

ARTICLE

Received 8 Jun 2010 | Accepted 22 Sep 2010 | Published 19 Oct 2010

DOI:10.1038/ncomms1097

Structural bases for the interaction of frataxin with the central components of iron–sulphur cluster assembly

Filippo Prischi¹, Petr V. Konarev², Clara Iannuzzi¹, Chiara Pastore¹, Salvatore Adinolfi¹, Stephen R. Martin¹, Dmitri I. Svergun² & Annalisa Pastore¹

Reduced levels of frataxin, an essential protein of as yet unknown function, are responsible for causing the neurodegenerative pathology Friedreich's ataxia. Independent reports have linked frataxin to iron–sulphur cluster assembly through interactions with the two central components of this machinery: desulphurase Nfs1/IscS and the scaffold protein Isu/IscU. In this study, we use a combination of biophysical methods to define the structural bases of the interaction of CyaY (the bacterial orthologue of frataxin) with the IscS/IscU complex. We show that CyaY binds IscS as a monomer in a pocket between the active site and the IscS dimer interface. Recognition does not require iron and occurs through electrostatic interactions of complementary charged residues. Mutations at the complex interface affect the rates of enzymatic cluster formation. CyaY binding strengthens the affinity of the IscS/IscU complex. Our data suggest a new paradigm for understanding the role of frataxin as a regulator of IscS functions.

¹ National Institute for Medical Research, The Ridgeway, London NW7 1AA, UK. ² European Molecular Biology Laboratory, EMBL c/o DESY, Notkestrasse 85, Hamburg D-22603, Germany. Correspondence and requests for materials should be addressed to A.P. (email: apastor@nimr.mrc.ac.uk).

Frataxin is an essential iron-binding protein that is highly conserved in most organisms, from bacteria to humans¹. Reduced expression levels of this protein are sufficient to induce Friedreich's ataxia, a relentless and currently incurable neurodegenerative disease. Absence of frataxin in knockout mice is also embryonic lethal². Frataxin has been implicated in one of the most important iron-related metabolic pathways through the observation of biochemical and genetic interactions with the highly conserved Nfs1/Isu complex (which corresponds to the IscS and IscU proteins in prokaryotes): Nfs1 and Isu, the two main components central to the iron–sulphur (Fe–S) cluster assembly machinery^{3–5}, are, respectively, a cysteine desulphurase enzyme (EC 2.8.1.7), which catalyses the reaction that converts l-cysteine into alanine and a highly reactive sulphide, and the scaffold protein on which the cluster is transiently formed⁶. In addition to having a role in Fe–S cluster assembly, IscS is also involved in other sulphur-related pathways, which include biotin synthesis, tRNA modifications and molybdopterin biosynthesis through interaction with proteins TusA, Thil and MoeB/MoaD^{7–9}.

Although evidence of a direct interaction between frataxin and the Nfs1/Isu complex has received independent confirmation^{3–5}, many questions, the answers to which would be crucial for understanding frataxin function, remain unanswered. For instance, it is unclear as to which of the IscS/IscU components frataxin binds. Calorimetric studies on human frataxin have suggested a direct 1:1 iron-mediated interaction with Isu1⁴, with nanomolar dissociation constants¹⁰. In yeast, frataxin (Yfh1) binds to Isu1, but the interaction seems to require Yfh1 oligomerization¹¹. In contrast, no direct interaction with isolated IscU was observed for prokaryotic frataxin (CyaY), which interacts instead with IscS^{12,13}. Another unanswered question is with regard to the role of a highly conserved tryptophan exposed on the frataxin surface^{14–16}. Tryptophans are relatively low-occurrence amino acids and their conservation on exposed surfaces strongly suggests a role in molecular recognition¹⁷. Finally, an even more crucial question is why these interactions take place and, ultimately, what is the primary function of frataxin. This point remains highly controversial.

Frataxin has been suggested to function as a chaperone that delivers iron to the IscS/IscU complex⁴, or as a scavenger¹⁸, which, through formation of large iron-loaded assemblies, prevents iron excesses in mitochondria while still keeping iron biologically available for being delivered to key acceptors. More recently, we have shown that, although not inhibiting desulphurase enzymatic activity, CyaY is able to slow down the speed of Fe–S cluster formation and have proposed a role for frataxin as the gatekeeper of the cluster assembly machinery¹³.

Probably the most effective way to test these hypotheses and resolve the debate is to study the frataxin function *in vitro*, which makes it possible to delimit the system to a minimal and well-controlled number of components. This strategy should allow us to formulate working hypotheses, which could then be tested *in vivo*. A similar approach has already proven successful, for instance, for the identification of functionally important residues on the frataxin surface¹⁹.

Here, we have dissected the interactions of CyaY, IscS and IscU by gaining information on the structure of their binary and ternary complexes using small-angle X-ray scattering (SAXS), a technique able to provide the overall shape of even large molecular assemblies. We have validated the surfaces of interaction by mapping them directly by nuclear magnetic resonance (NMR) chemical shift perturbation techniques and mutagenesis studies. Finally, we have proven the role of specific residues of either CyaY or IscS on the enzymatic activity of IscS and showed how the presence of CyaY modulates the binding affinities of the complex. As the residues involved in the interaction are conserved, we expect this function to be shared by all frataxin orthologues. The picture suggested by our results is that frataxin is a key element of a complex

network of interactions centred on the desulphurase. The interplay between competing interactions may be crucial for regulating cluster formation and selecting alternative metabolic pathways.

Results

CyaY forms a 1:1 complex with IscS. To quantify the nature of the interaction between *Escherichia coli* CyaY, IscS and IscU, we measured the stability and stoichiometries of the binary IscS/IscU and IscS/CyaY complexes. We used isothermal calorimetry (ITC), a technique that allows determination of binding constants with high accuracy. Titration of IscS with IscU results in an endothermic binding, which can be fit with a 1:1 binding model and provides a K_d of $1.3 \pm 0.2 \mu\text{M}$ (Fig. 1a). These results are in agreement with previous data²⁰ and with the recent structure of the *E. coli* IscU/IscS complex, which shows interaction of the IscS dimer with two IscU monomers⁹. Titration of IscS with CyaY results in an exothermic reaction, which can also be fit with a 1:1 binding model, leading to a K_d of $18.5 \pm 2.4 \mu\text{M}$ (Fig. 1b). A lower affinity of IscS for CyaY is in agreement with the evidence that (as opposed to IscU/IscS) the CyaY/IscS complex cannot be easily isolated by gel filtration²¹. As previously described, we could not observe any direct binding between IscU and CyaY either, in the presence of iron, both in aerobic and anaerobic conditions^{12,13}.

These data provide the first quantification of the affinities of the CyaY/IscS complex and direct evidence that the interaction also occurs without the need of large CyaY assemblies.

SAXS reveals the size and shape of the complexes. To obtain a molecular description of the IscS/CyaY and IscS/IscU/CyaY complexes, we tried to crystallize the binary complexes. We obtained good-quality crystals under several different conditions, but they invariably contained only IscS²¹. We resorted to a different strategy based on SAXS measurements and molecular modelling validated by experimental data. We acquired and compared SAXS data for each of the isolated individual components, as well as for the binary and ternary complexes (IscS/CyaY, IscS/IscU and IscS/CyaY/IscU). The estimated apparent molecular mass (MM_{exp}) and hydrated particle volume (V_p) for IscS agreed with the presence of a dimer in solution (Fig. 2 and Supplementary Table S1), as expected for the constitutive IscS homodimer, which has a P2 symmetry (>90 kDa)²². Both IscU and CyaY are monomeric in isolation, in agreement with previous characterizations^{23–25}. The MM_{exp} and V_p of the complexes are distinctly different from those of the isolated species and correspond to stoichiometries of 1:1:1 for the ternary complex and of 1:1 for both binary complexes. The envelopes of the complexes are clearly different from those of the isolated species, allowing us to identify the excess volumes in which monomeric IscU, CyaY or both could fit (Fig. 3). In agreement with the recently published crystallographic structure, we observed a more elongated shape for the IscS/IscU complex⁹. The shape of the binary complex IscS/CyaY is more globular, thus suggesting that CyaY is located close to the interface between the two IscS protomers. No changes were observed when iron (Fe^{2+}) was aerobically added to the sample to reach a molar ratio of 1:1:2 CyaY/IscS/ Fe^{2+} (data not shown).

