

Reaction of O₂ with a di-iron protein generates a mixed valent Fe²⁺/Fe³⁺ center and peroxide

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Abstract

The gene encoding the cyanobacterial ferritin *SynFtn* is upregulated in response to copper stress. Here we show that, while *SynFtn* does not interact directly with copper, it is highly unusual in several ways. Firstly, its catalytic di-iron ferroxidase center is unlike those of all other characterised prokaryotic ferritins and instead resembles an animal H-chain ferritin center. Secondly, as demonstrated by kinetic, spectroscopic and high resolution X-ray crystallographic data, reaction of O_2 with the di- Fe^{2+} center results in a direct, one electron oxidation to a mixed valent Fe^{2+}/Fe^{3+} form. Iron- O_2 chemistry of this type is currently unknown amongst the growing family of proteins that bind a diiron site within a 4 α -helical bundle in general and ferritins in particular. The mixed valent form, which slowly oxidized to the more usual di- Fe^{3+} form, is an intermediate that is continually generated during mineralization. Peroxide, rather than superoxide, is shown to be the product of O_2 reduction, implying that ferroxidase centers function in pairs via long-range electron transfer through the protein resulting in reduction of O_2 bound at only one of the centers. We show that electron transfer is mediated by the transient formation of a radical on Tyr40, which lies ~ 4 Å from the di-iron center. As well as demonstrating an expansion of the iron- O_2 chemistry known to occur in nature, these data are also highly relevant to the question of whether all ferritins mineralize iron via a common mechanism, providing unequivocal proof that they do not.

Significance

Enzymes that activate dioxygen at di-iron centers located within four α -helical bundles with carboxylate residue-rich coordination are abundant in nature. Mechanistic studies of the Fe^{2+}/Fe^{2+} forms of these proteins have shown that peroxo- Fe^{3+}/Fe^{3+} intermediates are common to most of their reactions. Remarkably, the O_2 reaction of the unusual ferritin *SynFtn* does not lead to the peroxo- Fe^{3+}/Fe^{3+} form but instead directly generates mixed valent Fe^{3+}/Fe^{2+} centers and hydrogen peroxide via a mechanism that involves long range electron transfer between di-iron centers and the transient formation of a Tyr• radical close to the di-iron center. This work reveals previously unrecognised catalytic iron- O_2 chemistry and demonstrates that ferritins do not all mineralize iron via a common mechanism.

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Ferritins constitute a ubiquitous family of proteins found in all domains of life. They are rhombic dodecahedral 24-mer protein assemblies that delimit a central cavity in which a ferric-oxo mineral can be reversibly deposited (1-3). They provide a means for storage, buffering and detoxification of iron, contributing to cellular iron management and the alleviation of oxidative stress. Animal ferritins are, in general, heteropolymers composed of two types of subunits, H- and L-chains. H-chain ferritins contain a di-iron site, termed the ferroxidase center, at which the rapid oxidation of Fe^{2+} to Fe^{3+} is coupled to O_2 reduction. L-chain ferritins lack this catalytic center but are thought to promote nucleation of the mineral core via a carboxylate-rich patch located on the inner surface of the protein nano-cage (4, 5). The subunit composition of animal ferritins is dependent on the tissue from which the protein was isolated (2).

The genomes of plants and prokaryotes often encode more than one predicted ferritin, of which all differ from their eukaryotic counterparts in that they are composed solely of H-chain like subunits containing iron binding catalytic centers. Other than in the distinct subset of heme binding bacterioferritins, the amino acid sidechains ligating these two iron atoms are highly conserved. The major structural difference between the catalytic centers of prokaryotic, animal and plant (phyto) ferritins is the presence of a third iron binding site, termed site C, approximately 6 Å from the di-iron site of the prokaryotic ferritin (Ftn) (6). Both phytoferritin and animal H-chain subunits lack this third iron binding site, and phytoferritin contains the carboxylate-rich patch associated with the animal L-chains, which is absent from both animal H-chains and Ftn (3). Therefore, the heme-free ferritins can be classified as animal-, prokaryotic- or plant-like by considering the subunit composition together with the presence or absence of a site C or a carboxylate-rich patch. However, despite differences in structure, composition and mechanistic detail, all of these proteins share a common first step in their reaction cycle; namely, the formation of an unstable diferric-peroxo (DFP) intermediate that decays to yield a ferric-oxo mineral precursor (7).

Cyanobacteria are photosynthetic organisms that play a critical role in the carbon cycle. Marine picocyanobacteria, including *Synechococcus* and *Prochlorococcus*, alone are estimated to account for a minimum of 20% of all global carbon fixation (8). Coastal dwelling *Synechococcus* strains exhibit greater tolerance to metal stress than their ocean dwelling counterparts. This tolerance is due to the presence of a greater number of genes involved in metal ion homeostasis, most likely as a consequence of the evolutionary pressures exerted by seasonal variations in the metal ion content of their habitat (9). An unusual example is the response of the coastal *Synechococcus* strain CC9311 to copper- and associated oxidative-stress, which involves the up-regulation of the predicted ferritin sync_1539, hereto referred to as *SynFtn* (9). Upregulation of a predicted ferritin was unexpected because they are typically expressed in response to iron (10). Furthermore, *SynFtn* was detected in the exoproteome of *Synechococcus* CC9311 (11), raising the possibility that this ferritin is localized, at least in part, outside of the cytoplasm, again suggestive of a role other than regulation of iron concentration within the cell.

Here we report optical and magnetic spectroscopic, kinetic and high-resolution structural studies of *SynFtn*, revealing that the O_2 -driven oxidation of the $\text{Fe}^{2+}/\text{Fe}^{2+}$ form does not result in a (per)oxo- $\text{Fe}^{3+}/\text{Fe}^{3+}$ intermediate as it does in all other characterised ferritins, and most other carboxylate bridged di-iron centers. Instead, *SynFtn* directly generates a mixed valent $\text{Fe}^{3+}/\text{Fe}^{2+}$ form along with hydrogen peroxide. Furthermore, the data demonstrate the formation of a

transient radical on Tyr40, which is essential for rapid Fe²⁺ oxidation. These data lead to a mechanistic model in which diiron centers cooperate via long range electron transfer to reduce a single O₂ molecule bound at one with formation of two mixed valent diiron centers. The physiological significance of these observations are discussed.

Results

SynFtn is a functional prokaryotic ferritin with a eukaryotic-like active site. Purified *SynFtn* behaved as a typical 24mer ferritin when examined by size exclusion chromatography and native PAGE (SI Appendix, Fig. S1A). However, alignment of its amino acid sequence with those of other ferritins showed that the residues comprising site C in all prokaryotic ferritins characterized to date are absent (SI Appendix, Fig. S1B). The 2.05 Å crystal structure of *SynFtn* (SI Appendix, Fig. S1C and D, Table S1) showed that each of the 24 subunits adopts the characteristic 4 α-helical bundle motif of the ferritin family and their assembly into a 24meric protein cage with the expected 4-3-2 symmetry. The protein as crystalized was free of metal ions but the disposition of the conserved ferroxidase center ligands suggested that it would be capable of binding two metal ions without major structural rearrangement (Fig. 1A).

The interactions of the protein with iron were assessed in solution. Absorbance monitored iron mineralization assays demonstrated that *SynFtn* was capable of laying down a mineral core containing up to 2000 Fe³⁺ per protein cage (Fig. 2A) at a rate greater than that of *E. coli* FtnA (12) or *Pyrococcus furiosus* Ftn (13), the most intensively studied of the Ftns, and similar to that of homopolymers of animal H-chains (4). The rate increased with pH consistent with the release of protons during mineralization, as reported for other ferritins (1). Interactions of *SynFtn* with copper were also investigated because expression of the gene encoding it was upregulated in response to this metal. 400 copper ions (as 200 μM Cu²⁺ or Cu⁺) were added per *SynFtn* and incubated for 60 min aerobically (Cu²⁺, Cu⁺) or anaerobically (Cu⁺) before removal of unbound copper via a desalting column. Residual copper concentrations determined were 2 ± 1 μM (anaerobic Cu⁺), 3 ± 1 μM (aerobic Cu⁺) and 5 ± 2 μM (aerobic Cu²⁺), while that determined for a protein free Cu²⁺ control was 3.0 ± 1.0 μM. Incorporation of copper into the mineral core during deposition was probed by the same assay method, but using 4.1 μM *SynFtn* incubated aerobically with 1.6 mM Fe²⁺ and 200 μM of either Cu⁺ or Cu²⁺. Copper concentrations determined were 5 ± 2 μM (Cu²⁺) and 6 ± 3 μM (Cu⁺) compared to 3 ± 1 μM in protein free controls, and we therefore conclude that copper is not incorporated into the ferric mineral core of *SynFtn*. Thus, in contrast to iron, there is no direct interaction between copper, either as Cu²⁺ or as Cu⁺ under aerobic or anaerobic conditions, and *SynFtn*. These data were unexpected because Cu²⁺ binding has previously been detected for ferritins not implicated in copper stress response (14, 15). Therefore, it seems that the function of *SynFtn* under copper stress is not associated with the direct sequestration of copper.

Crystals of *SynFtn* that were soaked under aerobic conditions in Fe²⁺ for either 2 min (Fig. 1B) or 20 min (Fig. 1D), revealed two additional areas of electron density at the ferroxidase center and a third in the 3-fold channel that penetrates the protein coat, all modeled as iron. Rearrangement of the side chain of Glu33, a ligand to one of the metal ions, was apparent in both iron soaked structures, as was bridging electron density between the metal centers. There was no evidence of electron density from a third metal bound at the ferroxidase center, and we conclude that site C is absent from *SynFtn*, as predicted from the sequence. As such, it is unique

among characterized prokaryote ferritins, and in this sense the *SynFtn* ferroxidase center bears far greater resemblance to animal H-chain proteins (Fig. 1C). Thus, like other prokaryotic ferritins, *SynFtn* is composed solely of H-chain like subunits lacking a carboxylate-rich patch, but the catalytic center lacks the ligands for site C, the defining feature of other prokaryotic ferritins, defying categorization according to the scheme described above.

