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Structure of the Thioredoxin-Like Domain of Yeast Glutaredoxin 3

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Structure of the thioredoxin-like domain of yeast glutaredoxin 3

Yeast glutaredoxin 3 (Grx3) is a cytosolic protein that regulates the activity of the iron-responsive transcriptional activator Aft1. This member of the monothiol glutaredoxin family contains a thioredoxin-like domain and a glutaredoxin-like domain, which both possess a monothiol active site. The crystal structure of the thioredoxin-like domain has been determined at 1.5 Å resolution and represents the first published structure of this domain for the monothiol glutaredoxin family. The loop containing the signature motif WAxxC is partially disordered, indicating a greater degree of flexibility in this region compared with classical dithiol thioredoxins with a WCGPC active-site motif.

1. Introduction

Thioredoxins (Trxs) and glutaredoxins (Grxs) are small oxidoreductases (usually 9–15 kDa) that are found in all types of organisms. Both proteins are members of the thioredoxin-fold superfamily and typically catalyze thiol–disulfide exchange reactions. The classical dithiol Trxs utilize a highly conserved WCGPC active-site motif to reduce disulfides in substrate proteins. The dithiol Grxs possess a CPYC active-site motif and primarily catalyze glutathionylation and deglutathionylation of target proteins (Berndt et al., 2008). A subfamily of Grxs were recently identified that are characterized by a conserved monothiol CGFS active-site motif (Herrero & de la Torre-Ruiz, 2007). *Saccharomyces cerevisiae* has three Grx-like proteins that belong to this subclass: yGrx3, yGrx4 and yGrx5. yGrx5 contains a single Grx-like domain and is localized to the mitochondrial matrix, where it plays an ill-defined role in Fe–S cluster biogenesis (Rodriguez-Manzaneque et al., 2002). However, recent evidence from studies of yGrx5 homologues in chloroplasts suggest that this protein may act as a scaffold for Fe–S cluster assembly and transfer (Bandyopadhyay et al., 2008). Homologues of yGrx5 with a single Grx-like domain are found in both prokaryotes and eukaryotes (Herrero & de la Torre-Ruiz, 2007).

Yeast Grx3 and Grx4 are highly homologous proteins that possess both an N-terminal Trx-like domain and a C-terminal Grx-like domain (Herrero & de la Torre-Ruiz, 2007). Both proteins are cytosolic and are required for iron-responsive inhibition of the transcriptional activator Aft1 (Kumanovics et al., 2008; Ojeda et al., 2006; Pujol-Carrion et al., 2006). The conserved cysteine in the CGFS active site located in the Grx-like domain of yGrx3 and yGrx4 is required for interaction with Aft1 and regulation of its activity (Ojeda et al., 2006).
Trx-like domains have a WAxPC motif in place of the signature WCGPC active-site motif found in classical dithiol Trxs. Monothiol Grxs with this hybrid Trx-Grx domain architecture are only found in eukaryotes (Herrero & de la Torre-Ruiz, 2007). In general, the Grx-like domains of this protein family are more highly conserved than the Trx-like domains (Isakov et al., 2000). The human homologue of yGrx3/4 (termed PICOT for PKC-interacting cousin of Trx) has an N-terminal Trx-like domain followed by two Grx-like domains (Isakov et al., 2000). The Trx-like domain specifically interacts with protein kinase C-1C (PKC-1C) and is proposed to play a role in regulating the function of this kinase and its downstream targets (Witte et al., 2000).

Here, we present the crystal structure of the N-terminal Trx-like domain of yGrx3 determined at 1.5 Å resolution. This report represents the first published structural information for the Trx-like domain for any member of the monothiol glutaredoxin/PICOT family.

2. Materials and methods

2.1. Protein expression and crystallization

The yGrx3 open reading frame containing both the Trx-like and Grx-like domains but lacking the first 35 amino acids (A1–35) was cloned into PET21a and expressed in Escherichia coli BL21 (DE3) (Novagen) as described previously (Kumanovics et al., 2008). Briefly, transformants were grown at 310 K until OD600 reached 0.6 and were induced with 1 mM isopropyl β-D-1-thiogalactopyranoside. Cells were harvested 3 h after induction and stored at 193 K. Cells were lysed using three freeze–thaw cycles and soluble protein was extracted with 50 mM Tris–HCl pH 8.0, 5 mM DTT. The extract was loaded onto a DEAE anion-exchange column (GE Healthcare) equilibrated with 25 mM Bis-Tris pH 6.0, 5 mM DTT and eluted with a salt gradient. The fractions containing yGrx3 were further purified by gel filtration using a Superdex 75 column (GE Healthcare) in 50 mM Tris–HCl pH 8.0, 150 mM NaCl, 5 mM DTT. The final pure protein was concentrated with the addition of 5% glycerol and stored at 193 K. The typical yield was ~30 mg of protein per litre of cell culture. Protein purity was checked by SDS–PAGE and MALDI–TOF, indicating a single species corresponding to 28 120 kDa (the calculated molecular weight is 28 130 Da upon removal of Met36).

