plot of the search profile of number (*N*) versus distance (*R*) for the proposed Fe–N interaction showing the minimum at ~ 2.0 Å and 1 atom (χ_o = observed EXAFS; χ_c = calculated EXAFS). (b) 3D plot of the search profile of Fe–N distance versus Fe–S distance. The dark line in profile b highlights the trough due to the Fe–N interaction.

We also investigated whether the EXAFS data supports the presence of a single interstitial light atom “X” within the 6Fe trigonal prism. As noted above, the most recent crystal structure for Mo-nitrogenase³ proposed such a C, N, or O atom at the center of FeMo-co, and we also note that our analysis of the NafY:FeMo-co EXAFS spectrum presented in Figure 4 and Table 1 readily accommodates this light atom. Clearly it is of key biosynthetic interest to know whether this atom is present in NifB-co. To address the potential presence of the interstitial atom, a multidimensional EXAFS “search profile” was conducted for different distances, numbers, and atomic numbers of an additional light atom. Figure 6 illustrates two results for a Fe–N interaction in the 6Fe model. The results for the number versus distance search for a Fe–X(N) interaction (Figure 6a) shows a well-defined minimum at ~ 1 atom at 2.06 Å. Similarly, a search profile comparing the Fe–S distance and Fe–X(N) distance (Figure 6b) clearly shows two independent troughs, with a weaker Fe–X(N) trough intersecting the deep Fe–S interaction. Both these profiles clearly support the presence of a single central light atom. To be thorough, multiple curve fits both with and without a single central light atom were trialed. In every case the fit was significantly improved by including the central atom. In addition, we note that very similar profiles were obtained from the 7Fe and 8Fe models (data not shown). We emphasize that the contributions of the Fe–X(N) interaction to the fits in table 1 are significant, about 6% of the total EXAFS (see Supporting Information). We therefore conclude that the EXAFS data demonstrate the presence of a one Fe–X (X = C, N, or O) interaction for each of the 6Fe of the trigonal prism of

NifB-co, consistent with the presence of a single C, N, or O interstitial atom.³¹

In summary, analysis of the EXAFS and near-edge spectra shows that NifB-co in NifX:NifB-co is composed of a 6Fe trigonal prism core very similar in geometry to that in FeMo-co. It is also clear that the interstitial atom is already present in the NifB-co precursor for FeMo-co. The current EXAFS analysis does not resolve whether or not NifB-co also contains an additional capping Fe (Fe1 in PDB file 1MIN).

NRVS. The NRVS-derived ⁵⁷Fe PVDOS for NifX:NifB-co is presented in Figure 7 together with calculated spectra, the spectrum of NafY:FeMo-co, and the previously reported spectrum of NMF:FeMo-co (FeMo-co isolated in *N*-methylformamide (NMF)). We note that, as expected, the NafY:FeMo-co NRVS PVDOS is very similar to that of NMF:FeMo-co, as well as that previously derived for the M-center of Monitrogenase,⁵ and while there are some differences in band intensity, there is an almost one-to-one peak correspondence (Figure 7d). This indicates that the vibrational structure of the FeMo-co core is not greatly affected by its coordination and environment.

The NifX bound NifB-co NRVS PVDOS spectrum in Figure 7a is also similar to those of the completed cofactor, showing strong well-defined features in both the Fe–S stretching and bending regions. This is in contrast with the spectra from other Fe–S clusters where the strongest features are in the Fe–S stretching region and individual bending modes are not well resolved.^{27,28} The strongest PVDOS features occur in a broad-band between ~ 183 and 198 cm^{–1}. This feature has been only previously observed by NRVS in Fe–S clusters of nitrogenase and isolated FeMo-co (Figure 7d) and is considered to arise from the shaking modes of the *c*₃ symmetric Fe₆X core and is thus dependent on the presence of the interstitial light atom.⁵ A doublet at 271 and 283 cm^{–1} apparently corresponds to a similar feature observed at 267 and 290 cm^{–1} in the NRVS spectrum of a [4Fe-4S] cluster model compound which has been assigned to the Fe–S_b A₁ stretching mode under *D*_{2d} symmetry.²⁸ There are additional peaks at 315 and 364 cm^{–1}, but their overall width suggests they are composed of multiple unresolved bands. There also are clear features up to 408 cm^{–1}, which are reminiscent of the B_{2u}^b stretch modes in [2Fe-2S] clusters.²⁷ In the lower frequency region, there are S–Fe–S

(31) It is interesting to note that the fitted Debye–Waller factor was most reasonable with a N interstitial atom compared with that for a C or O atom. For example, for the 6Fe model we obtained Fe–C of 0.00061 Å², Fe–N of 0.00272 Å², and Fe–O of 0.00620 Å². A reasonable minimum value for a Fe–light-atom interaction is ~ 0.002 Å².