Aerobic incubation with Fe^{2+} resulted in a color change, expected to be due to a di-ferric form of the ferroxidase center as it is in other ferritins. However the Fe-Fe distance in the 2 min soak structure (3.90 Å, Fig. 1B and SI Appendix, Table S1-3) was far longer than in di-ferric ferroxidase center structures (15). Increasing the soaking time to 20 min resulted in a contraction of the Fe-Fe distance to 3.46 Å (Fig. 1D and SI Appendix, Table S1-3). Concomitant with the contraction in Fe-Fe distance was an increase in the angle formed by these two iron ions and the bridging density, from 113.0 to 116.7 degrees. Both of these changes are consistent with those observed for oxidation of a hydroxy bridged mixed valent $\text{Fe}^{2+}/\text{Fe}^{3+}$ model complex to an oxo bridged di- Fe^{3+} state (16), suggesting that the unexpectedly long Fe-Fe distance in the 2 min soak results from the formation of a hydroxy bridged mixed valent ferroxidase center (MVFC) that is subsequently oxidized to the oxo bridged di- Fe^{3+} form upon prolonged exposure to O_2 .

Formation of a mixed valent ($\text{Fe}^{2+}/\text{Fe}^{3+}$) di-iron center in *SynFtn*. The kinetics of the initial iron- O_2 reaction at the catalytic sites following addition of Fe^{2+} to apo *SynFtn* were monitored using stopped-flow absorbance at a single wavelength of 340 nm (Fig. 2B, C). The observed dependence on the stoichiometry of Fe^{2+} to protein was complex (see *Materials and Methods* and SI Appendix, Fig. S2A-C). Rapid oxidation of Fe^{2+} ($t_{1/2} < 200$ ms) was observed only at loadings ≥ 18 $\text{Fe}^{2+}/\text{SynFtn}$. *SynFtn* variants E33A and E110A, in which iron binding residues from each ferroxidase center site were individually substituted, were virtually inactive (SI Appendix, Fig. S2D). Together, and in the absence of cooperative binding of Fe^{2+} at ferroxidase centers, these observations suggest that optimum catalytic activity requires binding of iron to both ferroxidase center sites, as in other ferritins. The amplitude of the absorbance change associated with rapid oxidation increased linearly with iron loading up to 48 $\text{Fe}^{2+}/\text{SynFtn}$, sufficient to saturate all available ferroxidase center sites. Whilst the rate of rapid oxidation continued to increase at greater iron loadings the amplitude of the absorbance change associated with this phase saturated at approximately half that expected from equivalent measurements of other ferritins (SI Appendix, Fig. S2A) (17-19). A slow phase of Fe^{2+} oxidation was also apparent, and, at iron loadings sufficient to saturate the ferroxidase center sites, the sum of the amplitudes of the absorbance increases associated with the rapid and slow phases accounted for the oxidation of all added Fe^{2+} . At low iron loadings (< 18 $\text{Fe}^{2+}/\text{Ftn}$) oxidation was dominated by a kinetically distinct phase characterized by an apparent rate constant intermediate between those of the rapid and slow processes (SI Appendix, Fig. S2A, C). The amplitude of this phase decreased with increasing iron loading such that it was only a minor component when the iron loading was ≥ 18 $\text{Fe}^{2+}/\text{Ftn}$, and not observed at all at ≥ 48 $\text{Fe}^{2+}/\text{Ftn}$. Importantly, spectro-kinetic measurements performed under similar conditions showed no evidence of formation of a DFP intermediate during rapid iron oxidation (SI Appendix, Fig. S2E) suggesting a mechanism of iron oxidation distinct from all previously reported ferritins.

Electron paramagnetic resonance (EPR) measurements of samples frozen 10 s after aerobic addition of Fe^{2+} to *SynFtn* resulted in detection of a paramagnetic species with g-values

all below 2.0 (Fig. 3A). The signal was remarkably similar to those observed previously upon partial reduction of the di-Fe³⁺ sites of *Pyrococcus furiosus* Ftn (20), human H-chain ferritin (21) and of model di-iron complexes (22). It was therefore assigned as arising from the formation of a Fe²⁺/Fe³⁺ MVFC. Importantly, none of the ferritins mentioned above give rise to a MVFC directly upon reaction with O₂; they all form a Fe³⁺/Fe³⁺ species that can be subsequently partially reduced to the MVFC. Direct MVFC formation in *SynFtn* was observed even for iron loadings (12 Fe²⁺/Ftn) below that at which rapid oxidation was detected by stopped flow absorbance. However, the intensity increased markedly at loadings ≥ 24 Fe²⁺/*SynFtn* (1 Fe²⁺ per ferroxidase center) (Fig. 3B), consistent with the requirement for simultaneous occupancy of both iron binding sites.

The kinetics of MVFC formation were followed by rapid-freeze quench EPR. The intensity maximized approximately 1 s after addition of saturating Fe²⁺, at which point it was present on approximately 70% of protein monomers (i.e. ~17 MVFCs per 24mer), before decaying away over the next 10 s (Fig. 3C, D). Simulation of the MVFC envelope (Fig. 3A and Table 1) could be achieved only by assuming two overlapping components with g-values at 1.943, 1.800 & 1.760 (major species, 80% of signal) and 1.955, 1.870 & 1.790 (minor species, 20% of signal), reminiscent of the result of one electron reduction of the di-ferric iron-oxo center of *P. furiosus* Ftn (20). The major species accounted for 56% of ferroxidase centers at peak intensity and the minor species a further 14%. Samples for rapid freeze quench EPR could not be prepared under identical conditions to stopped flow absorbance measurements (see *Materials and Methods*). However, the time dependence of the intensity of the MVFC signal was consistent with formation of this species during the rapid phase of iron oxidation detected using stopped flow, and subsequent decay during the slow phase (Fig. 3D).

The formation of a MVFC intermediate on the pathway to mineral core formation explains not only the unusually large Fe-Fe distance in crystals soaked aerobically in Fe²⁺ for 2 min, but also the observation that only 50% of the Fe²⁺ bound at ferroxidase centers is oxidized in the initial rapid phase of activity.

The MVFC forms and decays throughout mineralization in *SynFtn*. Incubation of *SynFtn* with larger excess of Fe²⁺ over ferroxidase center sites led to greater longevity of the MVFC signal (SI Appendix, Fig. S3B), indicating that the intermediate is continually generated during turnover and that the ferroxidase center continues to function as the principal catalytic center following the initial reaction of Fe²⁺ and O₂. To investigate whether ferroxidase center activity can be regenerated, *SynFtn* was incubated with 400 Fe²⁺/Ftn for four minutes (sufficient time for complete oxidation of all Fe²⁺) and immediately exposed to a further 72 Fe²⁺/Ftn. This sample exhibited ~30% recovery of the rapid oxidation phase measured by stopped-flow (with a rate constant indistinguishable from that for the apo-protein, SI Appendix, Fig. S3C, D). The extent of this recovery increased when the incubation time between the initial and subsequent Fe²⁺ additions was increased, attaining 80% at one hour, at which point the extent of recovery became invariant with increased incubation. Therefore, as with many other ferritins, the di-Fe³⁺ form of the ferroxidase center is unstable leading to translocation of Fe³⁺ into the central cavity and regeneration of the initial, rapid phase of Fe²⁺ oxidation.

O₂ reduction by *SynFtn*. Reduction of O₂ by a single electron, derived from formation of the MVFC, would yield the superoxide radical anion. The EPR spectrum of *SynFtn* frozen at any point

during turnover contained no features attributable to superoxide, nor was its presence detectable using the spin trap 5,5-Dimethyl-1-Pyrroline-N-Oxide (DMPO) (SI Appendix, Fig. S4A). Superoxide may be transiently formed during the initial reaction of iron and O₂ but the detection of near stoichiometric concentrations of MVFC in the EPR spectra demonstrates that it cannot remain bound here prior to the second, slower oxidation to the di-Fe³⁺ form. Together these observations suggest that any superoxide produced upon MVFC formation is consumed prior to the oxidation of the catalytic site to the Fe³⁺/Fe³⁺ form, indicating that the mechanism of MVFC formation is more complex than a simple one electron reduction of O₂.

The stoichiometry of the iron/O₂ reaction, measured using an O₂ electrode, was found to be ~2 Fe²⁺ per O₂ for the initial phase of iron oxidation (Fig. 4A), indicating that hydrogen peroxide is the final product of O₂ reduction. Indeed, peroxide ion was readily detected using the Amplex Red assay (Thermo Fisher), at close to the expected 2:1 ratio of Fe²⁺ to H₂O₂ (Table 2). An Fe²⁺:O₂ ratio of 2:1 is reported for eukaryotic H-chain homopolymers, but this ratio gradually increases towards 4:1 during mineralization due to peroxide-mediated oxidation of Fe²⁺ at the growing mineral surface (23). No evidence for peroxide involvement in *SynFtn* mediated Fe²⁺ oxidation was found: the di-Fe²⁺ form of the ferroxidase center did not react with exogenously added peroxide, and the Fe²⁺:O₂ ratio remained at ~2:1 even during mineralization (Fig. 4B).

