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Crystal Structure of Miner1: The Redox-active 2Fe-2S Protein Causative in Wolfram Syndrome 2

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The endoplasmic reticulum protein Miner1 is essential for health and longevity. Mis-splicing of CISD2, which codes for Miner1, is causative in Wolfram Syndrome 2 (WFS2) resulting in early onset optic atrophy, diabetes mellitus, deafness and decreased lifespan. In knock-out studies, disruption of CISD2 leads to accelerated aging, blindness and muscle atrophy. In this work, we characterized the soluble region of human Miner1 and solved its crystal structure to a resolution of 2.1 Å (R-factor = 17%). Although originally annotated as a zinc finger, we show that Miner1 is a homodimer harboring two redox-active 2Fe-2S clusters, indicating for the first time an association of a redox-active FeS protein with WFS2. Each 2Fe-2S cluster is bound by a rare Cys3-His motif within a 17 amino acid segment. Miner1 is the first functionally different protein that shares the NEET fold with its recently identified paralog mitoNEET, an outer mitochondrial membrane protein. We report the first measurement of the redox potentials (Em) of Miner1 and mitoNEET, showing that they are proton-coupled with Em ~ 0 mV at pH 7.5. Changes in the pH sensitivity of their cluster stabilities are attributed to significant differences in the electrostatic distribution and surfaces between the two proteins. The structural and biophysical results are discussed in relation to possible roles of Miner1 in cellular Fe-S management and redox reactions.

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Introduction

Miner1†, originally annotated as a zinc finger protein of unknown function, is actually a unique redox-active Fe-S protein. The gene encoding Miner1, CISD2 (CDGSH iron sulfur domain 2‡), found at chromosomal location 4q24, was identified recently as the carrier of a mutation that causes Wolfram Syndrome 2 (WFS2). The pre-mRNA of CISD2 has two splice sites, creating three exons and two introns (Fig. 1). Correct splicing is essential for the proper translation and function of Miner1. It localizes to the endoplasmic reticulum (ER) when properly expressed and folded. The mutation found in WFS2 patients is a G to C base-pair transversion in pre-mRNA CISD2 that causes a splicing error, resulting in removal of the second exon and introduction of a premature stop codon, which eliminates 75% of the protein transcript (Fig. 1). Indeed, the region binding the 2Fe-2S cluster is eliminated completely. Although initially healthy, patients with WFS2 experience a decreased life expectancy, early onset diabetes...
mellitus, optical atrophy, sensorineural deafness, and a significant bleeding tendency. Studies of structure and function of Miner1 are expected to provide critical information on the fundamental cause of the disease and reasons for the decrease in life expectancy.

CISD2 is on chromosome 4, a genetic locus associated with longevity. Investigations focused on the longevity locus where researchers disrupted CISD2 in mice found that the elimination of Miner1 resulted in decreased life expectancy and reduced general health. These knockout mice show accelerated aging, blindness, an abnormal skeleton, and muscle atrophy, effects that are very similar to those described in WFS2 patients. The mouse ortholog is 96% identical with the human Miner1 and is highly conserved (94–100%) in other mammals (Supplementary Data Fig. S1). Obtaining the Miner1 protein structure provides a basis for understanding the functional consequences of mutations in human and animal studies.

Miner1 belongs to a newly discovered family, which includes the 2Fe-2S-containing outer mitochondrial membrane (OMM) protein mitoNEET. In contrast, Miner1 localizes to the ER. It contains an atypical ER localization sequence, an N-terminal transmembrane domain, and a CDGSH 2Fe-2S domain. To facilitate yields and structural studies, we focused on the soluble domain of the protein and made a point mutation at the only non-conserved free cysteine (Supplementary Data Fig. S1). In this work, we show that the Cys mutation does not affect the properties and report the crystal structure of the soluble region of Miner1 (C92S) solved at 2.1 Å resolution using Fe-MAD phasing. The structure is homodimeric with a scaffold similar to that of the OMM protein mitoNEET, but has a distinct surface topology and differences in the charge distribution. We report and compare redox and stability measurements as functions of pH for both Miner1 and its paralog mitoNEET.