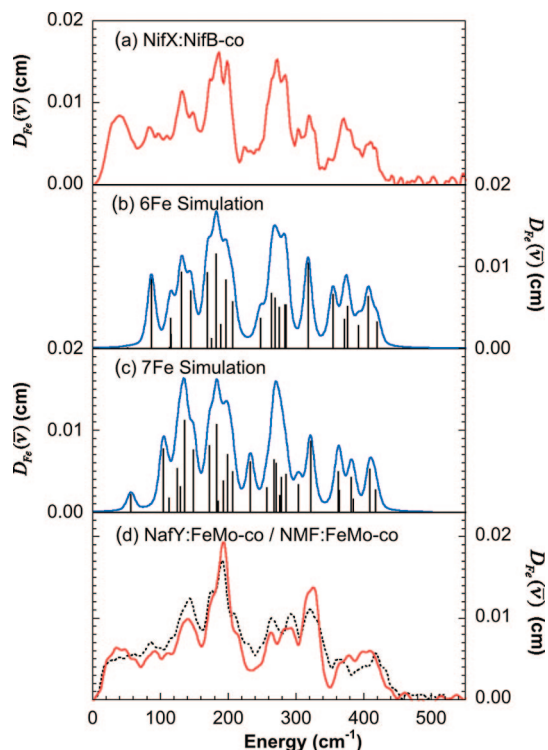


Figure 7. NRVS data and Urey–Bradley force field simulations for NifX:NifB-co compared to NafY:FeMo-co and NMF:FeMo-co: (a) data from NifX:NifB-co; (b) simulation using the 6Fe model; (c) simulation using the 7Fe model; (d) data from NafY:FeMo-co (solid line) compared with NMF:FeMo-co (broken line).⁵ The sticks in the simulated spectra show the calculated normal modes. The simulated spectra were obtained by broadening the stick spectrum with 7 cm⁻¹ Gaussian line shapes.

bend modes around 140 cm⁻¹ and significant NRVS intensity occurs between ~40 and 85 cm⁻¹ that presumably corresponds to cluster torsional modes and large-scale motion of the protein skeleton.

To quantitatively interpret the NRVS spectrum, we undertook an empirical normal-mode analysis using a Urey–Bradley force field. Cartesian coordinates for the Fe environments in NifB-co were derived from the distances in the EXAFS D_{3h} model. Initial force constants were taken from values used in the nitrogenase bound FeMo-co simulation,⁵ then adjusted to match the main peaks in the experimental spectrum. As with the EXAFS analysis, the 6Fe, 7Fe, and 8Fe models in Figure 5 were trialed. Figures 7b and 7c represent the calculated spectra obtained from the 6Fe and 7Fe models and compare them with the experimental spectrum in Figure 7a. Table 2 presents the calculated force constants. We note that to get a good fit in the 6Fe model it was necessary to differentiate the S atoms bridging the trigonal faces of the cluster (and which become tricoordinate in the completed FeMo-co) from those S bridging the two trigonal sides of the core. Despite extensive effort, it proved not possible to obtain a reasonable fit for the 8Fe model without using physically unreasonable force constants. This is demonstrated and discussed further in the Supporting Information.

Examination of Figure 7 shows good agreement between the experimental spectrum and the simulations from both the 6Fe and 7Fe models, especially over the strongest PVDOS region between 150 and 210 cm⁻¹. However, the 6Fe model gave a better fit, not only at this lower frequency region, but also over the Fe–S stretching region between 250 to 420 cm⁻¹. We note that modes below 100 cm⁻¹ mostly involve cluster torsional modes and large-scale motion of the protein skeleton, and hence

Table 2. NRVS Force Constants Used to Simulate the NifB-co Spectrum^a

	6Fe model	7Fe model
Stretching Mode (mdyn/Å)		
Fe–S ^U	1.136	1.200
Fe–S	1.065	0.960
Fe–N	0.265	0.300
Fe–Fe	0.225	0.155
Fe ^T –S		0.960
Bending Mode (mdyn Å/rad ²)		
S–Fe–S	0.130	0.140
S–Fe–S ^U	0.455	0.388
S–Fe–N	0.179	0.140
S ^U –Fe–N	0.453	0.400
Fe–N–Fe	0.056	0.250
Fe–S–Fe	0.536	0.450
Fe–S ^U –Fe	0.556	0.450
S–Fe ^T –S		0.120
Fe–S–Fe ^T		0.120
Nonbonding Mode (mdyn/Å)		
S–S	0.00	0.00
S–N	0.270	0.250
S–S ^U	0.047	0.118
S ^U –N	0.280	0.224
Stretch–Stretch Stretch–Bend Mode (mdyn/rad)		
Fe–S ^U /Fe–S ^U	0.114	0.074
Fe–S/Fe–S	0.124	0.070
Fe–N/Fe–N	0.265	0.136
Fe ^T –S/Fe ^T –S		0.113

^a S^U = trigonal face bridging S (three coordinate S atoms in 7Fe or 8Fe models). Fe^T = terminal Fe.

it is not possible to simulate this region with a cluster-based model alone.²⁹ We also attempted fits using models without the interstitial light atom, causing the strong broad band between 183 and 198 cm⁻¹ to be absent from the fit. As mentioned above, we note that we have previously assigned the analogous feature in FeMo-co as being characteristic of the interstitial light atom. We investigated routes to recover intensity in this region and found that one way was to use very large bending force constants for the Fe centered modes. This, however, is physically unrealistic and hence we conclude that the NRVS data also supports the presence of a light interstitial atom at the center of the 6Fe trigonal prism.