MVFC formation involves cooperation between pairs of ferroxidase centers. The lack of evidence for superoxide formation, the ~2:1 Fe²⁺:O₂ ratio and stoichiometric formation of hydrogen peroxide raise the key question of how these result from the oxidation of a single Fe²⁺ at each ferroxidase center of *SynFtn*. The simplest explanation is that the ferroxidase centers function in pairs, each delivering a single electron to an O₂ molecule bound at only one of them. This proposal requires that O₂ reacts directly with a maximum of 50% of ferroxidase centers during each catalytic cycle, the remainder of the observed MVFCs being formed by electron transfer away from di-Fe²⁺ sites with no O₂ bound.

In addition to the MVFC, mononuclear high spin Fe³⁺ was also detected by EPR, at *g* = 4.3 (SI Appendix, Fig. S4B), upon reaction of *SynFtn* with Fe²⁺ and O₂. Both species formed with similar rapid kinetics (Fig. 3D), and so the Fe³⁺ species is not a breakdown product of the fully oxidized ferroxidase center. As both E33A and E110A variants were inactive, it is apparent that, as for other ferritins, both ferroxidase center iron binding sites are required for catalytic activity (7, 24). Thus, it is highly unlikely that the mononuclear Fe³⁺ signal results from the reaction of Fe²⁺ and O₂ at singly occupied ferroxidase centers. Rather we propose that electron transfer from ferroxidase centers singly occupied by Fe²⁺, most likely at the predicted higher affinity site (site A in Fig. 1A), results in mononuclear high spin Fe³⁺ formation. Quantification by comparison with a high spin Fe³⁺-bound protein standard indicated that, at maximum intensity, high spin Fe³⁺ was present on ~8% of *SynFtn* monomers (2 high spin Fe³⁺ per 24mer). Therefore, overall EPR spin quantitation demonstrated that, during the initial, rapid reaction with O₂, a maximum of ~80% of ferroxidase centers are in a paramagnetic state, before quenching to antiferromagnetically coupled EPR-silent species during a subsequent slower phase.

A model in which pairs of ferroxidase centers cooperate for rapid formation of MVFC and hydrogen peroxide not only accounts for the simultaneous formation of MVFC and mononuclear Fe³⁺ (Fig. 3D), but is also consistent with the kinetic dependence of Fe²⁺ oxidation on the ratio of Fe²⁺:*SynFtn*. At low iron (e.g. 12 Fe²⁺/24mer), the MVFC was still formed (Fig. 3B), but the rapid

phase of Fe²⁺ oxidation was not observed because there was insufficient occupation of closely located pairs of ferroxidase centers for rapid electron transfer to occur.

The above model requires the presence of an electron transfer pathway between ferroxidase centers. Since it was first postulated for a diiron, O₂-activating protein (25) the transient oxidation of aromatic amino acid sidechains has been accepted as a means of such electron transport across large distances in proteins without need for redox active cofactors (26). In addition to the MVFC and Fe³⁺ signals, EPR spectra revealed the presence of a signal that is characteristic of a tyrosyl radical (Fig. 5A) with an intensity that increased linearly with added Fe²⁺ at loadings up to 80/Ftn and maximized within the first second of reaction (Fig. 5B, C). Quantification of this signal revealed that it is present on ~5% of monomers. Consequently, to assess the functional relevance, we sought to determine which of the six tyrosine residues in the *SynFtn* monomer hosts the radical.

Tyr40 is essential for rapid MVFC formation. An inter-ferroxidase center electron transfer pathway would most likely involve Tyr40, a residue that is approximately 4 Å away from the Fe_B site, is conserved among all ferritins containing a catalytic di-iron center (SI Appendix, Fig. S1B) and is reported to be the site of radical formation in several (12, 27, 28). The EPR spectrum of the radical species formed by *SynFtn* was simulated (Fig. 5A) using the Tyrosyl Radical Spectra Simulation Algorithm, TRSSA (29). The set of hyperfine splitting parameters used in the simulation suggested a specific tyrosyl sidechain conformation that was in good agreement with that of Tyr40 in the structures of *SynFtn* reported here (SI Appendix, Table S4, S5). Consequently, the structure, activity and spectroscopic properties of a Y40F variant of *SynFtn*, in which this residue is replaced by the non-oxidizable analogue phenylalanine, were investigated.

The crystal structure of this variant revealed no significant changes beyond the substituted side chain (SI Appendix, Fig. S5A, B). Iron soak experiments revealed only one iron bound at the ferroxidase center, at the high affinity site A. Rapid oxidation of Fe²⁺ was completely abolished in Y40F *SynFtn* (Fig. 6A –C, Fig S5C, D). However, binding of iron to both ferroxidase center sites was confirmed by detection of the characteristic MVFC EPR signal (Fig. 6D) although the MVFC formed at a rate much slower than in the wild type protein (Fig. 6E, F), consistent with the kinetics of Fe²⁺ oxidation deduced from the stopped flow data. As with the wild type protein, two species were required to simulate the observed signal (Fig. 6D, Table 1). Quantification revealed a maximal intensity corresponding to 60% of the subunit concentration (i.e. 14 MVFC per 24mer) 3 seconds after addition of Fe²⁺. Furthermore, EPR spectra of Y40F *SynFtn* showed no evidence of tyrosyl radical formation at any point during O₂ driven Fe²⁺ oxidation. However, absorbance measurements over a longer time period following addition of 400 Fe²⁺ per protein showed that mineralization proceeded at a rate similar to that of the wild type protein (SI Appendix, Fig. S2D). For protein concentrations of 0.5 μM, all Fe²⁺ oxidation in Y40F *SynFtn* proceeded at the rate observed for the slow oxidation of the MVFC to the di-ferric form for the wild type protein, suggesting that this process is the rate limiting step for core formation under these conditions. This rate is unaffected by substitution of Tyr40, the site of radical formation essential for *rapid* oxidation of Fe²⁺, and so we conclude that initial reaction of O₂ with di-Fe²⁺ *SynFtn* ferroxidase centers produces a species capable of oxidizing a Tyr residue, but the subsequent, slower oxidation of the MVFC does not.

The observed Fe:O₂ reaction stoichiometry for Y40F *SynFtn* was slightly greater than that for the wild type protein (SI Appendix, Fig. S5E, Table 2). This increase is most likely due to impaired ferroxidase center function; because Fe²⁺ persists in solution for longer it is susceptible to oxidation by peroxide released during the reaction cycle, a hypothesis supported by the detection of hydroxyl radical in spin trapping experiments (SI Appendix, Fig. S4A) and low levels of peroxy radical (SI Appendix, Fig. S4B) in EPR measurements at cryogenic temperatures conducted with this variant.

Discussion

The prokaryotic ferritin *SynFtn* contains a catalytic ferroxidase center that more closely resembles those of eukaryotic H-chain ferritins than ferritins isolated from other prokaryotes. An H-chain type mineralization mechanism with direct formation of a DFP intermediate was therefore anticipated, but instead *SynFtn* produced a mixed valent Fe²⁺/Fe³⁺ species upon reaction with Fe²⁺ and O₂. Mixed valent di-iron centers are known, for example in hemerythrins (30), purple acid phosphatases (31), rubrerythrins (32), ribonucleotide reductase (33), soluble methane monooxygenase (34), and the HD-domain protein superfamily members myo-inositol oxygenase (MIOX) (35) and PhnZ (36). However in all cases other than MIOX these mixed valent centers cannot be formed by reaction of the di-Fe²⁺ site with O₂ and are generally accessed by reduction of the di-Fe³⁺ state.

Many carboxylate bridged diiron proteins are characterized by a four α -helical bundle motif that coordinates the diiron center via a carboxylate-rich ligand set. This group includes ribonucleotide reductase, methane monooxygenase, plant fatty acyl-ACP desaturase, toluene/o-xylene monooxygenase, toluene 4-monooxygenase, and members of the ferritin family (37-39) and all activate O₂ to form a di-Fe³⁺-peroxo intermediate. MIOX is distinct from this group in that its diiron site is housed in an HD-domain, where five helices contribute a histidine-rich coordination environment. The catalytically active form of MIOX is the mixed valent Fe²⁺/Fe³⁺ state, which is formed in an activating step prior to further reaction with O₂ generating a highly oxidizing di-Fe³⁺-superoxo complex that enables the enzyme to cleave a C-H bond. The di-Fe²⁺ state of MIOX can react with O₂ to form the Fe²⁺/Fe³⁺ state but only either under limiting O₂ or in the presence of an exogenous reducing agent (40). Under other conditions the oxidation to a di-Fe³⁺ state common to all other carboxylate bridged diiron oxygenases occurs. Therefore, the formation of a MVFC in *SynFtn* is surprising given that it occurs regardless of the level of O₂ excess and in the absence of a reductant, and that the ligand set of the Fe_A and Fe_B sites of *SynFtn* are identical to those of the iron ions in animal ferritins, in which the first detectable intermediates are DFP species. Also, ferritins have no obvious need of highly oxidizing intermediates during their reaction cycles leading to mineral formation, and further reaction of the MVFC of *SynFtn* forms only a di-Fe³⁺ state similar to that produced in a concerted step by other ferritins.