Multiple runs of the program DAMMIF²⁶, a fast version of DAMMIN²⁷, were used to produce average low-resolution *ab initio* shapes for all the constructs. The shape of isolated IscS matches well with the crystal structure (1P3W)²² (Fig. 4a). Among the several NMR and X-ray structures of IscU, the best fit was observed with 1Q48 (ref. 28) (Fig. 4b), which corresponds to a highly flexible structure in which the N- and C-termini are unfolded. More compact structures such as 2Z7E yielded significantly worse fits (Fig. 4c). This is well in agreement with the observation that the N terminus of *E. coli* IscU is unstructured in solution^{21,29}. CyaY could be fit with the NMR structure of monomeric CyaY (ISOY)²⁵ (Fig. 4d).

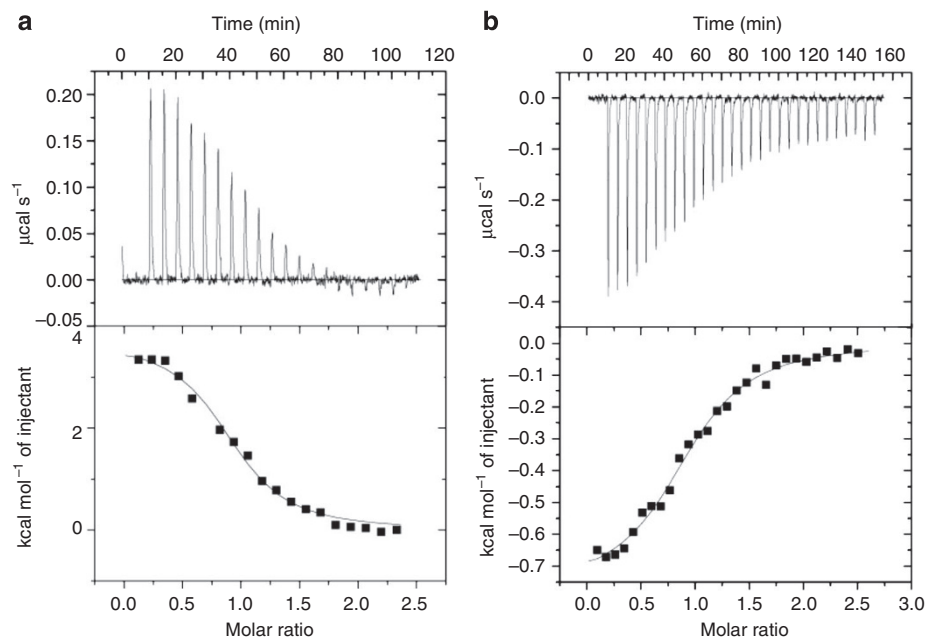


Figure 1 | Thermodynamic stability of the CyaY/IscS and IscU/IscS complexes. (a) Binding isotherm for the interaction of IscU with IscS. In the upper panel, peaks indicate the heat released after each addition of IscU into the protein solution. In the lower panel, the data were best fit using a single binding constant to calculate the thermodynamic parameters (continuous line). (b) Binding isotherm for the interaction of CyaY with IscS. In the upper panel, peaks indicate the heat released after each addition of CyaY into the protein solution. In the lower panel, the data were fit using a single binding constant.

For all the complexes, rigid body modelling was performed to fit the experimental SAXS data using the high-resolution structures of IscS, IscU and CyaY. The obtained rigid body models displayed good agreement with the *ab initio* shapes (Fig. 4e–g). For the IscS/IscU binary complex, we obtained a better fit with the structure (3LVL)⁹ than when using the individual models, thus reinforcing the observation that IscU undergoes a structural transition on binding (Fig. 4e). The binary and ternary complexes of CyaY reveal binding of this molecule at a site near the IscS dimer interface, as opposed to IscU, which is positioned towards the periphery of the IscS dimer (Fig. 4f,g). Accordingly, the radius of gyration R_g and the maximum size of the particle D_{max} for the IscS/IscU complex are significantly larger than for isolated IscS, whereas for IscS/CyaY complex there is only a small increase in these parameters (Supplementary Table S1). The rigid body modelling yielded excellent fits to the scattering data with discrepancies (χ -values) of < 1.45 . The fits for the IscS/CyaY and IscS/CyaY/IscU models displayed minor deviations at higher angles (beyond s -values of 0.15 \AA^{-1}). These deviations can be explained by the presence of small populations of free components in solution, in agreement with the lower affinity of the IscS/CyaY complex as compared with the IscS/IscU one (the difference is of approximately one order of magnitude).

Overall, the *ab initio* and rigid body models from SAXS are consistent with each other and give us the first direct picture of the structure of the IscS/IscU/CyaY complexes.

Model validation and collection of further restraints. To validate the SAXS models and obtain more restraints for further modelling, we resorted to NMR techniques that, by detecting spectral perturbation, are a sensitive tool to identify even residual binding and map the information to specific residues. We used ^2H , ^{15}N double-labelled CyaY and titrated it with IscS up to a 1:1 molar ratio. Per-deuteration was found to be necessary to decrease spin diffusion, resulting in much sharper spectra and therefore in a higher resolution. The residues of CyaY most affected by addition of IscS were Trp14, Leu15, Glu19, Asp22, Asp23, Trp24, Asp25, Asp27, Ser28, Asp29, Ile30, Asp31, Cys32, Glu33, Ile34, Leu39, Thr42, Phe43,

Glu44 and Gly46 (Fig. 5a). These residues map spatially close in the region of CyaY comprising $\alpha 1$, $\beta 1$ and the $\alpha 1\beta 1$ loop (blue side chains in Fig. 5b). We then added IscU to the solution up to a 1:1:1 molar ratio (CyaY/IscS/IscU) and observed new spectral perturbations (Fig. 5c). Some of the residues already perturbed by IscS, that is, 14, 19, 22–24, 27–31, 33–34, 39 and 43–44, were further affected by the addition of IscU. In addition, other residues, namely, Thr40, Ile41, Lys48, Ile50, Asp52, Arg53, Glu55, Trp61, Leu62, Ala63, Thr64, Gln66, Gly68, Tyr69 and His70, were either shifted and/or broadened. They mostly map to the β -sheet surface (red side chains in Fig. 4b). In particular, both the backbone and the indole amides of Trp61 were affected. The additional perturbation must be caused either by a direct contact of these residues with IscU or by a conformational change induced by the addition of this component.

Validation of the surface of interaction on IscS, as suggested by SAXS, came from the design of IscS mutants, chosen to target residues that could potentially affect interaction with CyaY (Fig. 5d). A IscS_R220E/R223E/R225E triple mutant was designed to invert the charge of a positively charged patch formed by residues 55, 67, 219–220, 223, 225 and 237 (all arginines) close to the dimer interface (Fig. 5e). These mutations affect solvent exposed residues. Therefore they do not alter the stability of the protein but are expected to abolish binding to the complementary charged surface of CyaY. IscS_I314E/M315E, IscS_K101E/K105E, IscS_E334S/R340S and IscS_R39E/W45E mutants were also produced as controls. Because affecting residues far from the CyaY/IscS interface, these mutations introduce charge inversion and/or perturb the surface but leave invariant the interaction.