Fig. 1. Elimination of functional Miner1 protein leads to multiple symptoms in both human and mice models. CISD2, shown at the top, codes for an endoplasmic reticulum (ER) protein Miner1 (left-hand side). Amr et al. showed recently that Wolfram Syndrome 2 is attributed to a single base pair conversion in the CISD2 gene. This mutation causes a splicing error that completely eliminates exon2 and creates a premature stop codon in exon 3 (right-hand side). For these reasons, 75% of the protein sequence is not translated, and no Miner1 is properly assembled in the ER (lower part of right-hand side). The absence of this protein causes a wide range of symptoms in WFS2 patients, including diabetes mellitus and optical atrophy. In an independent study, CISD2 knockout mice were constructed with a coding error that causes incorrect splicing, resulting in an inoperative gene. Knock-out mice are also symptomatic, with similar signs of early aging and optical atrophy.
Results

Isolation and UV-visible spectroscopy of the ER protein Miner1

In an effort to understand the properties of this protein, we produced a soluble form of recombinant human Miner1 that corresponds to amino acids 57–135 (lacking the amino-terminal targeting and transmembrane sequences). The protein was fused to a cleavable His tag to facilitate purification. As a result, the construct includes an additional four amino acids (GSHM) at the N-terminus. Early attempts at purification were hindered by protein instability and aggregation. Although the use of reducing agent decreased aggregation, yields remained insufficient for structural studies. Therefore, we replaced the one free non-conserved cysteine (C92S). The resultant mutant protein, Miner1 (C92S), has the same optical signature of the native Miner1 (Fig. 2). Furthermore, the construct had the desirable effects of improved yields of purified protein and a decreased tendency to aggregate. Since the point mutation does not change the spectra of Miner1, we refer to Miner1(C92S) simply as Miner1.

The overall fold of Miner1

The isolated purified recombinant protein showed CD spectra (Supplementary Data Fig. S2) indicative of a well-folded protein with a chiral center. The protein was amenable to crystallization. X-ray diffraction intensities were collected at Stanford Synchrotron Radiation Laboratory (SSRL) BL7-1 and BL9-2 to 2.1 Å (Table 1). Miner1 crystallized in the orthorhombic space group $P_2_1_2_1_2_1$ with unit cell parameters $a=49.90$ Å, $b=48.58$ Å, $c=74.10$ Å, $\alpha=\beta=\gamma=90^\circ$ (Table 1). The Matthews coefficient ($V_m$) of the crystal was 2.0 Å$^3$/Da with an estimated solvent content of 37% ($v/v$).

Experimental phases for model building were determined with Fe-MAD. The model was refined using 2.1 Å data to an R-factor of 17.0% ($R_{free}=21.6\%$). The high quality of the electron density is shown in Fig. 3a. The refined model reveals a parallel homodimeric structure that includes the cytoplasmic fragment of each protomer from Asp68 to Glu134 on protomer A and to Val135 on protomer B (Fig. 3a). A total of 15 amino acids at the N-termini are disordered and not resolved in the electron density. The homodimer is tightly packed with mean distances of 2.1 Å ($\langle R_M \rangle = 2.1\%$).

Table 1. Summary of crystal parameters, data collection, and refinement statistics for Miner1

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B. Model and refinement statistics

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Stereocchemical parameters

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<td>ESU based on $R_{free}$ (Å)</td>
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ESU, estimated overall coordinate error. $R_{symm} = \Sigma |I_i - \langle I \rangle| / \Sigma I_i$, where $I_i$ is the scaled intensity of the $i$th measurement and $\langle I \rangle$ is the mean intensity for that reflection.

$R_{cryst} = \Sigma |F_{calc} - F_{obs}| / \Sigma |F_{calc}|$, where $F_{calc}$ and $F_{obs}$ are the calculated and observed structure factor amplitudes, respectively. $R_{free}$ is $R_{cryst}$ for 5.0% of the total reflections chosen at random and omitted from the refinement. Values in parentheses are for the highest-resolution shell.