The product of oxygen reduction in the activating step of MIOX has not been determined but, given the absence of any potential electron donors other than the diiron site, presumably superoxide is formed. EPR spectra yielded no evidence of superoxide production by *SynFtn* whilst almost stoichiometric levels of peroxide were detected. The peroxide produced does not react further with the mixed valent center, nor with Fe²⁺ in the cavity, another unexpected feature given the propensity of peroxide to oxidize Fe²⁺. In further contrast to MIOX, three paramagnetic species

were detected by EPR for *SynFtn*, all forming too rapidly to be associated with the second, slower phase of iron oxidation detected by stopped-flow absorbance. The time dependence of the *SynFtn* MVFC EPR signal intensity suggests that it may form more slowly than iron is oxidized in the initial rapid phase. This discrepancy is most likely accounted for by the slightly lower temperature at which RFQ mixing experiments were conducted, but an alternative explanation is that the MVFC is formed by reduction of a di-Fe³⁺ intermediate. This possibility is unlikely because: (a) No spectroscopic evidence for a ferric-peroxo species was apparent; (b) It is unclear how a Fe³⁺/Fe³⁺ diiron site could be sufficiently oxidizing to generate a Tyr radical (such a reaction is without precedent); (c) There was no evidence for two phases in the initial part of the stopped-flow data; (d) The X-ray crystal structures of mixed valent and di-Fe³⁺ forms of the protein were obtained by incubation of apo-*SynFtn* crystals in aerobic Fe²⁺ solution for increasing time periods, reflecting the formation of the Fe²⁺/Fe³⁺ form prior to the di-Fe³⁺ form; and (e) Substitution of Tyr40 resulted in loss of rapid oxidation; the above alternative would be expected to result in unaffected oxidation to the di-Fe³⁺ form and its stabilization in the absence of the oxidizable Tyr.

Thus, the data are consistent with the direct formation of the MVFC upon reaction with O₂ followed by a slower oxidation to the di-Fe³⁺ state. Variability in the integrated intensities of EPR signals between independent RFQ samples meant that it was unclear whether the MVFC, tyrosyl radical and mononuclear ferric iron form at exactly the same rate. However, the data demonstrate unequivocally that these species are formed during the initial rapid phase of Fe²⁺ oxidation and are not formed sequentially.

Quantification of the EPR signals showed that only the MVFC forms in sufficient yield to be a reasonable candidate for the ultimate source of reducing equivalents for O₂. Oxidation of only one of two Fe²⁺ ions at each di-iron site, together with an overall Fe²⁺:O₂ ratio of 2:1 and peroxide as the product of O₂ reduction, indicates that pairs of ferroxidase centers (separated by ~24.5 Å) function together to deliver two electrons to a shared O₂ substrate. Therefore, the data point to the existence of a long-range electron transfer pathway between ferroxidase centers, and the transient formation of a radical on the strictly conserved residue Tyr40 is consistent with such a pathway; see Fig. 7. Here, the Tyr40 radical is formed simultaneously with the MVFC and almost immediately quenched by oxidation of Fe²⁺ at the second, remote ferroxidase center, generating a second MVFC or a mononuclear Fe³⁺, depending on the occupancy, the Tyr radical itself only being detected if the partner iron binding site is vacant. While the MVFC is the intermediate that drives mineralization, Tyr40 is essential for rapid oxidation, presumably because inter-center electron transfer is inhibited in the absence of an oxidizable sidechain close to the ferroxidase center. Whilst oxidation of a di-Fe²⁺ center to the Fe²⁺/Fe³⁺ state and inter subunit electron transfer mediated by oxidation of aromatic amino acids have both been previously reported, the occurrence of both of these phenomena simultaneously to facilitate reduction of a shared O₂ substrate represents novel chemistry not previously identified amongst the widespread family of carboxylate bridged diiron oxygenases.

A key question arising from this work is why this unique mechanism has evolved in this ferritin. Recalling that *SynFtn* is upregulated in response to elevated copper levels may provide an answer. Copper toxicity is known to be mediated through redox stress and by the displacement of native metals from protein sites, particularly iron-sulfur cluster proteins (41). Under such circumstances, the organism would have an urgent need to rapidly remove free iron to prevent it from engaging in Fenton chemistry. Thus, the extremely rapid Fe²⁺ oxidation observed in *SynFtn*

would serve to detoxify excess iron rapidly. Other ferritins achieve similar rates via oxidation of both Fe²⁺ ions and so the requirement to suppress Fenton chemistry alone cannot account for the mechanism reported here. Of further potential importance is that, in contrast to the DFP intermediate generated by H-chain ferritins, the MVFC of *SynFtn* contains a single unpaired electron and as such would generate closed shell molecules following single electron oxidation/reduction events with the reactive oxygen species superoxide- and hydroxyl- radical, of which both are generated under conditions of oxidative stress. Such quenching of toxic radicals may provide an additional mechanism of protection against oxidative stress.

Ferritins in general do not appear to be particularly widely distributed in cyanobacteria. Nevertheless, multiple sequence alignments of annotated ferritins from available cyanobacterial genomes (SI Appendix, Dataset S1) revealed >100 homologues of *SynFtn* (with no site C), which occur almost exclusively in the marine picocyanobacteria (*Synechococcus* and *Prochlorococcus*). Furthermore, while the more standard prokaryotic ferritin (containing a site C) is found in a genetically more diverse set of cyanobacteria, it is also much less common than *SynFtn* (only 55 homologues found in the genomes surveyed). For example, a few species of *Prochlorococcus*, such as *Prochlorococcus* sp. strain MIT9313 and MIT9303, have both standard and *SynFtn* types, but the majority of *Prochlorococcus* have only one ferritin, and it is almost always a *SynFtn* (SI Appendix, Dataset S1). *SynFtn* homologues appear to occur almost exclusively in cyanobacteria (we found only one example in other bacteria) suggesting that they fulfil a detoxification function that is somehow specialized for (marine) cyanobacteria. Furthermore, the prevalence of *SynFtn* in *Prochlorococcus*, a group known for their stringent genome streamlining (42), is indicative of the importance of this gene to fitness in picocyanobacteria.

Finally, a proposal that all ferritins mineralize iron via a common, 'universal' mechanism (43) has gained significant exposure in recent times. The data presented here show unequivocally that *SynFtn* utilizes a mechanism distinct from those of all previously characterized ferritins, demonstrating that a single mechanism that describes all ferritins does not exist. Rather, the data provide further evidence for mechanistic diversity amongst members of this super-family (6), and the important questions now focus on the reasons for it. *SynFtn* exhibits unique activity despite the ligands coordinating the di-iron catalytic center being identical to those of other ferritins (Fig. 1, SI Appendix, Fig. S1B). The substitution of several near ferroxidase center residues that are conserved in other ferritins may be the source of this difference (SI Appendix, Fig. S6) and a major challenge now is to elucidate the control of DFP versus MVFC reactivity exerted by such factors.

Materials and Methods

Protein overexpression and purification. *Sync_1539* was amplified from CC9311 genomic DNA (primers CCGAATTCATGGCTACCGACGTTGCTC and CCCCTCGAGGGTCAACCTCGCTGCGTTTGAC), resulting in a 569 bp product with *EcoRI* and *AvaI* cleavage sites on either end. Following purification and digestion with *EcoRI* and *AvaI*, PCR product was ligated into digested pET21b plasmid (Novagen), and used to transform One Shot TOP10 electrocompetent *E.coli* cells (Invitrogen) according to the manufacturer's specifications. Transformed cells were plated on LB plates containing 50 µg mL⁻¹ kanamycin, incubated overnight at 37 °C, colonies picked into LB containing 50 µg mL⁻¹ kanamycin and cultures incubated at 37 °C overnight. Plasmid was purified from cultures using a Qiagen miniprep kit (Qiagen) and

plasmid insert verified with Sanger sequencing. Plasmids for variants Y40F, E33A and E110A cloned into pET21a (Novagen) were purchased from Genscript (New Jersey, USA). The encoded proteins were overexpressed in *E. coli* strain BL21(DE3) (Promega). Cultures were grown in LB containing 100 $\mu\text{g mL}^{-1}$ ampicillin at 37 °C, 200 rpm shaking to $\text{OD}_{600\text{ nm}}$ of 0.6-0.8. Overexpression was induced by addition of isopropyl β -D-1 thiogalactopyranoside (IPTG, 20 μM) and cultures grown for a further 3 hr at 37 °C, 200 rpm shaking prior to harvesting by centrifugation. Cells were re-suspended in 20 mM HEPES pH 7.8, 100 mM KCl, 0.1 mM EDTA (buffer A), disrupted by sonication and debris removed by centrifugation at 40000 $\times g$ for 45 min. Thermally unstable proteins were precipitated from the supernatant by heating to 65 °C for 15 min and removed by centrifugation as before. Protein was precipitated from the supernatant via the addition of ammonium sulfate to a concentration of 0.55 g mL^{-1} . The precipitate was solubilized in the minimum volume of buffer A and dialyzed against 1 L of identical buffer for 12 hr. Contaminating proteins were removed by size exclusion chromatography (HiPrep 26/60 Sephacryl S-300HR, GE Healthcare) and contaminating DNA by anion exchange chromatography (HiTrap Q FF, GE Healthcare). For the latter, protein solutions were loaded in buffer A and eluted by stepping to 50% buffer B (20 mM HEPES pH 7.8, 100 mM KCl, 1.0 M NaCl, 0.1 mM EDTA). Protein as isolated contained small quantities of iron that was removed as previously described (5) before exchanging into 100 mM MES pH 6.5 by centrifugation over a 10 kDa molecular weight cut off cellulose membrane (Millipore). Sample purity was assessed using SDS-PAGE and proteins judged to be free of DNA contamination once the ratio of absorbance at 277 nm (wild type) or 276 nm (Y40F) and 260 nm reached 1.5.

