The D14C variant of *Pyrococcus furiosus* ferredoxin provides an extraordinary framework to investigate a [3Fe–4S] cluster at two oxidation levels and compare the results to its physiologic [4Fe–4S] counterpart in the very same protein. Our spectroscopic and computational study reveals vibrational property changes related to the electronic and structural aspects of both Fe–S clusters.

Iron–sulfur (Fe–S) clusters consist of iron atoms with bridging sulfides and play important roles in nature, e.g. in electron transport, as sensors, as catalytic heterometallic active sites, or through structural functions.1,2 The thermostable ferredoxin (Fd) from *Pyrococcus furiosus* (Pf) functions as an electron donor for many oxidoreductases involved in carbohydrate and peptide fermentation. Reduced Fd supplies electrons for proton reduction by a membrane bound respiratory hydrogenase (MBH) that both produces molecular hydrogen (H2) and generates an ion gradient.3 The well-studied wild-type (WT) Pf Fd contains a single [4Fe–4S] cluster and represents an excellent framework for studying Fe–S clusters. The [4Fe–4S] coordination to the protein matrix via three cysteines (C11, C17, C56) and one aspartate (D14) side-chains is unusual in comparison to other proteins commonly utilizing four cysteines, and has so far been found only in other ferredoxins and certain hydrogenases.4–6 The effects of this rare coordination with one aspartate have been scrutinized using Pf Fd site-directed mutagenesis. The redox potentials of the Pf Fd D14C and D14S mutants are, respectively, 58 and 133 mV more negative than the −368 mV of the WT protein.6,7

Pf Fd D14C is sensitive to oxidative damage, which reversibly converts its [4Fe–4S] cluster into a [3Fe–4S] cluster with a 213 mV more positive redox potential (∼155 mV) when compared to the WT protein.8 A 2.80 Å resolution X-ray diffraction (XRD) crystal structure of the ferricyanide-treated Pf Fd D14C reveals its [3Fe–4S] cofactor with the nearby Cys14 cysteine side chain facing away from the cluster (Fig. 1).8

The best-known multi-cofactor enzymes hosting a [3Fe–4S] cluster are succinate dehydrogenase and periplasmic [NiFe] hydrogenases.9,10 Their relatively high potential [3Fe–4S] clusters play an important role in electron transfer relays. In periplasmic hydrogenases, a [3Fe–4S]-to-[4Fe–4S] cluster conversion can be easily achieved by a single amino acid exchange.

Fig. 1 [3Fe–4S] cluster of the ferricyanide-treated D14C ferredoxin from *Pyroccocus furiosus*. The cofactor inorganic core is shown in ball-and-stick representation. Sulfur and iron atoms are in yellow and brown, respectively. The cysteine at position 14 is facing away from the cluster. The presented crystal structure is based on chain A from PDB entry 3PNI.8
from proline to cysteine at the missing fourth coordination site.11 This type of mutant-dependent cluster conversion (which is thus in contrast to Pf Fd D14C) results in oxygen sensitivity of the otherwise O₂-tolerant Hyd1 hydrogenase from Escherichia coli.12 A detailed understanding of redox-dependent vibrational properties of [3Fe-4S] clusters would certainly be valuable; only the oxidized WT [3Fe-4S] Pf Fd state has yielded useful resonance Raman (RR) spectra to date.13

Nuclear resonance vibrational spectroscopy (NRVS) is a relatively new synchrotron-based technique that selectively probes vibrational modes specific to Mössbauer-active nuclei.14,15 Several [2Fe-2S] and [4Fe-4S] cluster systems as well as multi-cofactor enzymes have so far been characterized by ⁵⁷Fe-NRVS.16–22 Density functional theory (DFT) has a long history of successful applications to Fe–S clusters, including those to Pf Fd and [3Fe-4S].19,21–23 In this study, we use ⁵⁷Fe-NRVS together with DFT to probe the vibrational dynamics of the [3Fe-4S] cluster in Pf Fd D14C at two oxidation levels, and compare our results to those from an earlier account on the [4Fe-4S] variant from the same protein.19