In summary, comparison of the NifX:NifB-co NRVS with the NRVS of FeMo-co and other Fe–S clusters, shows that the NifX bound NifB-co data has a pattern of bands similar to those of Mo-nitrogenase, NMF:FeMo-co and NafY:FeMo-co. This includes the strong signal near 190 cm⁻¹ that is diagnostic of the interstitial light atom within the 6Fe cage. Urey–Bradley force constant simulations support this observation and demonstrate that the experimental data is most consistent with the 6Fe model shown in Figure 5, although the 7Fe model cannot be completely excluded.

Discussion and Conclusions

FeMo-co is the product of a series of bioinorganic reactions performed by Nif proteins present in diazotrophic organisms. The complexities of synthesizing FeMo-co are underscored by the fact that a number of inorganic chemists have attempted to synthesize FeMo-co in vitro, but have so far not been successful. Knowledge of the structure of the NifB-co intermediate will greatly aid in understanding how nature accomplishes this chemical feat and may lead to better inorganic synthetic routes to manufacture complex Fe–S cluster moieties. To provide information for a structural model of FeMo-co, NifB-co bound

to its likely intracellular carrier protein NifX has been analyzed using the combination of two highly complementary synchrotron-based spectroscopic techniques: EXAFS spectroscopy and NRVS.³²

Taken together, the EXAFS and NRVS data show that NifB-co bound to NifX is likely to consist of a 6Fe atom cage having D_{3h} symmetry (Figure 5a), although the corresponding 7Fe model with a single capping Fe (Figure 5b) cannot be absolutely excluded. The Fe K-edge EXAFS of NifX:NifB-co shows that NifB-co is more complex than a simple Fe_4S_4 cubane, but is also distinct from FeMo-co. The observation of the long distance Fe–Fe interactions at ~ 3.7 Å, supported by the curve-fitting analysis, strongly suggests that NifB-co contains a cluster resembling the 6Fe central cage portion of FeMo-co. In addition, search profile analyses clearly indicate that the spectrum contains a Fe–X interaction at ~ 2 Å consistent with the presence of an interstitial light atom within the 6Fe cage. Measurements in the Mo K-edge region conclusively demonstrate the absence of Mo in the cluster. The Fe K-near-edge data indicate that the overall oxidation states and ligand environments of the Fe atoms in both NifX:NifB-co and NafY:FeMo-co are very similar. This is consistent with NifB-co comprising a substantially assembled FeMo-co core in the correct oxidation state. Presumably, it also indicates that capping the 6Fe core in Figure 5a with a Mo or Fe atom does not significantly affect the Fe near-edge spectrum. These conclusions are corroborated by NRVS analysis of the same sample of NifX:(⁵⁷Fe)NifB-co. The similarity of the NRVS spectrum of NifX:NifB-co to those of the completed FeMo-co is striking, especially as it includes the band at ~ 190 cm^{-1} which in FeMo-co has been assigned to modes involving the interstitial light atom within the 6Fe trigonal prism core. Vibratz simulation of the NRVS spectrum confirms this assignment. The analyses favor the 6Fe model over the 7Fe model, and clearly exclude the 8Fe model. To gain further insight into chemistry of NifB-co and how it is transformed by the FeMo-co biosynthetic pathway, it will be necessary to use EXAFS and NRVS to probe the structures of other biosynthetic precursors of the FeMo-co, such as the VK-cluster. These studies are underway and will be presented elsewhere.

(32) It is worth noting that one of the potential advantages of complementary EXAFS and NRVS studies is that a proper understanding of the normal-mode structure of a complex clusters like FeMo-co or NifB-co enables the potential use of NRVS-derived vibrational modes to estimate and constrain further the EXAFS Debye–Waller factors.

The conclusion that NifB-co contains a Fe_6X structure, which in turn becomes the central cage portion and interstitial atom of FeMo-co, rules out that the finished FeMo-co may be assembled without the X atom, which would be instead inserted at a late stage in the biosynthesis, such as initial enzyme turnover. This result is consistent with the suggestions that the subsequent enzymes involved in the FeMo-co biosynthetic pathway, NifEN and NifH, function to complete the cluster by adding the capping atoms of Mo, and perhaps the Fe, as well introducing the R-homocitrate ligand to the already constructed Fe_6S_9X core. Most interesting, however, is the implication that the function of the NifB enzyme is to perform the reaction that synthesizes this unique structure. The possible mechanistic details of this enzymatic reaction are intriguing. NifB is a SAM-radical enzyme, and it is reasonable to postulate a reaction that generates the Fe–S framework using SAM as a cosubstrate. However, the origin of the interstitial atom and the chemistry of its insertion remain real unknowns and establishing this chemistry will be the next major challenge in understanding the biosynthesis of the unique and remarkable bioinorganic catalyst, FeMo-co.

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Supporting Information Available: EXAFS deconvolution; NRVS analysis using the 8Fe model. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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