Gel filtration and native PAGE of SynFtn. The association state of purified SynFtn was assessed by loading 0.5 mL of 25 μM protein onto a HiPrep 26/60 Sephacryl S-200HR size exclusion column equilibrated with 100 mM MES pH 6.5. Column calibration under identical buffer conditions using protein standards of known molecular weight (Sigma) indicated a void volume of 106 mL and the expected dependence of elution volume on the logarithm of molecular weight. Assembly of SynFtn was also analysed by native PAGE. Samples were loaded onto 6% (v/v) polyacrylamide gels containing 60 mM HEPES, 40 mM imidazole (pH 7.2). Gels were conditioned by running in 60 mM HEPES, 40 mM imidazole at a current of 10 mA for 1 hr. Samples were then loaded in buffer containing 10% (v/v) glycerol and the gel run for a further 2 hr with a limiting current of 5 mA.

Crystallization and structure determination. Protein (10 mg mL^{-1}) exchanged into 20 mM MES pH 6.5 in 2 μL drops was mixed with an equal volume of well solution (0.1 M sodium acetate, 2.0 M NaCl pH 4.6) and equilibrated in sitting drops by vapour diffusion against 200 μL of the same well solution. Crystals of bi-pyramidal morphology appeared within 24 hours and grew to optimum size (100-150 μm) in \sim 1 week. Apo crystals were transferred to cryo-protectant comprising the well solution with pH adjusted to 6.5 containing 30% (v/v) glycerol prior to flash freezing in liquid nitrogen. Iron containing crystals were prepared by soaking for either 2 or 20 min in well solution containing 5 mM Fe^{2+} ions at pH 6.5. Crystals were then cryo-protected and frozen as above but using a solution containing 5 mM Fe^{2+} in addition to 30% (v/v) glycerol. Diffraction data were collected on beamline i04 of the Diamond Light Source except for the 20 min Fe^{2+} soak of wild type SynFtn, which was collected on beamline i03. The wavelength used was 0.9795 Å in all

cases. To reduce the likelihood of photoreduction of metal centers in iron-containing samples, exposure to X-rays was minimized subject to the requirement that the resulting diffraction datasets should be at least 95% complete and have a multiplicity of no less than 3.0 in the highest resolution bin. Additional, highly redundant anomalous scattering data were collected from crystals of similarly treated iron-containing crystals of wild type and Y40F *SynFtn* at wavelengths corresponding to the peak of the iron K-edge (around 1.7399 Å). All data were indexed and processed using XDS and Aimless as part of the automatic xia2 pipeline (44). Reprocessing was carried out as necessary using Aimless (45) as part of the CCP4 programme suite (46). Statistics are summarized in SI Appendix, Table S1 for X-ray data used for structure solution and refinement and in SI Appendix, Table S2 for data used for calculation of Bijvoet-difference Fourier (anomalous-scattering-density) maps.

Structure solution was performed by molecular replacement using phenix.phaser MR (47) with the 2.05 Å resolution structure of *E. coli* FtnA, pbd entry 1EUM (48), as the search model. In all cases the asymmetric unit contained a single copy of the protein monomer. Placement of metal ions was confirmed by reference to Bijvoet-difference Fourier maps calculated from anomalous scattering data (SI Appendix, Table S2). Model refinement employed iterative cycles using phenix.refine (48) and manual correction using COOT (49). No metal coordination restraints were applied to metal sites during refinement of iron-containing structures. Anisotropic temperature factor refinement was employed for all metal ions and their occupancies were manually adjusted to ensure that the average *B* factor of the metal fell within ± 14 % of the *B* factors of atoms of their environment (50). The coordination geometry of metal binding sites was analysed after refinement using the CheckMyMetal web server (51). Statistics relating to the metal binding sites in the refined structures can be found in SI Appendix, Table S3.

Kinetic analysis of iron oxidation and mineralization by *SynFtn*. Rates of iron oxidation were deduced from the rate of increase in $A_{340\text{ nm}}$ due to the resulting ferric-oxo species, be they iron bound at the ferroxidase center or in the mineral core. Typical assays employed 0.5 μM *SynFtn* in 100 mM MES pH 6.5 at 25 °C. Aerobic oxidation of ferroxidase center bound Fe^{2+} following addition to apo *SynFtn* was complete in ~ 20 s. Accordingly the kinetics of this process were monitored using stopped flow absorbance spectroscopy. 1.0 μM protein was mixed with an equal volume of Fe^{2+} of the appropriate concentration in 1 mM HCl using an Applied Photophysics Bio-Sequential DX.17MV spectrophotometer with a 1 cm path length observation cell. The time dependences of $A_{340\text{ nm}}$ increase were fitted to the sum of up to three exponential processes, encompassing rapid (r), intermediate (i) and slow (s) components, using OriginPro 8 (OriginLab):

$$\Delta A_{340}(t) = \Delta A_{340}^{(tot)} - \Delta A_{340}^r e^{-k_r t} - \Delta A_{340}^i e^{-k_i t} - \Delta A_{340}^s e^{-k_s t} \quad \text{Equation 1}$$

The extent to which oxidized iron vacates the ferroxidase centers of *SynFtn* was investigated by monitoring the regeneration of the rapid phase of iron oxidation associated with the apo protein. 1 μM protein was incubated with 400 μM Fe^{2+} at 25 °C for periods of 4, 5, 6 or 60 min prior to mixing with an equal volume of 48 μM Fe^{2+} in 1 mM HCl. An equivalent sample was incubated at 25 °C for 60 min followed by a further 19 hr at 4 °C. After re-equilibration at 25 °C the protein was mixed with an equal volume of 48 μM Fe^{2+} in 1 mM HCl as above. Data for comparison of iron oxidation kinetics to the time dependence of EPR intensities were obtained by mixing 8.33 μM *SynFtn* in 100 mM MES pH 6.5 with equal volumes of 0.6 mM Fe^{2+} in 1 mM HCl, giving a final

concentration of 4.17 μM *SynFtn* (100 μM in monomeric units) with an iron loading of 3.0 Fe^{2+} per monomer.

Attempts to detect formation and decay of a DFP intermediate during *SynFtn* catalyzed iron oxidation by monitoring at single wavelengths between 550 and 650 nm were inconclusive. Therefore, a SX20 stopped-flow spectrophotometer (Applied Photophysics) equipped with a diode-array multi-wavelength unit photomultiplier was employed to record spectra over the wavelength range 350-700 nm. SVD analysis of spectro-kinetic data acquired following mixing of 3 μM *SynFtn* in 100 mM MES pH 6.5 with 0.5, 1.0, 2.0 or 2.5 Fe^{2+} (in 1 mM HCl) per ferroxidase center at 25 °C was carried out using ProK (Applied Photophysics).

Iron mineralization assays employed a higher iron to protein stoichiometry of 400 Fe^{2+} per *SynFtn*. Dependences of absorbance on time were recorded on a Hitachi U-2900 spectrometer following manual mixing of 6.4 μL of a 50 mM Fe^{2+} solution in 50 mM HCl to a 1.6 mL sample of 0.5 μM protein in 100 mM MES pH 6.5. Initial rates of iron mineralization were deduced from the gradient of the initial linear region and an extinction coefficient for the mineral core calculated from the net absorbance change upon complete oxidation of the 200 μM Fe^{2+} added (active proteins) or an assumed extinction coefficient for the core of 2000 $\text{M}^{-1} \text{cm}^{-1}$ (E33A and E110A).

Electron paramagnetic resonance (EPR) spectroscopy. EPR spectra were recorded at 10 K or at room temperature on a Bruker EMX (X-band) EPR spectrometer. Low temperature measurements were performed with the use of an Oxford Instruments liquid helium system and a spherical high-quality ER 4122 SP 9703 Bruker resonator. Protein samples in EPR tubes were mixed with the appropriate volume of a 25 mM stock Fe^{2+} solution and frozen at least 10 s thereafter by plunging the tubes into methanol cooled with solid CO_2 . Samples frozen at times < 10 s (the Rapid Freeze-Quench technique, RFQ) were prepared by mixing equal volumes of an 8.33 μM protein solution (200 μM in subunit monomer) and a 0.6 mM Fe^{2+} solution in an Update Instrument 715 Syringe Ram Controller (Madison, WI) and ejecting the mixture onto the surface of a rapidly rotating aluminium disk maintained at liquid nitrogen temperature (52). Final protein concentration was 4.17 μM (100 μM in monomer) in all cases. Stopped flow absorbance measurements performed at comparable protein concentration to RFQ EPR showed that at ambient temperature the rapid oxidation phase of wild type *SynFtn* would be >50% complete in 50 ms, the shortest available freezing time, but that the rate of this phase could be slowed by lowering the reaction temperature (SI Appendix, Fig. S3A). Consequently, for RFQ measurements of wild type *SynFtn*, the chamber housing the sample syringes was packed with a water/ice mixture. The equivalent measurements on variant Y40F were conducted at ambient temperature because it lacks a rapid oxidation phase, enabling a more reliable comparison to stopped flow data.

Proteins were in 100 mM MES pH 6.5 and Fe^{2+} solutions in 50 mM HCl (slow freezing) or 1 mM HCl (RFQ). Instrument parameters for EPR measurements were as follows: microwave frequency $\nu_{\text{MW}} = 9.4657$ GHz, modulation frequency $\nu_{\text{M}} = 100$ kHz, time constant $\tau = 82$ ms, microwave power = 3.19 mW or 0.05 mW (for 100 G wide free radical scans), modulation amplitude $A_{\text{M}} = 5$ G or 3 G (for free radical scans), scan rate $\nu = 22.6$ Gs^{-1} or 0.6 Gs^{-1} (for free radicals). Quantification of paramagnetic species was achieved by comparison of integrated intensities to those of a 1 mM Cu[EDTA] solution (Tyr radical and MVFC) or a 35 μM solution of

Fe³⁺ containing Ferric Binding Protein from *Neisseria gonorrhoeae* (high spin mononuclear ferric iron with $g = 4.3$).