2.2. Crystallization and data collection

The yGrx3 crystallized in the monoclinic space group P21, with unit-cell parameters a = 47.91 Å, b = 48.61 Å, c = 55.29 Å, β = 116.8°, space group P21, and unit-cell parameters a = 47.91 Å, b = 48.61 Å, c = 55.29 Å, β = 116.8°. The crystals were cryoprotected in 50% glycerol, 25 mM Bis-Tris pH 6.0, 5 mM DTT, and flash-cooled in liquid nitrogen. The crystals were then cryo-protected in 50% glycerol, 25 mM Bis-Tris pH 6.0, 5 mM DTT, and flash-cooled in liquid nitrogen.

Table 1
Crystallographic data and refinement statistics.

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2.3. Structure determination

The structure was solved by molecular replacement using the program Phaser (McCoy et al., 2007) with the Trx-like domain of hTrx (PDB code 1ert) as the search model. The final model was refined using the program REFMAC (Murshudov et al., 2011) and validated using the program MolProbity (Chen et al., 2010).

Figure 1
Comparison of yGrx3 Trx domain and hTrx. (a) Sequence alignment of the crystallized form of yGrx3 (amino acids 37–148) and the complete sequence of hTrx. The hTrx active-site motif and cis-Pro75 are indicated by asterisks. (b) Superposition of yGrx3 Trx domain (green) with Cys72 in green atom type and hTrx (PDB code 1ert; blue) with Cys32 and Cys35 in blue atom type. The positions of the amino-acid inserts are marked by arrows.
The final crystallization conditions were optimized from initial crystallization-condition leads obtained using Crystal Screen Lite from Hampton Research. Crystals of yGrx3 were grown under high-salt conditions by the hanging-drop method at 300 K. Briefly, 2 μl protein solution (11 mg ml⁻¹ protein in 50 mM Tris pH 8.0 and 5 mM DTT) was mixed with 2 μl reservoir solution (0.1 M MOPS pH 7.5, 20 mM βME and 68% saturated ammonium sulfate).

2.2. Data collection and processing

A yGrx3 crystal was transferred to cryosolvent (60% ammonium sulfate, 20 mM βME, 0.1 M MOPS and 20% ethylene glycol) for a few seconds and subsequently flash-frozen in a liquid N₂ vapor stream. Data were collected on the SER-CAT 22ID beamline at Argonne National Laboratory at a 1.0000 Å wavelength in two passes: a first pass with a crystal-to-detector distance of 200 mm and a second pass with a crystal-to-detector distance of 300 mm, both using a 0.5° oscillation range. The data were indexed and processed using the HKL-2000 software (Otwinowski & Minor, 1997); processing parameters and statistics are summarized in Table 1.

2.3. Structure determination and refinement

The structure was solved by molecular replacement using AMoRe (Navaza, 1994) from the CCP4 system (Collaborative Computational Project, Number 4, 1994) with the structure of the reduced form of human thioredoxin (hTrx) as the search model (PDB code 1ert; Weichsel et al., 1996). The amino-acid sequence of hTrx is 32% identical to that of the Trx-like domain of yGrx3 (see Fig. 1a). The initial model was rebuilt using ARP/wARP (Perrakis et al., 1999) from the CCP4 suite, while subsequent interactive molecular-graphics and crystallographic refinements were carried out using TURBO FRODO (Roussel & Cambillau, 1991) and CNS (Brünger et al., 1998), respectively. The final refinement was carried out with REFMACS (Murshudov et al., 1997). Superpositions were calculated using the LSQKAB program (Kabsch, 1976) from the CCP4 suite. Figures were prepared using TURBO FRODO and PyMOL (DeLano, 2002).

3. Results and discussion

3.1. Crystallization

The recombinant yGrx3 protein was cloned from the second start site (encoding Met36) to the stop codon after determining that the first start site (encoding Met1) was not utilized in vivo (N. Dingra & C. Outten, unpublished data). Furthermore, MALDI–TOF analysis indicated that Met36 is removed upon overexpression, as is typically found when the penultimate amino acid is relatively small (Pro37 in this case; Miller, 1975). Crystallization experiments were performed using this 28 kDa protein that included both the Trx-like and Grx-like domains (amino acids 37–285). Two forms of crystals appeared after about four weeks under similar conditions. One form was tetragonal and diffracted poorly, while the other monoclinic form was superior and its characterization is the focus of this report. However, only the Trx-like domain is present in the monoclinic crystals, with two protomers per asymmetric unit and amino acids 37–148 identifiable in the electron density. The solvent content calculated with...
the two Trx-like domains in the asymmetric unit was 46%, which is a typical value, while the solvent content calculated for two molecules of the Trx-Grx chimera protein was below 0%. Therefore, we believe that the Grx-like domain was cleaved off during crystallization and the Trx-like domain was selectively crystallized. Dissociation of the Grx-like domain during crystallization suggests that the two domains do not maintain a strong interaction.