* Typically, the number of unique reflections used in refinement is less than the total number that were integrated and scaled. Reflections are excluded due to systematic absences, negative intensities, and rounding errors in the resolution limits and cell parameters.
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1800 Å² of buried surface area at the interface, as calculated using PISA. Model validation using the MOLPROBITY structure validation tool indicates that 97% of the amino acid residues are in the favored region of Φ/Ψ space.

Miner1 is folded into two spatially distinct subregions: a β-rich or β-cap domain and a helical 2Fe-2S binding or cluster binding domain (Fig. 3b). The β-rich domain contains a strand swap from opposite ends of the primary sequence to form the β-cap structure. This domain contains 28 residues within β strands with two antiparallel strands coming from one protomer and the third parallel coming from the other (Fig. 3b). These two strand-swapped sheets pack together to form the β-cap sandwich domain and form the narrowest part (~15 Å across) of the structure (Fig. 3b). The structure confirms the presence of two 2Fe-2S clusters expected from the optical spectrum (Fig. 2). These clusters are separated by approximately 16 Å (center-to-center) within the larger helical cluster binding domain (~30 Å across) (Fig. 3b). A structural similarity search using the DALI server revealed that this fold shows significant similarity only to the recently determined new structural NEET fold of mitoNEET. This is the first example of another protein that shares the NEET fold.

**Structure of the 2Fe-2S clusters**

The cluster binding domain consists of the sequences Asp68–Val83 and Cys99–Gly126 on each protomer (Fig. 3b). Each 2Fe-2S cluster is cradled by a relatively short polypeptide chain, Cys99–Gly112.

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Fig. 3. Structural organization and domain topology of dimeric Miner1. (a) The backbone tracing of homodimeric Miner1 (upper). Protomers are colored purple and cyan together with the observed 2F₀ – Fc electron density (gray) map contoured at 1.5σ. A 2Fe-2S cluster is present in each protomer (Fe, red spheres; S, yellow). The two protomers are related by a dyad axis along the vertical direction in the plane of the paper (see arrow). Lower: An expanded view of one 2Fe-2S cluster (rotated ~90° counterclockwise along the dyad axis) showing the cluster and iron ligands (O, red; N, blue; C, green) and the corresponding observed 2F₀ – Fc electron density (gray) map contoured at 2.0σ. The amino acid ligands are indicated. Rendered with PyMol (http://pymol.sourceforge.net/). (b) Ribbon diagram highlighting the two domains and protomer interactions within the Miner1 homodimer: a six-stranded β sandwich forms an intertwined β cap and a larger cluster-binding domain carries two 2Fe-2S clusters (one per protomer). Below is the amino acid sequence of the construct. Coded segments contributing to each domain are highlighted on the primary sequence with blue for the β cap and magenta for the cluster-binding domains. The 2Fe-2S binding cradle is located sequentially between two parts of the β cap domains.
This segment contains the three coordinating Cys ligands (Cys99, Cys101, Cys110) and the single coordinating His ligand (His114). His114 is near the N-terminus of the α-helix within the cluster-binding domain (Ala113–Thr121) (Fig. 3).

Cys110 and His114, which coordinate the outermost Fe, are solvent-accessible, as illustrated in Fig. 4 by their semitransparent surfaces. In contrast, Cys99 and Cys101, which coordinate the innermost Fe, are buried within the structure (Fig. 4). Located sequentially between the innermost Cys ligands is Arg100. It forms interprotomer interactions near the 2Fe-2S clusters. Its guanidinium group forms a direct interprotomer hydrogen bond with the amide side chain of Asn84 of the other protomer. In addition, Arg100 interacts with the backbone oxygens of Cys99 and Pro108 of the other protomer via an internal water molecule (Fig. 4) locking Pro108 into a cis conformation.

**The proton-coupled redox potential of the 2Fe-2S clusters of Miner1 and its paralog mitoNEET**

A property of Miner1 that relates to possible redox activity is its redox potential $E_m$. The $E_m$ values of both Miner1 and mitoNEET, i.e. $[2\text{Fe}-2\text{S}]^{2+}/[2\text{Fe}-2\text{S}]^+$, were measured spectroscopically by the change in visible absorbance that occurs upon reduction (Supplementary Data Fig. S3). The fraction oxidized was measured by the change in absorbance at 320 nm to 600 nm as a function of time at 35 °C (Supplementary Data Fig. S4). A half-life of 700 min (pH 7.1) was determined from the observed signal change as a function of time. Surprisingly, given the similar fold, Miner1 was more stable at pH 5.3 (Supplementary Data Fig. S4). In addition, the decay is more non-exponential, suggesting a more complex unfolding energy landscape. To further explore differences, we performed a more detailed investigation of the stability as a function of pH.