The effects of these mutants on the NMR spectrum of CyaY were tested and compared with that of wild-type IscS. The only mutant that caused no perturbation of the CyaY spectrum is IscS_R220E/R223E/R225E (data not shown), thus indicating that these mutations are sufficient to abolish the interaction completely. The other mutants produced the same perturbation as the CyaY spectrum, as observed for wild-type IscS. Following the same strategy, the mutants were tested for IscU binding as well, and yielded results fully consistent with the structure of the

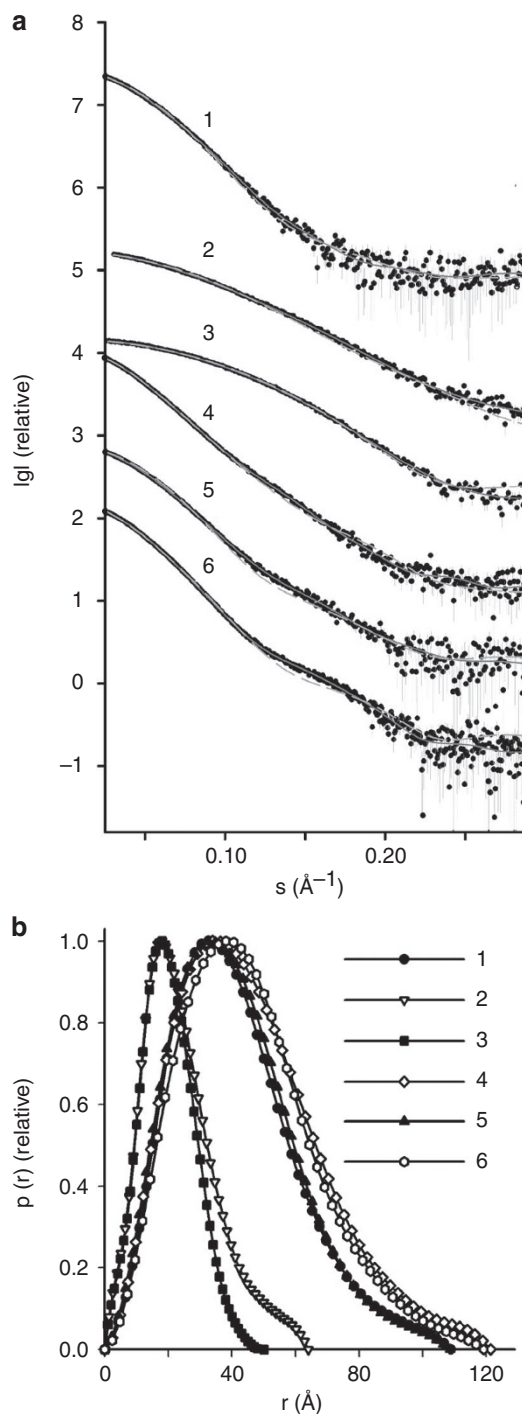


Figure 2 | Small-angle X-ray scattering data and fits. (a) The X-ray scattering patterns from IscS, IscU, CyaY binary complexes IscS/IscU and IscS/CyaY and ternary complex IscS/CyaY/IscU. The experimental data are displayed as dots with error bars, the scattering from typical *ab initio* models computed by DAMMIF as full lines and the calculated curves from the high-resolution (for proteins alone) and rigid body models (for complexes) computed by CRYSOLO/SASREF as dashed lines. Plots display the logarithm of the scattering intensity as a function of momentum transfer $s = 4\pi \sin\theta/\lambda$, where 2θ is the scattering angle and λ is the X-ray wavelength. The successive curves are displayed down by one logarithmic unit for clarity. (b) Distance distribution functions from IscS, IscU, CyaY, binary complexes IscS/IscU and IscS/CyaY and ternary complex IscS/CyaY/IscU (labelled 1–6, respectively).

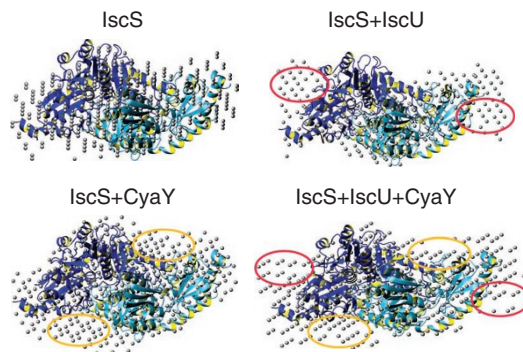


Figure 3 | Small-angle X-ray scattering shapes. Comparison of the densities superposed to the crystal structures of IscS (1P3W) for IscS alone, IscS/IscU, CyaY/IscS and CyaY/IscS/IscU. Regions with additional density in the binary and ternary complexes are highlighted in red or yellow ovals.

IscS/IscU complex⁹, thus providing an independent validation of our procedure (Supplementary Discussion).

Taken together, these results lend further support to the SAXS models, positioning CyaY in a site well distinct from that occupied by IscU.

The interaction between IscU and CyaY is mediated by IscS.

A hybrid approach combining SAXS-based rigid body modelling (SASREF³⁰) with the docking software HADDOCK³¹ was used to build the refined models of the complexes. Whereas the nominal resolution of the SAXS-based models is defined by the maximum momentum transfer in the fitting range of Figure 2 ($2\pi/s_{\max} \sim 10\text{--}20\text{\AA}$), the use of experimental docking restraints and high-resolution structures in hybrid modelling is expected to significantly push the resolution limit for the consensus model, thus allowing for meaningful analysis of protein–protein interactions in the complex.

The consensus model reflects most of the experimental observations and suggests several novel observations. Although occupying distinct surfaces of IscS, CyaY and IscU are next to each other, with CyaY fitting the cavity delimited by the IscS dimer interface and the IscU-binding site (Fig. 6a). This region overlaps only partially with that occupied by TusA, another protein partner of IscS involved in delivering sulphur for tRNA modification⁷ (Fig. 6b). Compared with the position of TusA⁹, CyaY is inserted less deeply into the cleft formed by IscU and the IscS dimer interface, as directly supported by the fact that IscS_R39E_W45E retains binding to CyaY, whereas these two residues were shown to be directly involved in the interaction with TusA (Supplementary Table S2). On the other hand, mutation of R220 and the surrounding positively charged residues of IscS abolishes CyaY but not TusA binding.

CyaY sits relatively close to the loop containing the catalytic Cys328, but far from the pyridoxal phosphate ($\sim 18\text{\AA}$), in agreement with our previous observation that CyaY does not interfere with enzymatic activity¹³ (Fig. 6c). Conversely, although we do not know its exact position, as it is not visible in the crystal structures, the catalytic loop is, in principle, long enough to protrude and contact the nearby CyaY. The experimental restraints that position the region 18–36 of CyaY to pack against the region of IscS containing Arg220, Arg223 and Arg225 automatically also impose the surface of the β -sheet of CyaY to be spatially close to IscU. Although insufficient to stabilize a stand-alone CyaY/IscU complex, the surface of interaction between these two proteins may stabilize the ternary complex. Among the residues of CyaY that are spatially proximal to IscU is the exposed Trp61, which is close to, albeit not necessarily in contact with, the conserved cysteines of IscU.