The [4Fe-4S] cluster of Pf Fd D14C protein purified and reconstituted with ⁵⁷Fe was converted by oxidation with ferri-cyanide and oxygen into a [3Fe-4S] cluster and the protein was subsequently treated with dithionite to obtain oxidized and fully reduced samples. The oxidation state and the conversion into the [3Fe-4S] cluster were confirmed by Mössbauer spectroscopy as expounded in the ESL.† The oxidized [3Fe-4S]⁺⁺ (S = 1/2) form of the cluster revealed doublets with isomer shift and quadrupole splitting corresponding to high-spin (S = 5/2) ferric (Fe⁴⁺) sites, and the reduced [3Fe-4S]⁰ (S = 2) form displayed hyperfine parameters corresponding to one ferric and two mixed-valent (Fe²⁺–³⁺) ions (Table S1 and Fig. S1†). Our DFT results employing the broken-symmetry (BS) concept (see the ESI† discussion on spin coupling and Table S2) are consistent with earlier spectroscopic indications on WT and mutant ferredoxins:27,28 these are sites Fe1 and Fe2 that form the redox-active pair, coupled antiferromagnetically to Fe3 which remains a ‘unique’ high-spin ferric in [3Fe-4S]²⁰¹⁵⁰ of Pf Fd D14C (see Fig. 1 and S2† for the Fe/S site labeling).

Previous NRVS data on Pf Fd D14C [3Fe-4S] samples suffered from inhomogeneity and were provided only in weighted-sum spectra.16 ⁵⁷Fe partial vibrational density of states ([⁵⁷Fe-PVDOS]) for oxidized and reduced samples from the new NRVS measurements are presented in Fig. 2A and B, respectively. The spectra are distantly similar to those from the Pf Fd D14C [4Fe-4S] variant (Fig. 2C and D),19 the nitrogenase Fe protein [4Fe-4S] cluster from Azotobacter vinelandii (Av2),18 and the [Fe₄S₄][SPh]₄ model compound.21 In what follows below, the DFT-computed spectra (Fig. 2A–L, S3 and S4†) aid in rationalization of the observed NRVS data.

In general, features range from ‘acoustic’ and torsional modes below 100 cm⁻¹ to ‘breathing’ (Fe–Fe) and S–Fe–S bending modes around 100–200 cm⁻¹, and to structurally-sensitive Fe–S stretching bands in the 200–420 cm⁻¹ region. Stretching vibrational modes of the inorganic bridging sulfides (Fe–S₈) dominate the bands in the 200–300 cm⁻¹ region, and
stretching modes of the terminal cysteinate sulfurs (Fe–S,) are mostly in the 300–420 cm$^{-1}$ higher end of the spectra, as indicated by kinetic energy distribution (KED) analysis from DFT in Fig. 2I–L (see ESI† for selected [3Fe–4S]$^{0/1+}$ normal modes animated).$^{18,19,29}$

In the mixed Fe–S$_{bn}$ stretching region, the strongest NRVS features of the [3Fe–4S]$^{1+}$ Pf Fd D14C species are broad peaks at 362 and 390 cm$^{-1}$ (Fig. 2A). RR bands from 250 to 300 cm$^{-1}$ of WT [3Fe–4S]$^+$ Fd$^{13}$ overlap well with the NRVS peaks. The greatest contrasts between the NRVS and RR spectra are at 347 cm$^{-1}$ with only a small NRVS signal and at 297 cm$^{-1}$ with no detectable RR intensity, which can be explained by a lack of Fe motion at 347 cm$^{-1}$, and less change of polarizability at 297 cm$^{-1}$. Upon reduction of Pf Fd with dithionite, a modified band pattern is observed for the spectral region between 240 and 420 cm$^{-1}$ (Fig. 2B), which is similar to the reduced [4Fe–4S] Pf Fd D14C (Fig. 2D) and Av2 proteins.$^{18,19}$ For reduced [3Fe–4S] clusters, RR spectra have not been reported. The pronounced high-frequency maximum for the oxidized [3Fe–4S] D14C at 390 cm$^{-1}$ downshifts to 378 cm$^{-1}$ upon reduction, and the band at 362 cm$^{-1}$ decreases in intensity; additionally, bands at 340 and 319 cm$^{-1}$ appear.