Room temperature EPR measurements were performed with the use of a Bruker AquaX system with a 4103TM Bruker resonator. A homemade mixing device was used to mix equal volumes of two liquid components and fill the AquaX capillary bundle. The dead volume of the mixing device was 130 μL and the minimal volume to be taken by each input syringe was 180 μL . The dead time of the mixing device was 2 s. 5,5-Dimethyl-1-pyrroline *N*-oxide (DMPO, Sigma) was used to spin-trap short lived radicals in the room temperature experiments. DMPO was dissolved in water at 228 mM concentration and filtered over activated charcoal. EPR spectra were recorded before and after this filtration step to measure the removal of paramagnetic impurities. In a typical experiment, a solution of protein with DMPO was mixed with equal volume of freshly prepared Fe²⁺ so that the final concentrations were as follows: 5.4 μM protein, 40 mM DMPO, 375 μM Fe²⁺ - in 100 mM MES buffer, pH 6.5. The reference spectra of the superoxide radical DMPO adduct and hydroxyl radical DMPO adduct were obtained in a xanthine oxidase / xanthine system (0.16 units xanthine oxidase (Sigma), 0.4 mM xanthine (Sigma), 100 mM DMPO - in 50 mM sodium phosphate buffer, pH 7.7) and in a Fenton reagent system (1 mM Fe²⁺, 1 mM H₂O₂, 40 mM DMPO), respectively. Instrumental parameters were: microwave frequency $\nu_{\text{MW}} = 9.799$ GHz, microwave power = 12.57 mW, modulation frequency $\nu_{\text{M}} = 100$ kHz, modulation amplitude $A_{\text{M}} = 1$ G, time constant $\mu = 40.96$ ms, scan rate $v = 0.83$ Gs⁻¹.

Spectral simulations of MVFC EPR signals were performed using WinEPR SimFonia (Bruker Biospin) using the parameters reported in Table 1. The free radical EPR spectra were simulated using SimPow6 (53) while the simulation parameters were found by using the Tyrosyl Radical Spectra Simulation Algorithm, TRSSA (29). Assignment of the radical in *SynFtn* to a specific tyrosine site was performed on the basis of correspondence drawn between the rotation angle in the phenoxyl ring in the radical, found from the simulation, and the angles of the ring in the tyrosine residues as seen in the crystal structure. The analysis of these angles is reported in SI Appendix, Table S5.

Oximetry measurements. The stoichiometry of the *SynFtn* catalyzed iron-O₂ reaction was deduced by monitoring the decrease in the dissolved O₂ concentration in a 1 μM solution of protein in 2 mL of 100 mM MES pH 6.5 following additions of 4 μL aliquots of 24 mM Fe²⁺ in 50 mM HCl (48 μM final concentration) using a Clark electrode (Hansatech Instruments Oxygraph⁺). The iron:O₂ ratios reported are the mean of three independent measurements under each reaction condition together with the standard error of the mean. The ability of *SynFtn* to reduce O₂ at the surface of a growing mineral core was assessed by adding 33 μL of 24 mM Fe²⁺ to apo protein as above (400 Fe²⁺/*SynFtn*). Reaction was judged complete once the O₂ concentration became invariant within error for 20 s and a further 4 μL of Fe²⁺ added.

Detection of hydrogen peroxide. Release of H₂O₂ was detected using the Amplex Red assay (Thermo Fisher Scientific). The 560 nm absorbance of samples containing 0.1 μM wild type or Y40F *SynFtn*, incubated with varying ratios (24-72 equivalents) of Fe²⁺, were compared to a standard plot ($y = 0.009x + 0.0379$, $R^2=0.997$) constructed using 0-5 μM H₂O₂ prepared from dilution of a 20 mM standard solution into identical buffer (sodium phosphate pH 7.4).

SynFtn interactions with copper. The possibility of copper binding to *SynFtn* was investigated under two conditions. Direct interaction with *SynFtn* was probed by incubating 0.5 μM protein with 200 μM copper either added aerobically as Cu^{2+} or anaerobically as Cu^+ for 60 minutes. Separate samples containing Cu^+ were either maintained in an anaerobic environment or exposed to O_2 during incubation. Weakly associated or unbound copper was then removed using a desalting column (Sephadex G-25M, GE Healthcare). 0.5 mL aliquots of the eluted sample were then digested by mixing with an equal volume of 68% HNO_3 and heating to 80 $^\circ\text{C}$ for 16 hours. Cooled samples were neutralized by adding 2.5 mL of saturated sodium acetate and incubated at 37 $^\circ\text{C}$ for 15 min following the addition of 0.5 mL each of NH_2OH , 3.2 mg mL^{-1} , and bathocuproinedisulfonic acid (BCS), 5.0 mg mL^{-1} . Copper content was determined by comparison of the absorbance at 484 nm to a standard plot constructed using copper solutions (0-100 μM) treated identically to protein solutions eluted from the G-25 column ($y = 0.0014x - 0.0034$, $R^2=0.998$). The same assay was used to probe incorporation of copper into the mineral core during deposition but with 4.1 μM *SynFtn* incubated aerobically with 1.6 mM Fe^{2+} and 200 μM of either Cu^+ or Cu^{2+} . Absorbance monitored mineralization assays conducted on 0.5 μM protein demonstrated that incorporation of iron into the protein core proceeds at 90% of the rate observed in the absence of copper. A standard plot constructed from solutions containing both Fe^{2+} and Cu^{2+} indicated that the presence of the former did not interfere with copper detection using BCS.

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Figure Legends

Figure 1. Comparison of the ferroxidase centers of *SynFtn* and Human H-chain ferritin (HuHF). (A) The Apo ferroxidase center of *SynFtn* at pH 6.5, pdb entry 5OUW with the double difference Fourier ($2mF_o-DF_c$) map in the vicinity of the iron binding sites contoured at 1.1σ represented as a blue mesh. Metal binding sites, labelled A and B, are occupied by water molecules. Site A is predicted to have higher affinity for iron than site B. (B) The ferroxidase center of *SynFtn* following a 2 min soak in a solution containing 5 mM Fe^{2+} at pH 6.5 with the anomalous difference map contoured at 8σ in the same region, pdb entry 6GKC. (C) The ferroxidase center of HuHF following a 30 min. soak in aerobic Fe^{2+} containing solution at pH 8 from pdb entry 4YKH(19). (D) As in (B) other than the soaking time for the crystal was increased to 20 min and the anomalous difference map contoured at 10σ , pdb entry 6GKA. Iron and water oxygen atoms are shown as orange and red spheres, respectively. Side chains of selected residues are shown in stick representation. Dashed lines in panels B and D indicate metal ligand bonds to the higher occupancy, Fe_A , and lower occupancy, Fe_B , iron sites. We note that the Fe-O-Fe core of the HuHF structure is more similar to the 20 min soak structure of *SynFtn* than it is to the 2 min soak structure, in which Fe_B and the bridging O(H) are in different positions, consistent with the assignment of the 2 min soak structure to a mixed valent form.

Figure 2. Absorbance monitored iron- O_2 reactivity of *SynFtn*. (A) pH dependence of the rate of mineral core formation on consecutive addition of aliquots of 400 Fe^{2+} per *SynFtn* up to the maximum core capacity of 2000 $Fe/SynFtn$. (B) iron oxidation during the first second following mixing of increasing ratios of Fe^{2+} to 0.5 μM *SynFtn* at pH 6.5. Data shown in gray, black traces represent progress curves calculated from Equation 1 using the values of ΔA_{340nm} and k specified in SI Appendix, Fig. S2. Broken red line indicates the predicted increase in absorbance at 340 nm for oxidation of 24 Fe^{2+}/Ftn . (C) equivalent traces to (B) over the first twenty seconds following mixing of increasing ratios of Fe^{2+} to 0.5 μM *SynFtn* at pH 6.5 showing the second, slower phase of oxidation.

Figure 3. EPR monitored iron- O_2 reactivity of *SynFtn*. (A) The EPR signal attributed to a mixed valent (Fe^{2+}/Fe^{3+}) form of the ferroxidase center (MVFC) in *SynFtn* frozen 12 s after initiating iron oxidation (iron added at 80 Fe^{2+}/Ftn) and its modeling as a sum of two signals simulated with the parameters listed in Table 1. Experimental data is truncated to omit the region of overlap with the tyrosyl radical (Fig. 5). (B) Intensity of the MVFC signal at increasing iron loading in samples of 4.17 μM *SynFtn* in 100 mM MES pH 6.5 frozen 10 ss after mixing. Response from the apo protein and protein loaded with 80 Fe^{2+} per Ftn are shown in black, intermediate loadings (12, 24 and 48 equivalents of Fe^{2+}) are shown in gray. (C) Increase in signal intensity in the high field region ($g < 2$) assigned to the MVFC in samples of 4.17 μM *SynFtn* frozen 50 – 365 ms after mixing with 72 equivalents of Fe^{2+} . Thick traces are the shortest (50 ms) and longest (365 ms) ageing times shown, the latter corresponding to maximum signal intensity. Traces at intermediate freezing times are shown in gray. The EPR signal minimum at ~ 3800 G shifts in both sets of spectra, for iron load (B) and reaction time (C), due to increasing relative input of the ‘minor’ MVFC component (A), which has a greater g_z value than the ‘major’ one (Table 1). (D) Intensity plots showing the formation and decay of the MVFC (red circles), mononuclear Fe^{3+} (black squares) and tyrosyl

radical (blue triangles) detected by low temperature EPR spectroscopy over the first 10 s of iron oxidation (72 Fe²⁺/Ftn) by *SynFtn*. Solid lines represent fits of the data, yielding apparent first order rate constants of 3.0 s⁻¹ (MVFC), 2.0 s⁻¹ (Fe³⁺) and 1.2 s⁻¹ (Tyr radical) for the formation phase, and 0.2 s⁻¹ (MVFC), 0.15 s⁻¹ (Fe³⁺) and 0.17 s⁻¹ (Tyr radical) for the decay phase. Data for the MVFC can be compared with those for Fe²⁺ oxidation measured by stopped-flow absorbance at 10 °C (SI Appendix, Fig. S3A), for which apparent first order rate constants of 8.3 s⁻¹ (rapid phase) and 0.3 s⁻¹ (slow phase) were obtained.