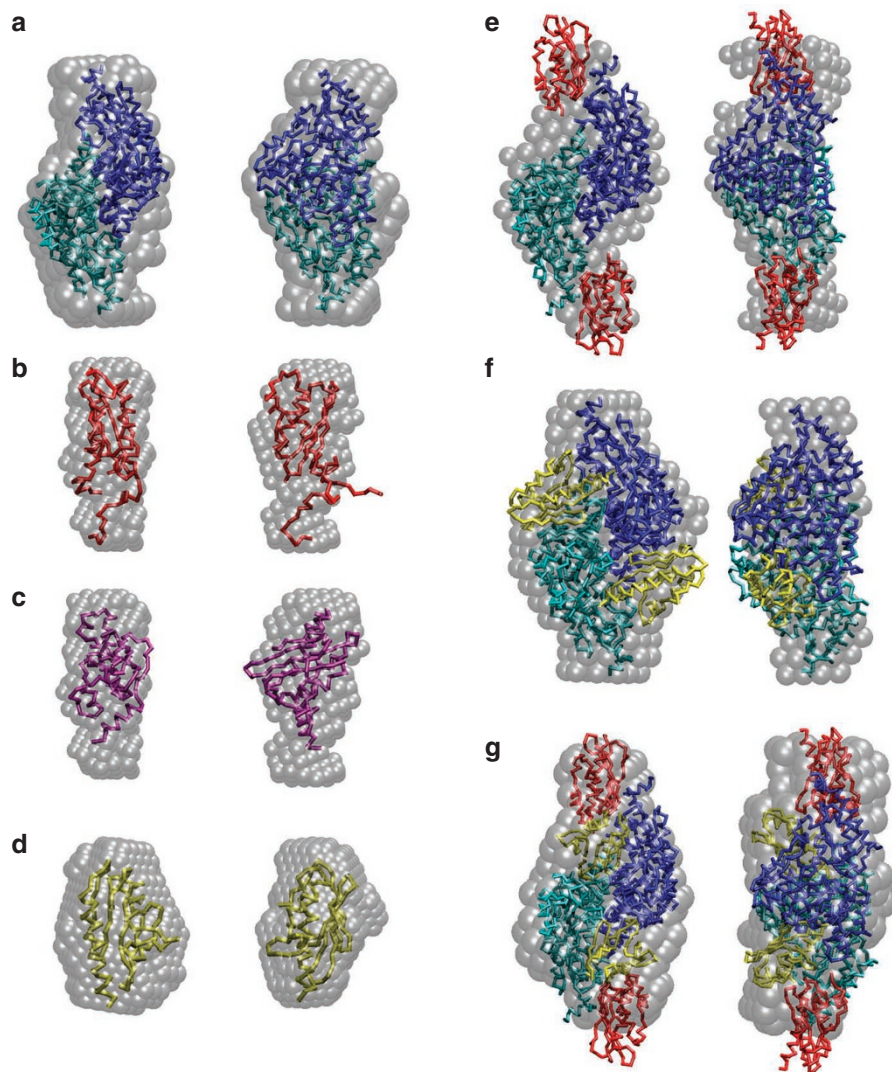


Figure 4 | Molecular modelling based on small-angle X-ray scattering. *Ab initio* bead models of IscS (a), the solution (b) and X-ray (c) structures of IscU, CyaY (d), IscU/IscS (e), CyaY/IscS (f) and CyaY/IscS/IscU (g) complexes. The models were obtained from DAMMIF (grey semitransparent spheres) superimposed with rigid body models by the software SASREF. The two components of the crystallographic dimer model of IscS are displayed as cyan and blue C_α traces. For each model, the right views are rotated counterclockwise around the vertical y axis.

CyaY influences the kinetics of enzymatic Fe–S cluster assembly.

We have previously shown that the presence of CyaY slows down the enzymatic rates of Fe–S cluster formation, suggesting a role of frataxin as an IscS inhibitor¹³. To investigate whether the interaction of CyaY with the IscS/IscU complex has a direct influence on these inhibitory properties, we tested the effect of replacing wild-type IscS with its mutants on the rates of cluster formation. We followed this process by analysing, under strict anaerobic conditions, the absorption spectrum at 456 nm, a wavelength typical of the cluster²⁰. For comparison, we also tested two CyaY mutants, CyaY_E18K/E19K/E22K and CyaY_W61R. The first should affect the surface of interaction with IscS and therefore have complementary effects to those of IscS_R220E/R223E/R225E. The second should influence the surface of interaction with IscU.

The presence of CyaY has a strong inhibitory effect as compared with the assay carried out in its absence, and appreciably reduces the initial reaction rate (Fig. 7a, a red curve as compared with a green curve). The IscS_R220E/R223E/R225E and CyaY_E18K/E19K/E22K mutants behave as the control experiment in the absence of CyaY (blue and magenta curves), as expected from abolishing the interaction between CyaY and the enzyme. CyaY_W61R has

an intermediate behaviour, suggesting that this mutation causes a lower degree of disruption of the complex. IscS_I314E/M315E, which weakens without abolishing the interaction between IscS and IscU, only results in a reduction of the kinetic rates (cyan curve). The other IscS mutants that should have no effect on the complex behave similar to the wild-type IscS (data not shown).

Taken together, these results provide a strong and convincing link between the interaction of the three components and the enzyme activity.

CyaY enhances the affinity of IscU on IscS. Finally, we wondered whether binding of one component on IscS has an influence on the affinities of the other interactions and whether the presence of the cluster on IscU could decrease the affinity of the IscU/IscS complex, thus suggesting how Fe–S cluster formation could be regulated. We measured the dissociation constants by Biolayer interferometry (BLI), a novel technique that measures molecular interactions and also provides information on the on/off rates. When we immobilized CyaY on the surface and measured the affinities of the CyaY/IscS complex in the absence of IscU, we obtained a K_d of $23 \pm 3 \mu\text{M}$ (Fig. 7b), whereas when IscU was

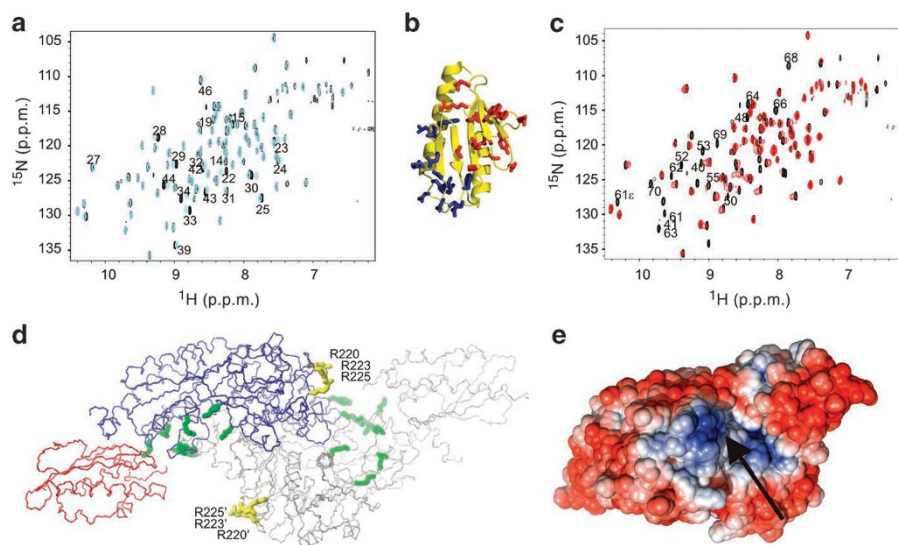


Figure 5 | Experimental validation of the models. (a) Superposition of the hetero single quantum coherence spectra of ^{15}N uniformly labelled CyaY (light blue) with the final point of a titration with unlabelled IscS (1:1 molar ratio, black). (b) Residues of CyaY (1SOY) involved in the interaction with IscS (blue) and IscU (red). (c) Superposition of the hetero single quantum coherence spectra of the CyaY/IscS complex (black) with the final point of titration with unlabelled IscU (up to 1:1:1 molar ratios, red). (d) Mapping of IscS mutations into the IscS/IscU crystal structure (3LVL). To highlight the symmetry relation, the backbone of one of the IscS protomers is shown in blue with the complexed IscU shown in red, whereas the backbones of the corresponding symmetry related to IscS/IscU are shown in grey. A linear fog effect was applied. The side chains of mutated residues, which have no effect on the interaction with CyaY, are shown in green, those that abolish interaction are shown in yellow. The dimer interface and the presence of IscU delimit a small cavity in which CyaY is accommodated. (e) Electrostatic surface of unbound IscS. An arrow indicates the position of the positively charged patch involved in binding. The patch comprises residues arginines 55, 67, 219–220, 223, 225 and 237. The residues 220, 223 and 225 are those that, if mutated to glutamates, abolish CyaY binding. IscS is oriented as in **d** and rotated by 90° around the horizontal x axis.

present, the K_d is 35 ± 6 nM (Fig. 7c). Vice versa, by immobilizing IscS on the surface, the affinity value of the IscU/IscS interaction in the absence of CyaY is 1.5 ± 0.3 μM . When the experiment was repeated in the presence of CyaY, we measured a K_d of 400 ± 30 nM, which is appreciably lower than the value observed in the binary complex. In all cases, we observed that the dissociation rates (k_{off}) of the ternary complex were lower than those observed for the binary complexes (for example, the k_{off} for the disassembly of the IscS/IscU complex is 0.8 s^{-1} in the absence of CyaY, versus 0.006 s^{-1} in the presence of CyaY). The values obtained by BLI are in excellent agreement with ITC results, which also provide a K_d for the IscU/IscS interaction in the presence of CyaY that is approximately one order of magnitude tighter than that observed in the absence of CyaY (250 ± 18 nM data not shown).

We then tried to measure the affinities of the IscU/IscS interaction for cluster-free and cluster-loaded IscU (in the absence of CyaY) to check whether the presence of the cluster is able to modulate the interaction with IscS. Unfortunately, the cluster is too unstable when bound to wild-type IscU, especially under non-anaerobic conditions, to allow us to perform interferometry assays. We thus used an IscU_D39A mutant that is known to stabilize the cluster that persists for several hours, even under aerobic conditions^{32,33}. We measured by BLI a K_d of 41.7 ± 4 nM for the cluster-free IscU mutant. The presence of CyaY further increases the affinity yielding to a K_d of 12.00 ± 2 nM. Conversely, we obtained a K_d of 0.86 ± 0.04 μM for the cluster-loaded protein (Fig. 7d).