NRVS of [3Fe–4S] revealed a drop in the intensities of the ‘breathing’ and S–Fe–S bending bands in the 140–180 cm$^{-1}$ region, otherwise very intensive in the [4Fe–4S] form (Fig. 2). Another remarkable feature of the [3Fe–4S] protein spectra is the split-band pattern of the stretching Fe–S$_{b}$ mode region (267, 286 and 297 cm$^{-1}$ for [3Fe–4S]$^{1+}$) in comparison to the well-defined peaks at 269/281 cm$^{-1}$ for the [4Fe–4S]$^{1+}$ counterpart. This type of 57Fe-PVDOS band splitting has been also demonstrated for multi-cluster Desulfovibrio vulgaris Miyazaki F (Dv ME) [NiFe] hydrogenase with a [3Fe–4S] cluster in its electron relay.$^{16}$ The dispersal of features in the [3Fe–4S] spectra can be explained by the loss of symmetry as an outcome of metal extraction from one of the [4Fe–4S] corners. The [4Fe–4S] core structure can be approximated as a $T_{d}$ point group or $D_{2h}$, while the [3Fe–4S] structure is reminiscent of a lower symmetry group such as $C_{nv}$, which would produce the observed breakdown in the pseudo-degeneracy of the bending and breathing modes.$^{18,19}$ High-resolution crystal structures of some [3Fe–4S]-containing ferredoxin indeed reveal distortions due to the absence of the fourth iron in contrast to [4Fe–4S] clusters, such as the more open $\mu_{5}$-Fe–$\mu_{5}$S ($\mu_{5}$S = S$_{0/1/2/3}$, Fig. 1) conformation.$^{30,31}$ Consistently, our computational results predict that the mean optimized Fe–$\mu_{5}$S ($\mu_{5}$S = S$_{0}$A, D) vs. Fe–$\mu_{5}$S bonds are longer by 0.07 Å (Table S3†).

Breakdown of the 57Fe-PVDOS spectra from DFT into contributions from individual Fe sites offers an alternative rationalization of the [3Fe–4S] vs. [4Fe–4S] spectral differences, see Fig. 2E–H. The major collective bands of the inorganic core calculated for [4Fe–4S]$^{1+}/2+$ at 145/150 (Fe–Fe modes) and 246/262 (Fe–S$_{b}$ modes) cm$^{-1}$, respectively, result from consolidated displacements of the four Fe sites at these vibrational energies, while in the case of [3Fe–4S]$^{0/1+}$ the corresponding individual bands of the three metal sites are distributed and therefore lack their cooperative effect. Notably, in both cluster types the degree of Fe vibrational cooperation is lower among the Fe–S modes above 300 cm$^{-1}$, which represent interactions outside the inorganic core.

Consistently at both the oxidized [3Fe–4S]$^{1+}/[4Fe–4S]^{2+}$ and reduced [3Fe–4S]$^{0}/[4Fe–4S]^{1+}$ levels, prominent 57Fe-PVDOS bands of the [3Fe–4S] cluster display a blue shift within ~20 cm$^{-1}$ when compared to the counterpart bands of the [4Fe–4S] form of Pf Fd D14C (Fig. 2A vs. C and B vs. D). A similar effect has been observed for the [3Fe–4S] vs. [4Fe–4S] derivatives of the WT Fd in a previous RR study.$^{15}$ Comparable in their magnitude, band shifts also apply to the [3Fe–4S]$^{1+}/2+$ and [4Fe–4S]$^{1+}/2+$ redox couples of the same cluster type (Fig. 2A vs. B and C vs. D). The latter observation is in line with a systematic 10–30 cm$^{-1}$ stepwise decrease of Fe–S stretching frequencies upon each one-electron reduction to the [4Fe–4S] cluster from Av2 Fe protein.$^{18}$ The mean oxidation level per Fe site, which averages to +3/2.6/2.5/2.5 respectively for the [4Fe–4S]$^{1+}/[3Fe–4S]^{2+/1+}$, [4Fe–4S]$^{1+}/[4Fe–4S]^{2+/1+}$ species, is a qualitative parameter which rationalizes the trends displayed in Fig. 2: upon a gradual decrease of the mean Fe charge, the red shift increases in both the experimental and calculated vibrational spectra. This tendency is mapped by the reduction-dependent expansion of both clusters,$^{19,23,32}$ as reflected by elongation of the mean DFT-optimized Fe–Fe/S$_{b}$/S$_{c}$ internuclear distances in the [3Fe–4S]$^{1+}$ and [4Fe–4S]$^{2+/1+}$ pairs (Table S3†). The mean Fe–Fe distance of 2.61/2.68/2.68/2.70 Å respectively for the [3Fe–4S]$^{1+}/[4Fe–4S]^{2+/1+}$ species is found to be the most relevant structural parameter which follows the above-mentioned general trend in all four Pf Fd D14C 57Fe-PVDOS spectra.