Investigation of the stability as a function of pH provides additional information on interactions between titrating (charged) residues and the cluster. In contrast to mitoNEET, the lifetime for Miner1 varies by only ∼5-fold from pH 5.3 to 7.1 compared to ∼300-fold for mitoNEET (Fig. 5b). Thus, the release of the 2Fe-2S clusters of Miner1 is orders of magnitude less sensitive to pH as that of mitoNEET.

**Prominent differences in the charge distribution of Miner1 and mitoNEET and in the packing of the β-cap domain**

The two proteins display prominent differences in the distribution of charges. Overall, Miner1 has both fewer charged (Asp and Lys) and His residues (Fig. 6b). The homologous sites of polar Asn87 and apolar Leu93 and Leu120 in Miner1 are acidic (Asp61, Asp67 and Asp93) in mitoNEET. Other differences include: (i) polar Thr106 in Miner1 replaced with a basic Lys79 in mitoNEET; (ii) acidic Glu85 by neutral Ala59; and (iii) Lys74 and Asn84 by His48 and His58, respectively. These changes result in a more positively charged soluble domain for Miner1.

![Fig. 4. The 2Fe-2S cluster binding cradle. View of the 2Fe-2S cluster (Fe as brown and S as yellow spheres) from a perspective rotated ~90° along an axis perpendicular to the dyad axis from that shown for the cluster binding site in Fig. 3a. The amino acids belonging to the individual protomers are shown in purple and cyan, respectively. The two 2Fe-2S cradles are related to each other via a 180° rotation along the dyad symmetry axis of the dimer located approximately perpendicular to the page. Cys110 and His114 bind the outermost Fe, while the innermost Fe is bound by Cys99 and Cys101 (as indicated). The solvent-accessible His114 is located at the end of the prominent α-helix in the cluster binding domain (Fig. 3). Two additional residues, Arg100 and Asn84, form an interprotomer bifurcated hydrogen bond within the interior of the protein dimer. Each Arg100 also forms a potential hydrogen bond with one of two internal water molecules (shown as green spheres and labeled as Wat) that form additional interprotomer hydrogen bonds with the backbone oxygen atoms of Cys99 and Pro108.](image-url)
The similarities and differences in the properties of Miner1 and mitoNEET are evident when the two structures are superimposed (Fig. 6). Minera1 is the second known member of the family established by the determination of the mitoNEET structure (Fig. 6).6,10,11 The cluster-binding domains of Miner1 and mitoNEET share the same Cys 3-His 2Fe-2S coordination and are structurally the same within uncertainty (rmsd ∼0.3; PDB ID 2QH7).6 However, superposition of the dimeric structures reveals significant differences in the outermost strands. The backbone rmsd of the β-cap domain is ∼1.1 Å, ~4-fold greater than that of the cluster binding domain (Fig. 6a). These differences result in a more open V shape appearance in the β-cap of Miner1. As an additional test of the significance of this apparent difference, we superimposed the protomer structures of Miner1 and mitoNEET (Fig. 7a). Strikingly, the central strand along with the cluster binding domain of the protomers are superimposable. However, the outer strands (circled) are shifted with respect to one another. Interestingly, Miner1 has an amino acid insertion at the top of the β-cap domain (Thr 94; Fig. 7, highlighted) that appears to act as a spacer extending the distance between the central and the outer strand. In addition, several key variations in aromatic residues (Phe60/Ile86 and Phe82/Ala109) create differences in side chain orientation and appear to affect the position of Tyr71/Tyr98, with a relative displacement of 2.2 Å.