Taken together, these results indicate that the presence of the third component and/or the iron state of IscU modulate the affinities of the binary IscU/IscS and CyaY/IscS complexes.

Discussion

We have reported a description of the structural determinants of the interaction between bacterial frataxin and IscS and IscU, the two essential components of *isc* operon. We have shown that

IscS binds CyaY with a 1:1 stoichiometry that reflects the twofold symmetry of the dimeric enzyme. We then resorted to SAXS to obtain a more detailed, although low-resolution, description of the complexes. The information was validated by NMR and by judiciously designed mutations and finally translated into a comprehensive three-dimensional model. These results help us to address several open questions.

As the interaction with IscS occurs at the level of CyaY monomers, there is no need for the large assemblies that have been suggested to be involved in the recognition of yeast frataxin (Yfh1) by the corresponding Nfs1/Isu1 complex¹¹. The surfaces of interaction that we have mapped both on desulphurase and CyaY are anyway incompatible with the binding of a large assembly: CyaY is accommodated in a relatively narrow cavity of the IscS dimer, which could not accommodate larger species without a major rearrangement. Even more important is the consideration that the residues involved in the Yfh1 trimer interfaces (the trimer being the smallest building block for larger spherical assemblies) are the same that mediate IscS recognition³⁴.

Our findings also resolve the long-standing debate on whether frataxin interacts with IscS and/or with IscU. A direct interaction between human frataxin and Isu was reported as early as 2003⁴. These results could not, however, be reproduced in pulldowns of the bacterial orthologues, either using overexpressed proteins or endogenous extracts^{12,13}, suggesting the possibility that IscU could be detected solely as a consequence of its interaction with IscS. Direct titrations of the two proteins using biophysical techniques also failed to detect an interaction^{13,21}. We now show that CyaY packs mainly against IscS, which functions as a mould. Contacts with IscU are possible but only in the context of the ternary complex. However, it is possible that the differences observed between prokaryotic and eukaryotic proteins may be explained by the N-terminal extension present only in eukaryotic frataxins. These additional residues, which are a flexible spacer between the mitochondrial localization signal and frataxin, could contribute to

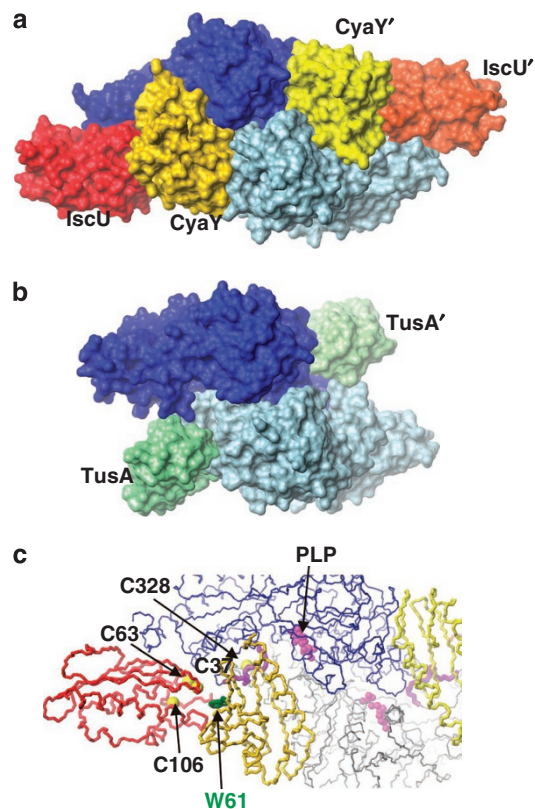


Figure 6 | Structure of the ternary complex CyaY/IscS/IscU. (a) Surface representation of the model obtained by combining the small-angle X-ray scattering and NMR information, showing in blue and cyan the two IscS protomers; in red and orange red, IscU; and in gold and yellow, CyaY. (b) Surface representation in the same orientation as in (a) of the IscS complex (blue and light blue) with TusA⁹ (different gradations of green, 3LVJ). (c) Blowup of Figure 6a but in a ribbon representation showing the relative position of pyridoxal phosphate (PLP, magenta), the catalytic loop (purple), the three conserved cysteines of IscU (yellow) and Trp61 of CyaY (green). The catalytic loop was built by homology using the 3GZC47 coordinates the sequence of which has a sequence similarity/identity of 30 and 50%, respectively, and superposes to 1P3W, with a lower root mean square deviation of 1.21 Å. The resulting model was energy minimized by the Gromacs package⁵⁵.

stabilizing the interaction with IscU, thus explaining why this can be observed in the absence of desulphurase only in eukaryotes.

Interestingly, IscU recognition seems to involve CyaY Trp61, as directly suggested by the SAXS models and the NMR data. This interaction could explain the high degree of conservation of the Trp throughout the frataxin family and the severe effects that mutation of this residue have *in vitro* and *in vivo*^{35,36}.

Even though we should not at this stage overspeculate, as the resolution we are working at is insufficient to discuss atomic details, it is clear from our data that formation of the IscS/IscU/CyaY complex does not require the presence of iron. This is at variance with previous reports. Pull-down experiments on the yeast complex have suggested iron to be essential for the interaction³. In another study, it was suggested that binding of Yfh1 requires oligomerization and occurs in an iron-independent manner¹¹. However, the surface of interaction involves a direct recognition of the negatively charged region of CyaY by a positively charged patch on IscS, thus rendering an active role of iron in complex formation unlikely.

We confirmed our previous observation that CyaY slows down Fe-S cluster formation and proved that the effect is directly related to the interaction between CyaY and IscS¹³. This behaviour could be

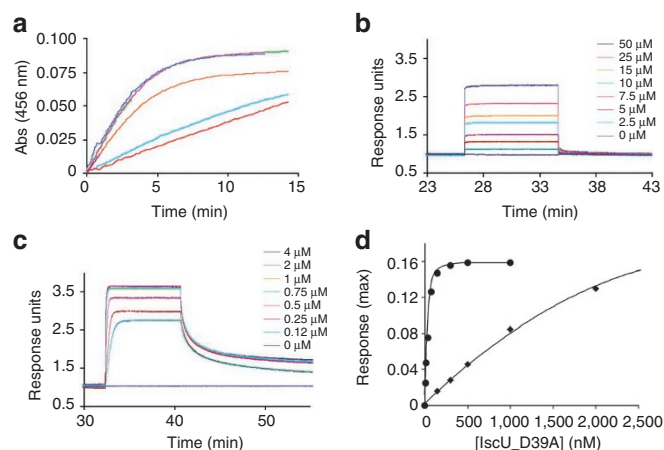


Figure 7 | Influence of complex formation on the kinetics of cluster formation and on the complex affinities. (a) Kinetics rates of Fe-S cluster formation on IscU. The experiments are all conducted under the same conditions (1 μM IscS or IscS mutants, 25 μM Fe²⁺, 50 μM IscU, 250 μM Cys, 2 mM DTT and 5 μM of CyaY or CyaY mutants). Green curve: control experiment in the absence of CyaY; red curve: control experiment in the presence of wild-type CyaY; blue, magenta, orange and cyan curves: experiments carried out by substituting wild-type proteins with CyaY_E18K_E19K_E22K, IscS_R220E_R223E_R225E, CyaY_W61R and IscS_I314E_M315E, respectively. (b) Biolayer interferometry profile of the titration of CyaY with increasing concentrations of IscS in the absence of IscU. (c) As in b, but in the presence of IscU. Because the complex is stabilized, lower concentrations of IscS are needed to reach saturation. (d) Plot of the interferometry response obtained by immobilizing IscS as a function of increasing concentrations of cluster-free (circle) and cluster-loaded (diamond) IscU_D39A mutant.

explained in several ways. First, CyaY could directly affect the enzymatic properties of IscS. However, the catalytic site is in a region close but quite distinct from those hosting IscU and CyaY in our structure, in full agreement with our previous findings that conversion of cysteine into alanine is unperturbed by the presence of CyaY¹³. Second, CyaY could interfere with the cluster transfer from IscS to IscU, for instance, by impairing the motions of the flexible catalytic loop containing Cys328. Evidence that nucleotide modification, which is one of the alternative functions of IscS, is not impaired by CyaY-overproducing conditions, seems, however, to also disprove this possibility⁹. A different explanation is directly suggested by our data. We observe that formation of the ternary complex increases the affinities of the interaction between IscU (and of IscU_D39A) and IscS and decreases the dissociation rate constants. The inhibitory effect of CyaY on the enzymatic kinetics could then be the consequence of an impairment of the detachment of IscU from IscS, because of the stabilizing effect of CyaY. This hypothesis implies that, once the cluster is formed, cluster-loaded IscU detaches from IscS to transfer the cluster to the next acceptor and regenerate the reaction, a possibility that is reasonable because the surface of IscU that hosts the cluster is buried in the complex⁹. Accordingly, we have observed an appreciably different affinity of the cluster-free and cluster-loaded forms at least of the IscU_D39A mutant. IscU is also known to interact with the chaperone HscA, another component of the *isc* operon required for Fe-S cluster assembly or delivery from IscU to target proteins, with a site at least partially overlapping with the surface involved in IscS recognition³⁷.