The redox-dependent structural shift of the [3Fe–4S] cluster is markedly superior to that of the [4Fe–4S] cluster (by ~0.07/ 0.10/0.08 Å for [3Fe–4S] vs. +0.02/0.05/0.01 Å for [4Fe–4S] in terms of the mean Fe–Fe/S$_{b}$/S$_{c}$ distances, respectively, see Table S3†), which can be readily accepted in view of less efficient electron delocalization in the smaller cluster. This structural shift has a tight connection with reorganization energy $\lambda$ required to relax the structure and its environment following the redox event. In semi-classical Marcus theory of electron transfer, the activation barrier for electron transfer $\Delta G^\ddagger$ is predicted to depend on $\lambda$ and the driving force $\Delta G^0$ (difference in reduction potentials) by the relationship $\Delta G^\ddagger = (\lambda + \Delta G^0)^{2}/4\lambda$. Our computed inner-sphere contribution to reorganization energy $\lambda$ (see the ESI† discussion on reorganization energy, Table S2† and earlier theoretical studies$^{19,25,35–38}$) of the [3Fe–4S]$^{1+}$ redox couple is 111.3 kJ mol$^{-1}$, which is about 6 times the 18.8 kJ mol$^{-1}$ value$^{19}$ obtained for the [4Fe–4S]$^{1+}/2+$ variant within the same Pf Fd D14C protein framework. The significantly higher reorganization energy of [3Fe–4S] explains its less frequent occurrence in electron-transfer proteins as compared to [4Fe–4S]. Interestingly, Asp (D14) coordination to [4Fe–4S] in the native Pf Fd WT protein has been indicated to increase the [4Fe–4S]$^{1+}/2+$ reorganization energy by 80 kJ mol$^{-1}$, as compared to that of conventional Cys-only ligated [4Fe–4S] in the Pf Fd D14C mutant.$^{39}$ In the native protein, a unique capability of aspartate to switch
between mono- and bidentate coordination modes to [4Fe–4S] ("carboxylate shift")\(^2\) was proposed to allow for conformational control over the electron transfer, at the expense of its reduced rate.\(^3\)

In context of the discussion above (and also in the absence of high-resolution XRD structures of the [3Fe–4S] cluster hosted specifically by Pf Fd D14C), it is remarkable to note that 1.4 Å resolution determinations of the [3Fe–4S]\(^{0/1+}\) structures from Azotobacter vinelandii Fd (Av FdI) displayed deviations within experimental error only (see the ESI\(^\dagger\) discussion on structural aspects and Table S3\(^\dagger\)).\(^{30,31}\) The Av FdI protein has been suggested to enforce an entatic state\(^10\) upon the cluster in order to facilitate electron transfer. Our DFT calculations on the reduced [3Fe–4S]\(^0\) state using the equilibrium [3Fe–4S]\(^{1+}\) structure show that stabilization of the entatic state should be backed by energetic input of as much as 14 kcal mol\(^{-1}\) from the protein (Table S2\(^\dagger\)). Further, \(^{57}\)Fe-PVDOS spectra computed for the entatic state produce high-intensity extensions in the 400–460 cm\(^{-1}\) region (Fig. S3\(^\dagger\)), which are not consistent with the NRVS experiment. Collectively, this indicates that the entatic state of the [3Fe–4S] cluster core is not realized in Pf Fd D14C.

Conclusions

In conclusion, our results highlight the power of the combined application of NRVS and DFT for understanding Fe–S clusters.\(^{18,19,41}\) The gradual red shift of vibrational bands, both upon cluster reduction and the [3Fe–4S]–to-[4Fe–4S] variant transition in the very same Pf Fd D14C protein, is explained by the steadily increasing ferrous character of the inorganic core; this trend is well mapped by expansion of the Fe–Fe internuclear distances. The reduction-dependent expansion of the [3Fe–4S] cluster is markedly superior to that of the [4Fe–4S] cluster, which ultimately translates to higher reorganization energy of [3Fe–4S] and rationalizes its less frequent occurrence in electron-transfer proteins. The dispersal and splitting of the NRVS bands caused by conversion from [4Fe–4S] to [3Fe–4S] are related to the electronic and structural aspects of both cluster types, in particular to the symmetry loss in the smaller cluster with one open corner. A new approach in the breakdown of the collective DFT-computed NRVS spectra into contributions from individual \(^{57}\)Fe nuclei shows that the band dispersal results from the inhomogeneity of the metal site properties in [3Fe–4S], which is an alternative indication for a less efficient electron delocalization, as compared to [4Fe–4S]. To the best of our knowledge, this is the first comprehensive NRVS analysis of a [3Fe–4S] cluster in a protein and a pioneering vibrational study of its reduced [3Fe–4S]\(^0\) form, which therefore completes the list of typical Fe–S clusters. An extension of the present approach is expected to shed light on the structural and functional properties of electron relays and active sites of enzymes with multiple Fe–S cofactors. Applicable examples include membrane-bound \(\text{O}_2\)-tolerant [NiFe] hydrogenase with its unique [4Fe–3S] proximal cluster,\(^10,42\) and CO dehydrogenase/acetyl-CoA synthase hosting a heterometallic NiFe–S cluster.\(^43\)

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Notes and references