Discussion

We report here the crystal structure determined to 2.1 Å resolution of the ER protein Miner1. We show that, contrary to its original annotation, it contains redox-active 2Fe-2S clusters with an $E_m \sim$ 0 mV at pH 7.5. Miner1 is a member of the novel 2Fe-2S CDGSH protein family and the first FeS protein localized to the ER. Mis-splicing of the mRNA that codifies this protein is causative in the genetic disease Wolfram Syndrome 2 (WFS2). FeS proteins participate in many types of biochemical reactions, the most predominantly characterized are oxidation-reduction reactions, but they also participate in Fe-S cluster assembly and gene expression regulation.16–19 Although not yet determined in vivo, Miner1 has biophysical properties necessary to participate in the functions that have a vital role in homeostasis required for proper health and longevity. The importance of Miner1 in cellular health is evidenced in the deleterious effects of its absence in WFS2 and knockout mice.1,2 The potential role this protein plays in human health and defense against environmental stresses is suggested by the fact that patients with WFS2 can live to 15 years before becoming symptomatic.

Miner1 is the first 2Fe-2S protein found in the ER

Miner1 was identified as one of a small family of proteins that share the CDGSH sequence domain.4 On the basis of strong perinuclear staining and its localization to a lacy network in V5 epitope studies performed in transiently expressed COS-7 cells, Wiley et al. suggested that Miner1 localizes to the ER.7 Amr et al. subsequently confirmed this suggestion in mouse P19 cells in which N-FLAG tagged Miner1 colocalized with calnexin, a known ER marker.3 Taken together with our structural studies (Fig. 3), Miner1 is to our knowledge the first 2Fe-2S-containing protein localized to the ER membrane.

![Fig. 5. Redox potential and stability of the 2Fe-2S clusters of Miner1 and mitoNEET. (a) pH-dependence of the redox potentials ($E_m$) of Miner1 and mitoNEET. Both display pH-dependent $E_m$ values from pH 7.5 to 10.0 that decrease by 50 mV per pH unit, indicating that the reduction is proton-coupled. (b) pH-dependence of the 2Fe-2S cluster stabilities of Miner1 and mitoNEET. The half-life for Miner1 is orders of magnitude less pH-dependent varying by only ~5-fold from pH 5.3 to 7.1 compared to the ~300-fold change for mitoNEET.](image-url)
Proton-coupled redox activity of Miner1

One of the best characterized functions of FeS proteins is their role in redox reactions. A primary property that relates to this function is the redox potential $E_m$. The $E_m$ of Miner1 and mitoNEET were measured via equilibrium titration to be essentially the same at $\sim 0$ mV at pH 7.5. The similarity of their $E_m$ values is expected as both Miner1 and mitoNEET have nearly superimposable structures of the cluster binding domains that include the rare Cys3-His coordination. The $E_m$ values vary by $\sim 50$ mV/pH unit indicating that the reductions are proton-coupled (Fig. 5a). Since both Miner1 and mitoNEET display the same behavior, the site of protonation is also likely the same. Given the magnitude of the change in redox potential ($>150$ meV), the protonation site must be located near the 2Fe-2S cluster. In addition to the inorganic sulfide of the cluster, the 2Fe-2S clusters share ligation by Cys110 and His114, and interact with the nearby Asp111 and Ser104. Further localization of the proton requires additional investigation.

The most common redox-active motifs for 2Fe-2S proteins are ferredoxin-like, whose clusters are coordinated by four Cys, and Rieske-like, whose clusters are coordinated by two Cys and two His. The Fd-like proteins tend to have negative $E_m$ ($\sim -300$ mV) and the Rieske-like positive $E_m$ ($\sim +300$ mV), although some can be in the same range as Miner1.17 Miner1 and mitoNEET have redox potentials similar to those of known electron transfer species such as cytochrome $b$ and quinones.20,21 Thus, their potentials are within the physiological range consistent with Miner1 and mitoNEET participating in electron transfer or oxidation/reduction reactions in the cell.