Taken together, these observations suggest the following cycle, even though still awaiting a much deeper description of the exact time sequence of the events. CyaY and IscU are likely to be simultaneously bound to IscS, thus stabilizing the complex and allowing cluster formation. Once the cluster is loaded

on IscU, this would detach from IscS, having a reduced affinity (as hinted by our results with the IscU_D39A mutant) and the interaction being competed out by the chaperone HscA. The complex IscU/HscA, further stabilized by the cochaperone HscB^{38,39}, would deliver the cluster to the final acceptors. Once freed from the cluster, IscU and CyaY would bind to IscS again, thus restarting the cycle. Other competitors might interfere with CyaY and further regulate the process.

What does all this tell us about the cellular function of frataxin?

It is increasingly clear that the role of frataxin cannot simply be that of an iron chaperone, as it would be hard to understand in such a case why frataxin binds the IscU/IscS complex with the same surface that transports iron or why its presence affects the complex stability and the kinetics of Fe-S cluster formation. We suggest a role for frataxin as the regulator of a complex network, centred on the desulphurase of multiple and competing interactions, the nature of which we are just starting to understand. IscS is known to bind not only IscU and CyaY but also the small component of the *isc* operon YfhJ (also known as IscX)^{40,41} and various other proteins involved in completely different pathways, such as TusA, Thil and MoeB/MoaD^{6,7,9}. With the only exception of MoeB/MoaD, which is still undetermined, these interactions have all been mapped in distinct but partially overlapping surfaces of IscS⁹. The different affinities of these proteins for IscS may determine which of these partners will be bound at any one time. The iron-loaded state of at least some of the proteins (for example, iron loaded/iron free, cluster loaded/cluster free) and the relative cellular concentrations may add additional opportunities for achieving regulation.

In conclusion, although much more work is needed to completely clarify the cellular role of frataxin, establishment of the structural bases of its interaction with IscS/IscU opens a new perspective to further advance the field. The information provided in this work may be crucial for understanding both, one of the most essential metabolic pathways and the molecular basis of Friedreich's ataxia.

Methods

Protein purification. *E. coli* CyaY, IscS, IscU and IscU_D39A mutants were purified as previously described^{14,21,23}. In short, they were produced as fusion proteins with a His-tagged glutathione-S-transferase and purified by affinity chromatography using Ni-NTA agarose gel (Qiagen). The cells overexpressing IscU were grown in Luria broth-enriched medium containing 8.3 μM ZnSO₄ to stabilize the protein fold^{21,28,29}. All purification steps were carried out in the presence of 20 mM β-mercaptoethanol. The collected proteins were cleaved overnight from glutathione-S-transferase by Tobacco Etch Virus protease and further purified by gel-filtration chromatography on a Superdex 75 26/60 column (GE Healthcare). Protein purity was checked by SDS-polyacrylamide gel electrophoresis and by mass spectrometry.

SAXS measurements and modelling. SAXS data were collected on the EMBL X33 beamline on the storage ring DORIS III (DESY)⁴². IscS, CyaY and IscU solutions and their complexes were measured at 15 °C in a concentration range of 2.5–5.0 mg ml⁻¹. As CyaY binds both Fe(II) and Fe(III), ferrous ammonium sulphate was added to reach a 1:1:2 CyaY/IscS/Fe molar ratio under aerobic conditions. The data were recorded using a 1 M PILATUS detector (DECTRIS) at a sample-detector distance of 2.7 m and a wavelength of 1.5 Å, covering the range of momentum transfer 0.012 < *s* < 0.6 Å⁻¹. No measurable radiation damage was detected by comparison of four successive time frames with 30 s exposures. The data were averaged after normalization to the intensity of the transmitted beam and the scattering of the buffer was subtracted using PRIMUS⁴².

Forward scattering *I*(0) and the radius of gyration *R*_g were evaluated using the Guinier approximation⁴³. These parameters were also computed from the entire scattering patterns using the program GNOM⁴⁴, which provides the distance distribution functions *p*(*r*) and the maximum particle dimensions *D*_{max}. The solute MM_{exp} was estimated by comparison of the forward scattering with that from reference solutions of bovine serum albumin (MM 66 kDa). The excluded volume of the hydrated particle (the Porod volume *V*_p) was computed as reported by Porod⁴⁵.

Low-resolution *ab initio* models of IscS, IscU, CyaY and their complexes were generated by the program DAMMIF⁴⁶, which represents the protein by an assembly of

densely packed beads. Simulated annealing was used to build a compact interconnected configuration of beads that fits the experimental data *I*(*s*) to minimize discrepancy:

$$\chi^2 = \frac{1}{N-1} \sum_j \left[\frac{I(s_j) - I_{\text{calc}}(s_j)}{\sigma(s_j)} \right]^2$$

where *N* is the number of experimental points, *c* is a scaling factor, *I*_{calc}(*s*) and σ(*s*) are the calculated intensity and the experimental error at the momentum transfer *s*_{*j*}, respectively²⁷. The results from 10 separate runs were averaged to determine common structural features using the program DAMAVER⁴⁶.

The models were further refined by rigid body modelling using the program SASREF³⁰, which uses a simulated annealing protocol to generate an interconnected assembly of subunits without steric clashes that fits the scattering data. The coordinates 1P3W and 1SOY were assumed for isolated IscS and CyaY^{22,25}. As the loop containing Cys328 is not visible in the 1P3W, this was built by homology from the 3GZC⁴⁷ structure and energy minimized. Several structures of IscU were tested. The coordinates of the 1Q48 entry²⁸ were finally selected as those giving the best fit to the data and the best overlap to the *ab initio* envelope of isolated IscU. The scattering amplitudes of the individual subunit structures were calculated using the program CRY SOL⁴⁸.

Docking was performed with a combination of SASREF³⁰ and HADDOCK 1.3 (ref. 31) software, using 3LWL and 1SOY as the starting structures. To improve the resolution, we incorporated the following information. We assumed P2 symmetry for the complexes. Arginines 220, 223 and 225 of IscS were imposed to interact with residues 22–48 of CyaY on the basis of mutagenesis and NMR titrations. For testing purposes, modelling was also achieved without these restraints and provided similar quality fits and similar positions of IscU/CyaY relative to the IscS structure. The accuracy of the resulting models was validated by considering not only our own results (mutagenesis and NMR studies) but also literature data⁹ (Supplementary Table S2). Solutions that did not fulfil all the mutagenesis data were discarded.

NMR titrations. NMR spectra were acquired on ¹⁵N uniformly labelled CyaY or IscU samples and ¹⁵N-²H-¹³C uniformly labelled CyaY samples in 90%/10% H₂O/D₂O. All proteins used in the titration experiments, both labelled and non-labelled, were in 20 mM HCl-Tris at pH 8.0, in 150 mM NaCl and in 20 mM β-mercaptoethanol. All spectra were recorded at 25 °C on a Varian spectrometer operating at 800 MHz ¹H frequencies or on a Bruker spectrometer operating at 700 MHz ¹H frequencies and equipped with 5 mm triple-resonance probes or cryoprobes. Water suppression was achieved by the WATERGATE sequence⁴⁹. The spectra were processed with NMRPipe⁵⁰ and analysed with XEASY⁵¹.