Figure 4. O₂ consumption during Fe²⁺ oxidation by *SynFtn*. (A) Change in dissolved O₂ concentration following successive additions of 48 equivalents (48 μM) of Fe²⁺ to 1 μM apo *SynFtn* in 100 mM MES pH 6.5 at 20 °C. (B) As for (A) but following addition of 48 equivalents of Fe²⁺ to 1 μM *SynFtn* containing a freshly prepared mineral core of 400 Fe³⁺/*SynFtn*.

Figure 5. The transient tyrosyl radical of *SynFtn*. (A) EPR signal due to a tyrosyl radical in *SynFtn* frozen 12 s after initiating iron oxidation (iron added at 80 Fe²⁺/Ftn). The simulation parameters are reported in SI Appendix, Table S4. (B) Intensity of the tyrosyl radical signal at increasing iron loading in samples of 4.17 μM *SynFtn* in 100 mM MES pH 6.5 frozen 10 s after mixing. Response from the apo protein and protein loaded with 80 Fe²⁺ per Ftn are shown in black, intermediate loadings (12, 24 and 48 equivalents of Fe²⁺) are shown in gray. (C) Increase in signal intensity due to the radical at approximately g = 2 in samples of 4.17 μM *SynFtn* frozen 50 – 365 ms after mixing with 72 equivalents of Fe²⁺. Thick traces are the shortest (50 ms) and longest (365 ms) ageing times shown. Traces at intermediate freezing times are shown in gray.

Figure 6. Absorbance and EPR monitored iron-O₂ reactivity of the Y40F variant of *SynFtn*. (A) Increase in A_{340 nm} over the first second following aerobic mixing of between 6 and 96 Fe²⁺ per *SynFtn*. Data in gray, black traces show the progress curves calculated from the parameters plotted in SI Appendix, Fig. S5C, D. Red trace shows the response following addition of 96 Fe²⁺ to wild type *SynFtn* for comparison. (B) data as in (A) showing the response over the first 20 s following mixing. Over this time period, only a single exponential was required to fit the traces at all iron loadings and the apparent rate constants describing this process were in good agreement with those for the slow phase of oxidation observed in wild type *SynFtn*. (C) A_{340 nm} as a function of time following mixing of a solution of 4.17 μM Y40F *SynFtn* with 72 equivalents of Fe²⁺ at 20 °C. Data shown as gray circles, black trace represents a bi-exponential function fitted through the data, described by apparent first order rate constants of 1.5 and 0.16 s⁻¹. (D) Low temperature EPR signal attributed to the formation of a mixed valent (Fe²⁺/Fe³⁺) form of the ferroxidase center (MVFC) in Y40F *SynFtn* frozen 15 s after aerobic addition of 72 Fe²⁺ per *SynFtn* and its modeling as a sum of two EPR signals simulated with the parameters listed in Table 1. Experimental data is truncated to omit the region of overlap with intensity from the peroxy (ROO[•]) radicals observed in place of the tyrosyl radical of the wild type protein (SI Appendix, Fig. S4B). (E) and (F) Time dependence of Y40F *SynFtn* EPR. (E) Increase in signal intensity in the high field region (g < 2) assigned to the MVFC in solutions equivalent to that in (C) frozen 50 – 2965 ms after mixing with 72 equivalents of Fe²⁺. Thick traces are the shortest (50 ms) and longest (2965 ms) ageing times shown, the latter corresponding to maximum signal intensity. Traces at intermediate freezing times are shown in gray. (F) Normalized intensity of observed EPR signals as a function of time

after mixing of Y40F *SynFtn* and 72 equivalents of Fe^{2+} . Sample conditions were as in (E). MVFC intensity is shown as black circles, and mononuclear Fe^{3+} as blue. The black trace shows the predicted kinetics of a species formed in a first order reaction at the faster rate derived in (C) and consumed at the slower.

Figure 7. Proposed catalytic cycle of *SynFtn*. (A) Apo ferroxidase center binds two Fe^{2+} ions from solution. (B) Two electron reduction of O_2 to hydrogen peroxide leads to oxidation of a single Fe^{2+} ion, thus yielding the MVFC, and a radical on Tyr40. (C) The Tyr radical is transient and only observed in the subset of subunits where the partner subunit's ferroxidase center is unoccupied. In the remainder, rapid electron transfer from $\text{Fe}^{2+}/\text{Fe}^{2+}$ or monomeric Fe^{2+} bound at a second ferroxidase center results in quenching of the radical. (D) Slow reaction of the MVFC with a second molecule of O_2 , accompanied by transfer of a second electron from the paired ferroxidase center results in formation of the unstable bridged di-ferric center observed in other ferritins and a second molecule of hydrogen peroxide. The kinetics of this process were unaltered in a Y40F variant, suggesting that Tyr40 is not required for transfer of the second electron. (E) Hydration and translocation of oxidized iron from the ferroxidase center to the internal cavity results in formation of mineral core and regeneration of apo ferroxidase centers. Overall four ferroxidase center bound Fe^{2+} ions are oxidized reducing two molecules of O_2 to H_2O_2 accounting for the observed 2:1 stoichiometry.

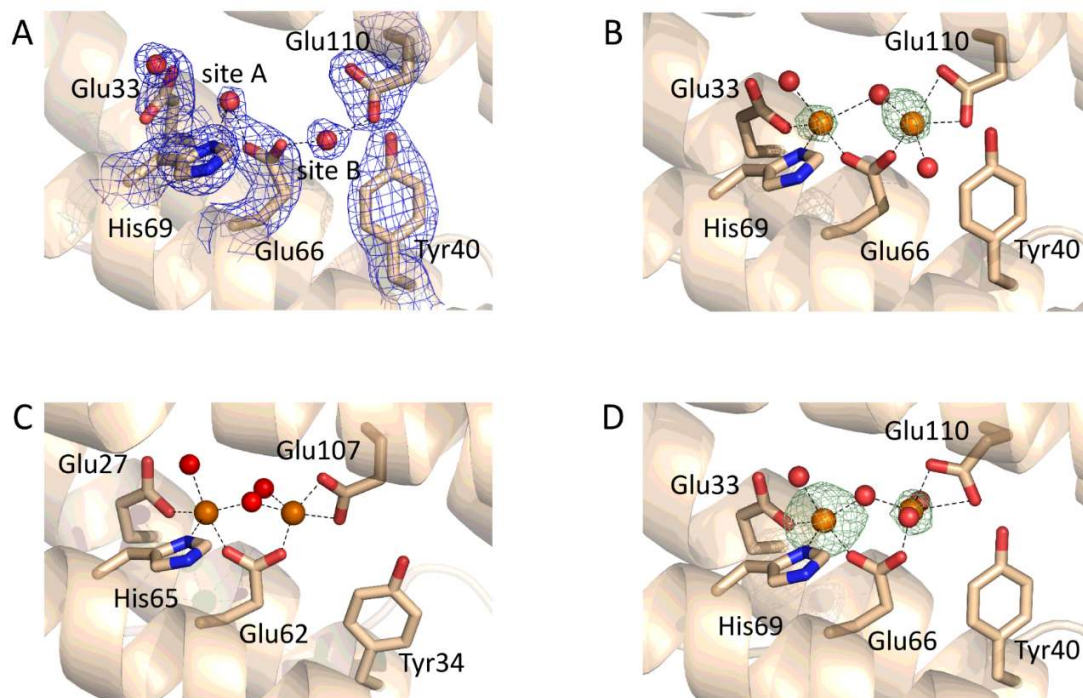


Fig. 1.

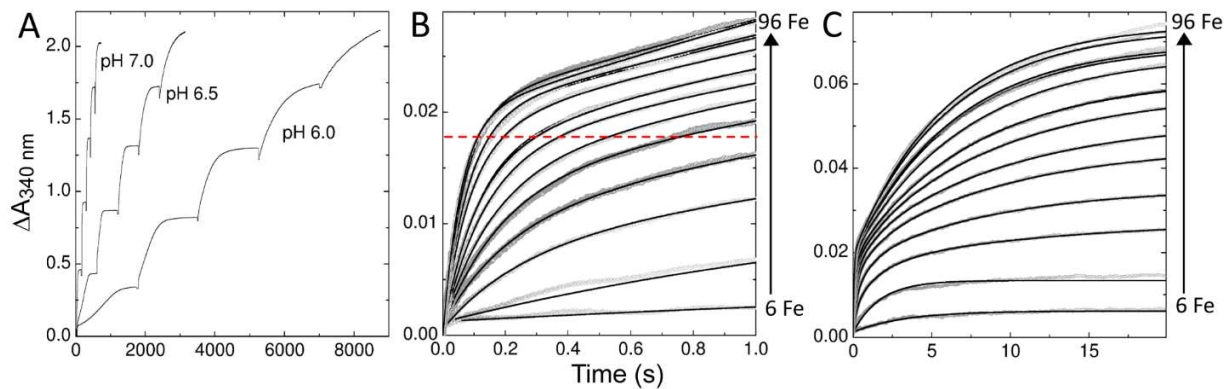


Fig. 2.