Different pH sensitivity of the 2Fe-2S clusters of Miner1 and mitoNEET

Another known functional role for FeS proteins is cluster assembly and transfer.19 The Miner1 family shares the unusual Cys3-His coordination with only one other known protein structure—that of an IscU D38A mutant from a hyperthermophilic bacteria,22 a protein that participates in cluster assembly. In...
Fig. 7. Protomer superposition of Miner1 and mitoNEET highlighting distinct backbone and side chain conformations. (a) The backbone superposition of the Miner1 protomer A on the mitoNEET protomer A. Specific side chains showing different conformations are indicated, and the region around them is expanded in b. The 2Fe-2S centers are essentially identical in each superposition and are shown as lines. The backbones of the two proteins are nearly indistinguishable in the lower cluster binding domain. Difference in the backbone are encircled. (b) The largest difference in the backbone occurs in the vicinity of an additional amino acid in Miner1, Thr94 (blue box). Side chain differences in the β-cap domain occur near Ile86 and Ala109 of Miner1, which correspond to Phe60 and Phe82 of mitoNEET. Tyr98 is located in the vicinity and is displaced from the corresponding Tyr71 of mitoNEET (red box). (c) Sequence alignment of Miner1 and mitoNEET. Changes to side chains shown in b are boxed in blue and red. The 16 amino acid cluster binding motif is boxed in orange. Secondary structures are indicated above the sequences by a thick arrow for a β strand and by a cylinder for an α helix.
order for this function to be plausible for Miner1, the Fe-S cluster must be easily transferable or labile. The measured lifetime of the 2Fe-2S clusters of Miner1 was ∼10-fold shorter at pH 7.1 than that of mitoNEET. This lability is accelerated by orders of magnitude in cellular extracts (data not shown), suggesting a possible role of Miner1 in FeS cluster assembly and/or transfer.

Surprisingly, given the highly similar structure of Miner1 and mitoNEET, the lability of the two proteins displayed drastically different pH dependence. Whereas the lability of Miner1 varied by only ∼5-fold from pH 5.3 to 7.1, mitoNEET varied ∼300-fold over the same pH range (Fig. 5). This difference in the lability is likely due to differences in the charged or titratable groups of the proteins. A site-specific comparison revealed seven differences between the proteins (Fig. 6) — Lys74, Asn84, Glu85, Asn87, Leu93, Thr106, and Leu120. One or more of these changes is likely responsible for the altered pH sensitivity.

Miner1, ER function and health and disease

Miner1 is targeted to the ER membrane, the organelle in which lipid synthesis, protein folding and protein maturation take place. A number of post-translational modifications are performed here, including lipiddation, hydroxylation, glycosylation, methionine and cysteine oxidation. The ER participates in quality control by facilitating the recognition and targeting of aberrant proteins for degradation. When its capacity is exceeded, a signaling network called the unfolded protein response is activated. The redox ability of Miner1 could allow it to participate in protein modification and possible ER stress responses. Recent studies on the tethering of the ER and mitochondria via the protein mitofusin indicate that a direct interaction between the organelles is essential for Ca2+ transfer and cellular function. Increasing evidence also links altered communication between the ER and mitochondria to apoptosis and disease. Mitochondrial–ER crosstalk is implicated in type 2 diabetes. Interestingly, Miner1 is situated on the surface of the ER and mitoNEET is on the surface of the OMM. The similarity of their Em values indicates that they would respond to similar redox stimuli and could possibly communicate through electron transfer, thus providing a means of ER–mitochondrial crosstalk.

Proper biological function of Miner1 is necessary for cellular health, as the absence of Miner1 has deleterious effects, such as WFS2 or decreased longevity (Fig. 1). The similarities of Miner1 with its paralog mitoNEET also strengthen the link between proper function of this novel family of 2Fe-2S proteins and general health. Thus, further investigation of the interplay between the functional and biophysical properties of this family is anticipated to advance our understanding of these complex mechanisms involved in health and metabolic disease.