ITC measurements. Complex formation between IscU and IscS was measured by ITC using a MicroCal VP-ITC microcalorimeter (MicroCal)⁵². Proteins were dialysed overnight into the same buffer (20 mM Tris-HCl at pH 8.0, 150 mM NaCl and 10 mM β-mercaptoethanol). When anaerobic conditions were necessary (that is, for testing the CyaY/IscU interaction in the presence of Fe²⁺), the solutions were thoroughly degassed and prepared in an anaerobic chamber (Belle technology). The experiments were conducted at 25 °C. The sample cell typically contained IscS (16–25 μM or 180–200 μM concentrations) and the syringe contained either IscU (180–250 μM) or CyaY (1.7–1.9 mM). For the studies of the IscS/IscU/CyaY ternary complex, the sample cell contained IscS (16–25 μM) saturated by CyaY (380–420 μM).

Heat of dilution, as determined by titrating IscU or CyaY into the buffer alone and subtracted from the raw titration data before analysis, was always negligible as compared with the heat signal observed for the initial titrant additions. Data were fit by least-squares procedures using Microcal Origin (version 7.0) provided by the manufacturer. All titrations using IscU were fit using the 'ligand in the cell' assumption. *K*_d values were averaged over three measurements.

Biolayer interferometry. All experiments were conducted aerobically in 20 mM HEPES pH 7.5, 150 mM NaCl, 2 mM tris(2-carboxyethyl)phosphine and 0.5 mg ml⁻¹ bovine serum albumin on an Octet Red instrument (ForteBio), operating at 25 °C. Streptavidin-coated biosensors with immobilized biotinylated IscS were exposed to different concentrations of IscU (0–5 μM) in the presence and absence of 100 μM CyaY or to cluster-free and cluster-loaded IscU_D39A mutant (0–4 μM). Alternatively, streptavidin-coated biosensors with immobilized biotinylated CyaY were exposed to different concentrations of IscS (0–50 μM) in the presence and absence of 10 μM IscU.

Fe-S cluster reconstitution on IscU or IscU_D39A. Cluster reconstitution was performed under strict anaerobic conditions in a chamber (Belle technology) kept under nitrogen atmosphere. The reaction was followed by absorbance spectroscopy using a Cary 50 Bio spectrophotometer (Varian). Variations in the absorbance at 456 nm were measured as a function of time. A solution of 50 μM IscU was incubated in sealed cuvettes typically using 3 mM DTT and 25 μM Fe(NH₄)₂SO₄ for 30 min in 50 mM Tris-HCl buffer, pH 7.5 and 150 mM NaCl. The reaction was initiated by adding IscS (1 μM) and the reaction substrate cysteine (250 μM).

The relative concentrations of the assay components were chosen as previously detailed⁸. In short, the concentration of IscU, which is the final cluster acceptor in the assay, was set to 50 μM, that is, a value that allowed us to easily measure the

ultraviolet signal of the cluster. IscS was set to catalytic concentrations (1 μM) to avoid even minimal contributions of unspecific iron-thiolate polysulphides bound to IscS⁵³. The IscS/IscU molar ratio also determines the time scale in which the kinetics are completed⁵⁴. Similarly, we used small concentrations of CyaY (typically 5 μM). The presence of the antioxidant DTT is necessary to provide reducing equivalents required for cluster generation and to regenerate the prosthetic group pyridoxal phosphate²⁰.

Fe-S cluster reconstitution on the IscU_D39A mutant was obtained using the same protocol as for wild-type IscU and incubating the protein in anaerobic conditions for 4h at room temperature.