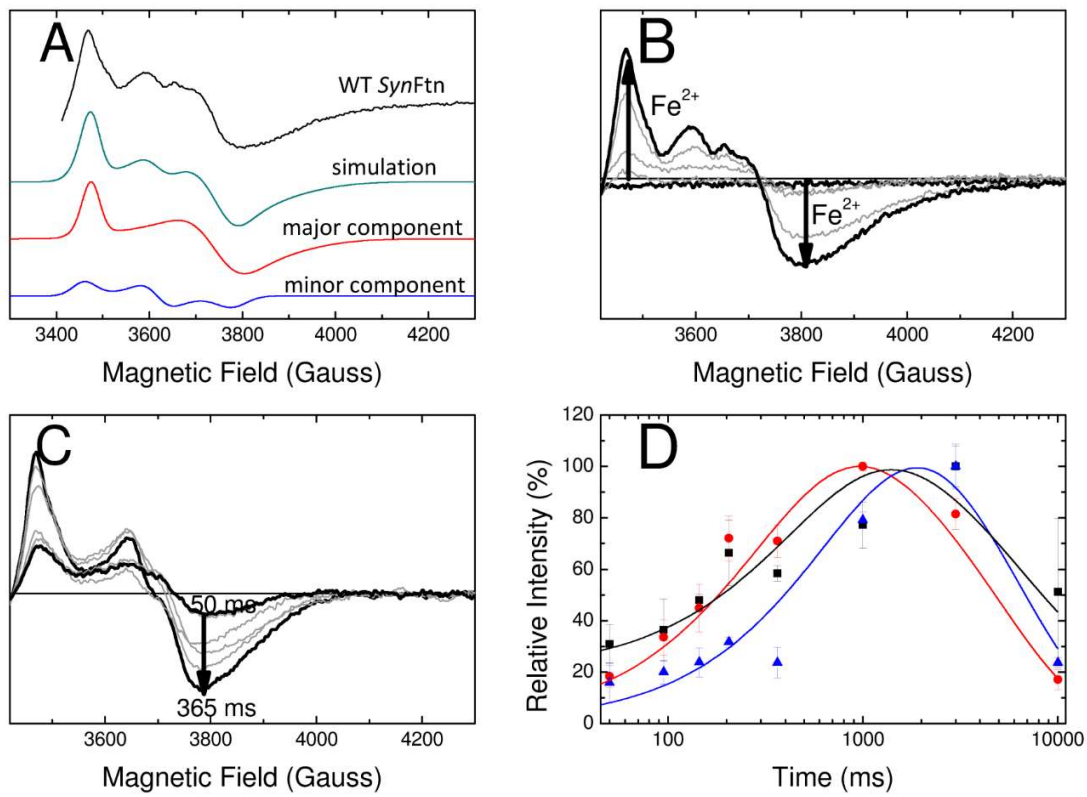


Fig. 3.

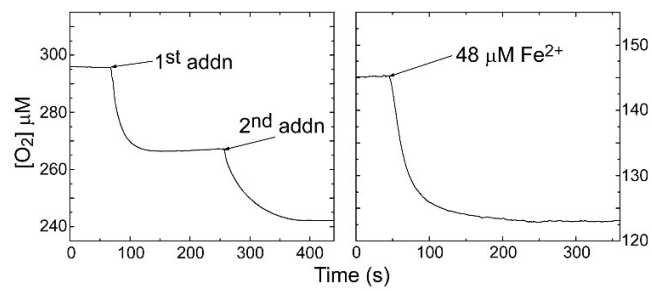


Fig. 4.

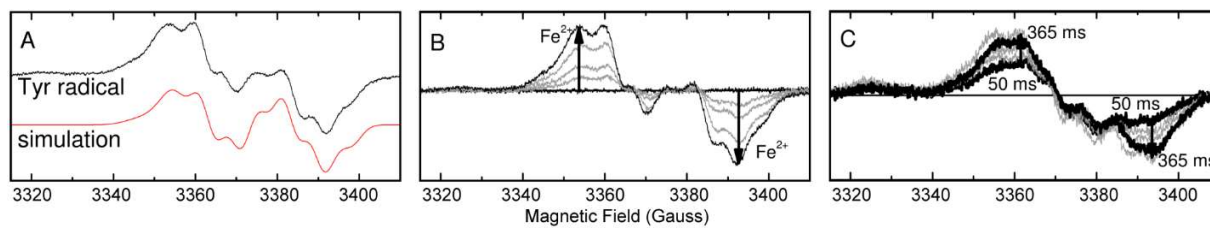


Fig. 5.

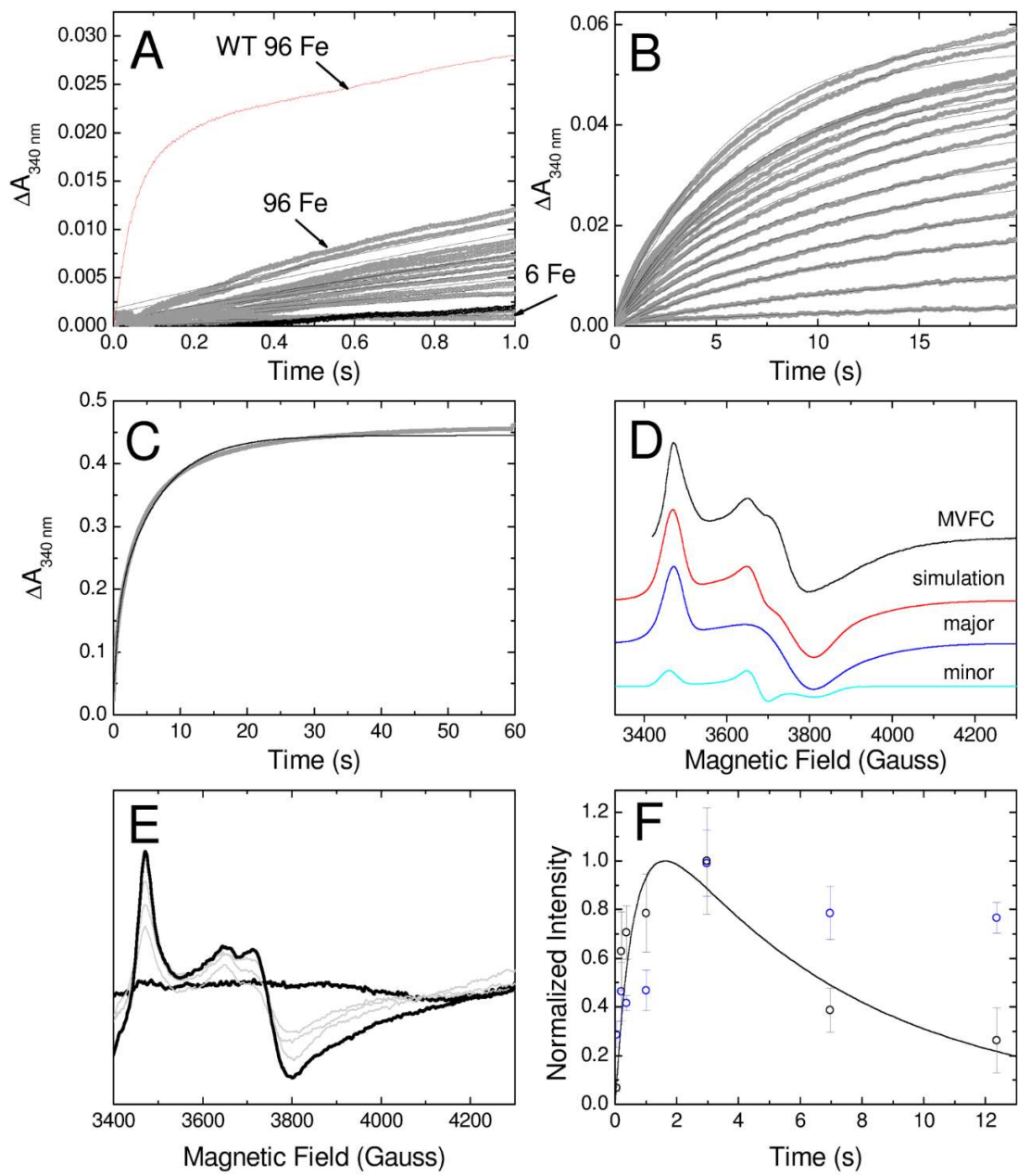


Fig. 6.

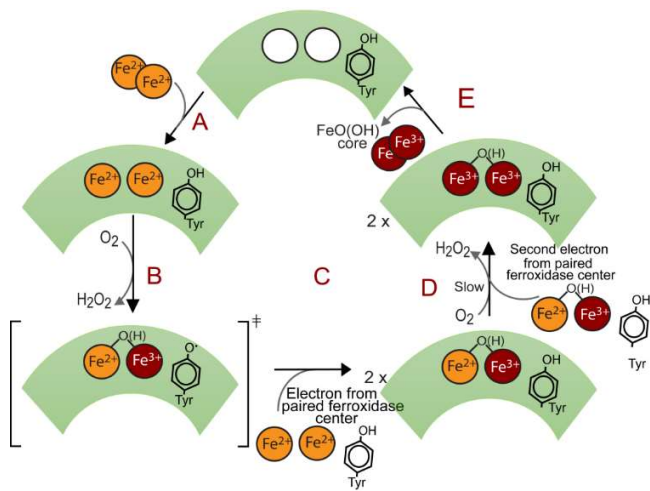


Fig. 7.