Materials and Methods

Expression and purification

The soluble domain of miner1 (residues 57–135; cDNA purchased from Open Biosystems) was amplified by PCR and subcloned into the bacterial expression vector pET28a (+) (Novagen) containing an N-terminal, thrombin cleavable His tag. The C92S mutation was made by site-directed mutagenesis using PCR. This plasmid was transformed into BL21-RIL(DE3) (Stratagene) and grown as described. Harvested cells were suspended in binding buffer (20 mM Tris–HCl pH 7.9, 5 mM imidazole, 500 mM NaCl). After centrifugation, the lysate was added to Ni-NTA resin, which was washed with 10 column volumes of wash buffer (20 mM Tris–HCl pH 7.9, 30 mM imidazole, 500 mM NaCl), equilibrated in thrombin cleavage buffer (25 mM Tris–HCl pH 8.0, 100 mM NaCl, 2.5 mM CaCl2), and cleaved upon addition of thrombin at room temperature for 16–24 h. Miner1 was eluted with wash buffer and further purified by size-exclusion chromatography (S-200 GE Healthcare) in 25 mM Tris–HCl pH 8.0, 100 mM NaCl. For crystallographic experiments, the protein was further purified by cation-exchange chromatography (HiTrap, GE Healthcare) as described.

Optical spectroscopy and stability measurements

All UV-visible absorption spectra were measured from the near-UV to the near-IR (250 – 700 nm) at 35 °C with a Cary50 spectrometer (Varian Inc, Palo Alto CA) (10–20 μM protein in 25 mM Tris–HCl pH 8.0, 100 mM NaCl).

The stabilities of the 2Fe-2S clusters were determined from monitoring the characteristic absorbance at 460 nm as a function of time following dilution of the protein into the assay buffer (∼20 μM protein in 100 mM buffer at 35 °C) (Supplementary Data Fig. S3). The following buffers were used at the pH indicated: Tris–HCl pH 7.1 and 6.8, and bis-Tris–HCl at pH 6.3, 5.8 and 5.3. Both the buffer and the protein were filtered before the start of the measurement and a layer of paraffin oil was placed over the solution to minimize evaporation. The half-life corresponding to 50% decay was determined at each pH value.
midpoint/redox potential ($E_m$) from a fit of the fraction oxidized to the Nernst equation:

$$A = A_{ox} \left[ 1 + 10^{\left( (E_m - E) / (n \cdot 59.1 \text{mV}) \right)} \right] + A_{red}$$

where $A_{ox}$ is the absorbance of the fully oxidized sample, $E_m$ is the midpoint potential in mV, $E$ is the measured cell potential in mV, $n$ is the number of electrons being transferred in the redox reaction, and $A_{red}$ is the lowest absorbance level corresponding to the fully reduced Fe-2S center in either native or mutant mitoNEET. Potentials were adjusted to SHE for presentation. Samples were tested for their ability to re-oxidize upon exposure to ambient oxygen.

**Crystallization**

Crystallization was achieved as described for mitoNEET. The final conditions were 100 mM Tris–HCl pH 8.0, 100 mM NaCl, and 15–20% (w/v) PEG 3000 in the protein well, equilibrating against 100 mM Tris–HCl pH 8.0 and 30–35% PEG 3000 in the reservoir. Crystals were frozen at 77 K after soaking for 1 min in 100 mM Tris–HCl pH 8.0, 40% PEG3000 and sent frozen (77 K) to SSRL in an SSRL-supplied cassette system for X-ray data collection and analysis.

**X-ray data collection and structural determination**

Frozen crystals were screened using the Stanford Automated Mounter operated by Blu-ice. The data were recorded using a Rigaku R3B CCD detector at BL9-2 and an ADSC Q315R CCD detector at BL7-1. Data were processed with XDS. The structure of Miner1 was determined by MAD phasing. Data reduction and primary phasing at a resolution of 2.1 Å were accomplished as described.

**Protein Data Bank accession numbers**

The atomic coordinates have been deposited in the Protein Data Bank: PDB ID code 3FNV; PDB ID code 3FNV.

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**Supplementary Data**

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.jmb.2009.06.079

**References**


Note added in proof: Since the work was submitted for publication, a Taiwanese consortium published an extensive paper on the characteristics of cisd2 knockout mice (Chen, Y. F., Kao, C. H., Chen, Y. T., Wang, C. H., Wu, C. Y., Tsai, C. Y. et al. (2009). Genes Dev. 23, 1183–1194). Furthermore, they showed that the knockout mice had a reduced electron transport activity of complex I-III, complex II-III and complex IV. They localized Miner1 to both the OMM and the ER with a 6:1 ratio. In our work, we present the structure of the protein that is key to their model for longevity and WFS2.