References

- Pandolfo, M. & Pastore, A. The pathogenesis of Friedreich ataxia and the structure and function of frataxin. *J. Neurol.* **256** (Suppl 1), 9–17 (2009).
- Cossee, M. *et al.* Inactivation of the Friedreich ataxia mouse gene leads to early embryonic lethality without iron accumulation. *Hum. Mol. Genet.* **9**, 1219–1226 (2000).
- Gerber, J., Muhlenhoff, U. & Lill, R. An interaction between frataxin and Isu1/Nfs1 that is crucial for Fe/S cluster synthesis on Isu1. *EMBO Rep.* **4**, 906–911 (2003).
- Yoon, T. & Cowan, J. A. Iron-sulfur cluster biosynthesis. Characterization of frataxin as an iron donor for assembly of [2Fe-2S] clusters in ISU-type proteins. *J. Am. Chem. Soc.* **125**, 6078–6084 (2003).
- Ramazzotti, A., Vanmansart, V. & Foury, F. Mitochondrial functional interactions between frataxin and Isu1p, the iron-sulfur cluster scaffold protein, in *Saccharomyces cerevisiae*. *FEBS Lett.* **557**, 215–220 (2004).
- Johnson, D. C., Dean, D. R., Smith, A. D. & Johnson, M. K. Structure, function, and formation of biological iron-sulfur clusters. *Annu. Rev. Biochem.* **74**, 247–281 (2005).
- Kessler, D. Enzymatic activation of sulfur for incorporation into biomolecules in prokaryotes. *FEMS Microbiol. Rev.* **30**, 825–840 (2006).
- Zhang, W. *et al.* IscS functions as a primary sulfur-donating enzyme by interacting specifically with MoEB and MoaD in the biosynthesis of molybdopterin in *Escherichia coli*. *J. Biol. Chem.* **285**, 2302–2308 (2010).
- Shi, R. *et al.* Structural basis for Fe-S cluster assembly and tRNA thiolation mediated by IscS protein-protein interactions. *PLoS Biol.* **8**, e1000354 (2010).
- Huang, J., Dizin, E. & Cowan, J. A. Mapping iron binding sites on human frataxin: implications for cluster assembly on the ISU Fe-S cluster scaffold protein. *J. Biol. Inorg. Chem.* **13**, 825–836 (2008).
- Li, H., Gakh, O., Smith, D. Y. IV & Isaya, G. Oligomeric yeast frataxin drives assembly of core machinery for mitochondrial iron-sulfur cluster synthesis. *J. Biol. Chem.* **284**, 21971–21980 (2009).
- Layer, G., Ollagnier-de Choudens, S., Sanakis, Y. & Fontecave, M. Iron-sulfur cluster biosynthesis: characterization of *Escherichia coli* CyaY as an iron donor for the assembly of [2Fe-2S] clusters in the scaffold IscU. *J. Biol. Chem.* **281**, 16256–16263 (2006).
- Adinolfi, S. *et al.* Bacterial frataxin CyaY is the gatekeeper of iron-sulfur cluster formation catalyzed by IscS. *Nat. Struct. Mol. Biol.* **16**, 390–396 (2009).
- Musco, G. *et al.* Towards a structural understanding of Friedreich's ataxia: the solution structure of frataxin. *Structure* **8**, 695–707 (2000).
- Cho, S. J. *et al.* Crystal structure of *Escherichia coli* CyaY protein reveals a previously unidentified fold for the evolutionarily conserved frataxin family. *Proc. Natl Acad. Sci. USA* **97**, 8932–8937 (2000).
- Dhe-Paganon, S., Shigetani, R., Chi, Y. I., Ristow, M. & Shoelson, S. E. Crystal structure of human frataxin. *J. Biol. Chem.* **275**, 30753–30756 (2000).
- Creighton, T. E. Protein folding. Up the kinetic pathway. *Nature* **356**, 194–195 (1992).
- Cavadini, P., O'Neill, H. A., Benada, O. & Isaya, G. Assembly and iron-binding properties of human frataxin, the protein deficient in Friedreich ataxia. *Hum. Mol. Genet.* **11**, 217–227 (2002).
- Foury, F., Pastore, A. & Trincal, M. Acidic residues of yeast frataxin have an essential role in Fe-S cluster assembly. *EMBO Rep.* **8**, 194–199 (2007).
- Urbina, H. D., Silberg, J. J., Hoff, K. G. & Vickery, L. E. Transfer of sulfur from IscS to IscU during Fe/S cluster assembly. *J. Biol. Chem.* **276**, 44521–44526 (2001).
- Prischi, F. *et al.* Of the vulnerability of orphan complex proteins: the case study of the *E. coli* IscU and IscS proteins. *Protein Expr. Purif.* **73**, 161–166 (2010).
- Cupp-Vickery, J. R., Urbina, H. & Vickery, L. E. Crystal structure of IscS, a cysteine desulfurase from *Escherichia coli*. *J. Mol. Biol.* **330**, 1049–1059 (2003).
- Adinolfi, S. *et al.* Bacterial IscU is a well folded and functional single domain protein. *Eur. J. Biochem.* **271**, 2093–2100 (2004).
- Adinolfi, S., Trifuoggi, M., Politou, A. S., Martin, S. & Pastore, A. A structural approach to understanding the iron-binding properties of phylogenetically different frataxins. *Hum. Mol. Genet.* **11**, 1865–1877 (2002).
- Nair, M. *et al.* Solution structure of the bacterial frataxin ortholog, CyaY: mapping the iron binding sites. *Structure* **12**, 2037–2048 (2004).
- Franke, D. & Svergun, D. I. DAMMIF, a program for rapid ab-initio shape determination in small-angle scattering. *J. Appl. Crystallogr.* **42**, 342–346 (2009).
- Svergun, D. I. Restoring low resolution structure of biological macromolecules from solution scattering using simulated annealing. *Biophys. J.* **76**, 2879–2886 (1999).
- Ramelot, T. A. *et al.* Solution NMR structure of the iron-sulfur cluster assembly protein U (IscU) with zinc bound at the active site. *J. Mol. Biol.* **344**, 567–583 (2004).
- Kim, J. H. *et al.* Structure and dynamics of the iron-sulfur cluster assembly scaffold protein IscU and its interaction with the cochaperone HscB. *Biochemistry* **48**, 6062–6071 (2009).
- Petoukhov, M. V. & Svergun, D. I. Global rigid body modeling of macromolecular complexes against small-angle scattering data. *Biophys. J.* **89**, 1237–1250 (2005).
- Dominguez, C., Boelens, R. & Bonvin, A. M. HADDOCK: a protein-protein docking approach based on biochemical or biophysical information. *J. Am. Chem. Soc.* **125**, 1731–1737 (2003).
- Foster, M. W. *et al.* A mutant human IscU protein contains a stable [2Fe-2S]²⁺ center of possible functional significance. *J. Am. Chem. Soc.* **122**, 6805–6806 (2000).
- Shimomura, Y., Takahashi, Y., Kakuta, Y. & Fukuyama, K. Crystal structure of *Escherichia coli* YfhJ protein, a member of the ISC machinery involved in assembly of iron-sulfur clusters. *Proteins* **60**, 566–569 (2005).
- Schagerlof, U. *et al.* Structural basis of the iron storage function of frataxin from single-particle reconstruction of the iron-loaded oligomer. *Biochemistry* **47**, 4948–4954 (2008).
- Leidgens, S., De Smet, S. & Foury, F. Frataxin interacts with Isu1 through a conserved tryptophan in its beta-sheet. *Hum. Mol. Genet.* **19**, 276–286 (2010).
- Correia, A. R., Ow, S. Y., Wright, P. C. & Gomes, C. M. The conserved Trp155 in human frataxin as a hotspot for oxidative stress related chemical modifications. *Biochem. Biophys. Res. Commun.* **390**, 1007–1011 (2009).
- Cupp-Vickery, J. R., Peterson, J. C., Ta, D. T. & Vickery, L. E. Crystal structure of the molecular chaperone HscA substrate binding domain complexed with the IscU recognition peptide ELPPVKIHC. *J. Mol. Biol.* **342**, 1265–1278 (2004).
- Hoff, K. G., Silberg, J. J. & Vickery, L. E. Interaction of the iron-sulfur cluster assembly protein IscU with the Hsc66/Hsc20 molecular chaperone system of *Escherichia coli*. *Proc. Natl Acad. Sci. USA* **97**, 7790–7795 (2000).
- Bonomi, F., Iametti, S., Morleo, A., Ta, D. & Vickery, L. E. Studies on the mechanism of catalysis of iron-sulfur cluster transfer from IscU[2Fe2S] by HscA/HscB chaperones. *Biochemistry* **47**, 12795–12801 (2008).
- Tokumoto, U. & Takahashi, Y. Genetic analysis of the isc operon in *Escherichia coli* involved in the biogenesis of cellular iron-sulfur proteins. *J. Biochem.* **130**, 63–71 (2001).
- Pastore, C. *et al.* YfhJ, a molecular adaptor in iron-sulfur cluster formation or a frataxin-like protein? *Structure* **14**, 857–867 (2006).
- Roessle, M. W. *et al.* Upgrade of the small-angle X-ray scattering beamline X33 at the European Molecular Biology Laboratory, Hamburg. *J. Appl. Crystallogr.* **40** (Suppl), s190–s194 (2007).
- Guinier, A. La diffraction des rayons X aux très faibles angles: Applications à l'étude des phénomènes ultra-microscopiques. *Ann. Phys. Paris* **12**, 161–237 (1939).
- Svergun, D. I. Determination of the regularization parameter in indirect-transform methods using perceptual criteria. *J. Appl. Crystallogr.* **25**, 495–503 (1992).
- Porod, G. *In Small-Angle X-ray Scattering* (Academic Press, 1982).
- Volkov, V. V. & Svergun, D. I. Uniqueness of ab initio shape determination in small-angle scattering. *J. Appl. Crystallogr.* **36**, 860–864 (2003).
- Tirupati, B., Vey, J. L., Drennan, C. L. & Bollinger, J. M. Jr. Kinetic and structural characterization of Slr0077/SufS, the essential cysteine desulfurase from *Synechocystis* sp. PCC 6803. *Biochemistry* **43**, 12210–12219 (2004).
- Svergun, D. I., Barberato, C. & Koch, M. H. J. CRYSOLO—a program to evaluate X-ray solution scattering of biological macromolecules from atomic coordinates. *J. Appl. Crystallogr.* **28**, 768–773 (1995).
- Piotto, M., Saudek, V. & Sklenar, V. Gradient-tailored excitation for single-quantum NMR spectroscopy of aqueous solutions. *J. Biomol. NMR* **2**, 661–665 (1992).
- Delaglio, F. *et al.* NMRPipe: a multidimensional spectral processing system based on UNIX pipes. *J. Biomol. NMR* **6**, 277–293 (1995).
- Bartels, C., Xia, T., Billeter, M., Güntert, P. & Wüthrich, K. The program XEASY for computer-supported NMR spectral analysis of biological macromolecules. *J. Biomol. NMR* **6**, 1–10 (2004).
- Wiseman, T., Williston, S., Brandts, J. F. & Lin, L. N. Rapid measurement of binding constants and heats of binding using a new titration calorimeter. *Anal. Biochem.* **179**, 131–137 (1989).
- Bonomi, F., Pagani, S. & Kurtz, D. M. Jr. Enzymic synthesis of the 4Fe-4S clusters of *Clostridium pasteurianum* ferredoxin. *Eur. J. Biochem.* **148**, 67–73 (1985).
- Agar, J. N. *et al.* IscU as a scaffold for iron-sulfur cluster biosynthesis: sequential assembly of [2Fe-2S] and [4Fe-4S] clusters in IscU. *Biochemistry* **39**, 7856–7862 (2000).
- Van Der Spoel, D. *et al.* GROMACS: fast, flexible, and free. *J. Comput. Chem.* **26**, 1701–1718 (2005).

Acknowledgments

We acknowledge financial support from the Medical Research Council, as well as support from the MRC (Grant ref. U117584256) and from the EFACTS EU grant. We are also thankful to the MRC NMR Centre for technical support and to Pierandrea Temussi for critical reading of the manuscript.

Author contributions

F.P. and P.V.K. designed, conducted and analysed the SAXS experiments under the guidance of D.I.S.; C.I. and S.R.M. conducted the interferometry experiments and analysed the data. S.A. produced the original clones and carried out the enzyme kinetics experiments. C.P. contributed to the NMR data titration together with F.P. A.P. analysed the data and wrote the paper. All authors discussed the results and commented on the paper.

Additional information

Supplementary Information accompanies this paper on <http://www.nature.com/naturecommunications>

Competing financial interests: The authors declare no competing financial interests.

Reprints and permission information is available online at <http://npg.nature.com/reprintsandpermissions/>

How to cite this article: Prischi, F. *et al.* Structural bases for the interaction of frataxin with the central components of iron–sulphur cluster assembly. *Nat. Commun.* 1:95 doi: 10.1038/ncomms1097 (2010).

License: This work is licensed under a Creative Commons Attribution-NonCommercial-Share Alike 3.0 Unported License. To view a copy of this license, visit <http://creativecommons.org/licenses/by-nc-sa/3.0/>