Subsequently, the Trx domain (amino acids 36–157) of yGrx3 was overexpressed (N. Dingra & C. Outten, unpublished data). Its crystallization under similar conditions yielded both crystal forms. Tetragonal crystals appeared in 2 d and belonged to the P4 space group with unit-cell parameters a = 76.42, c = 37.16 Å. Despite good morphology and a size of ~0.3 mm, their diffraction only extended to ~3.5 Å resolution and thus this crystal form was not investigated further. Crystals of the monoclinic form appeared in 4 d (in contrast to the month needed to grow them from yGrx3). They were isomorphic to the crystals obtained from proteolysis of yGrx3 and had similar scattering power. Their structure was essentially the same and thus is not discussed here in detail.

3.2. Overall structure

Overall, both protomers have good electron density as shown in Fig. 2. A sequence alignment of the crystallized protein with hTrx is given in Fig. 1(a). The model contains residues 37–148, but both subunits have a gap in the density in the putative active-site region (Fig. 1b). In subunit A the gap is between residues 68 and 71, while in subunit B the gap is somewhat larger, between residues 68 and 74. This region is located on a surface-exposed loop. The two protomers are very similar: the r.m.s. distance between the Cα positions is 0.58 Å and they have similar corresponding temperature factors.

The yGrx3 Trx-like domain differs from hTrx by having one amino acid inserted after position 52 and two amino acids inserted after position 85 (Fig. 1a). Despite these differences, the overall fold of the yGrx3 Trx-like domain remains very similar to other known thioredoxins with the typical ββαβαβαβα β topology. The r.m.s. distance between the Cα positions in the Trx-like domain A and in hTrx (PDB code 1ert) is 1.14 Å.

3.3. Putative active-site region

The signature active-site WCGPC motif found in classical dithiol thioredoxins is different in the yGrx3 Trx-like domain and aligns with the sequence WAEPC in the putative active site. Density is present for the single Cys72 in subunit A, which is moderately good for the main chain. It is likely that its side chain populates two conformations with some shift of the main chain involved. There is no density for Cβ, but there are two distinct peaks in electron density that are 2.7 Å apart at positions that approximately correspond to two rotamers of the thiol. This distance is too short for a simple SH—H2O contact, which should be ~3.2 Å, but some combination of water/thiol disorder is clearly a possibility. Interestingly, the two positions of the partially disordered thiol group roughly correspond to the positions of the two cysteines found in dithiol thioredoxins (Fig. 3). The position of Pro71 is different research papers

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but well defined, while the residues corresponding to the rest of the signature motif (Ala69–Glu70) are disordered, indicating flexibility in this part of the structure. The electron density for Trp68 indicates that it is only slightly shifted in comparison to the corresponding residue in hTrx. There is a question whether the catalytic loop disorder might be related to the proteolysis of yGrx3, since this is a surface loop. This appeared to be possible when only the structure of the proteolytically cleaved Trx domain was available. However, the structure of the recombinant Trx domain also has the catalytic loop disordered. The recombinant Trx protein showed limited proteolysis after storage for about a month, but mass spectrometry showed that none of the major fragments corresponded to a cut in the disordered region.

The lack of conservation of active-site residues is not limited to the CxxC motif. Asp26 in hTrx, which acts as a general acid–base catalyst and deprotonates the N-terminal cysteine in the WCGPC active site, is not present. The residues corresponding to Asp26 in monothiol Trx proteins are quite variable (histidine, tyrosine and leucine have been observed; Fig. 4), so its role is not likely to be significant. The lack of conservation of the active-site residues suggests that yGrx3 does not catalyze disulfide-bond reduction and that its regulatory function involves a different mechanism. However, it is possible that another cysteine from the Grx-like domain combines with Cys72 to form a pair of thiols with reducing capabilities. If that is indeed the case, the flexibility of the region containing Cys72 suggests that its thiol could penetrate substrate proteins more efficiently to reach a buried disulfide bond. For instance, human thymidylate synthase contains a highly reactive catalytic cysteine in the active-site cavity that readily forms a covalent bond with oxidized glutathione (L. M. Gibson, work to be published). To reduce such a buried bond, an oxidoreductase would have to penetrate the active-site cavity. This hypothesis can be perhaps verified when the structure of the recombinant Trx domain also has the catalytic loop disordered. The recombinant Trx protein showed limited proteolysis after storage for about a month, but mass spectrometry showed that none of the major fragments corresponded to a cut in the disordered region.

References


