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Structure of the Human FANCL RING-Ube2T Complex Reveals Determinants of Cognate E3-E2 Selection

Charlotte Hodson, Andrew Purkiss, Jennifer Anne Miles, and Helen Walden

INTRODUCTION

Ubiquitination is a reversible posttranslational modification in which ubiquitin (Ub) is covalently attached via its C terminus, typically to a substrate lysine. Ubiquitination is required for the strict regulation of a wide range of essential cellular processes, from protein degradation to DNA repair and cell-cycle control (Pickart and Eddins, 2004). Consequently, defects that arise in the regulation of ubiquitination can lead to a variety of diseases, such as cancers and neurodegeneration.

Substrate ubiquitination is achieved through an enzyme cascade involving an E1 activating enzyme, an E2 ubiquitin-conjugating (UBC) enzyme, and an E3 ligase. The E3 ligase, in combination with its partnered E2 enzyme, coordinates the transfer of ubiquitin onto a specific lysine residue. The E3-E2 pair also dictates the type of modification, ranging from monoubiquitination to Ub polymers (Ye and Rape, 2009). The human genome encodes two E1 enzymes, approximately 40 E2s and over 600 E3 ligases, giving rise to thousands of possible permutations of E3-E2 pairings.

RESULTS AND DISCUSSION

FANCL and Ube2T coelute as a 1:1 stoichiometric complex by size-exclusion chromatography and have an affinity with a dissociation constant (K_d) of ~0.5 μM (Hodson et al., 2011). The experimentally determined structures of E3-E2 complexes have revealed a well-conserved hydrophobic interaction surface, encompassing loops 1 and 2 and the first helix of the E2. (Bentley et al., 2011; Dou et al., 2012; Huang et al., 1999; Plechanovalova et al., 2012; Pruneda et al., 2012; Yin et al., 2009; Zheng et al., 2000). Furthermore, the conservation of the E2 UBC fold, along with the conservation of the hydrophobic residues for the E3-interacting interface, suggests that all E3s could function with all E2s (van Wijk and Timmers, 2010). Yet, this is not what is observed in nature, as there is selectivity in E3-E2 pairs with some pairs being exclusive (Bailly et al., 1994; Chen et al., 2006). There have been great efforts using yeast-two-hybrid screens, computational biology, and modeling methods to determine E3-E2 pairs (Kar et al., 2012; Markson et al., 2009; van Wijk et al., 2009). A recent proteome scale modeling study, aimed at identifying determinants of E3-E2 specificity, predicts residues on loop 1 of the E2 to be important for E3 selection (Kar et al., 2012). Additionally, there is much interest in creating new E3-E2 pairs or enhancing specificity (Starita et al., 2013; Winkler and Timmers, 2005), both for understanding ubiquitin biology and from a therapeutic perspective. However, these aims are hampered by the lack of molecular details and structural data as to what constitutes a specific E3-E2 pair.

An example of an exclusive E3-E2 pair is the catalytic center of the Fanconi Anemia (FA) pathway, FANCL-Ube2T (Alpi et al., 2008; Machida et al., 2006). The FA pathway is required for DNA interstrand crosslink repair. Mutations in the FA pathway result in the genetic disorder known as Fanconi Anemia, where patients have high predispositions to cancers because of their genomic instabilities (Alter, 1996). FANCL is a monomeric RING E3 ligase (Cole et al., 2010; Meetee et al., 2003), which specifically interacts with the E2, Ube2T (Alpi et al., 2008; Machida et al., 2006), for the strict monoubiquitination of FANC2 (Garcia-Higuera et al., 2001; Timmers et al., 2001). This monoubiquitination event is key in signaling the recruitment of downstream DNA repair factors (Kottemann and Smogorzewska, 2013).

SUMMARY

The combination of an E2 ubiquitin-conjugating enzyme with an E3 ubiquitin-ligase is essential for ubiquitin modification of a substrate. Moreover, the pairing dictates both the substrate choice and the modification type. The molecular details of generic E3-E2 interactions are well established. Nevertheless, the determinants of selective, specific E3-E2 recognition are not understood. There are ~40 E2s and ~600 E3s giving rise to a possible ~24,000 E3-E2 pairs. Using the Fanconi Anemia pathway exclusive E3-E2 pair, FANCL-Ube2T, we report the atomic structure of the FANCL RING-Ube2T complex, revealing a specific and extensive network of additional electrostatic and hydrophobic interactions. Furthermore, we show that these specific interactions are required for selection of Ube2T over other E2s by FANCL.
complex crystallizes with a diffraction limit of ~11 Å. In order to obtain high-resolution data to observe the interface, we fused human Ube2T to the C terminus of the human FANCL RING domain with a linker between the proteins. Subsequently, we determined the structure for the E3-E2 pair, the FANCL RING domain (residues 299–373), and Ube2T (residues 1–153) to 2.4 Å resolution (Figure 1A; Table 1).

The human RING domain contains two zinc atoms coordinated by a (Cys)₄, His, (Cys)₃ arrangement in a cross-brace structure. The arrangement of cysteine and histidines differs slightly to the (Cys)₃, His, (Cys)₄ arrangement observed in other RING domains. This unusual arrangement is also noted in the Drosophila FANCL structure (Cole et al., 2010) and is conserved across all other FANCL homologs. An overlay with the Drosophila FANCL RING domain reveals that the homologs are highly similar with a root-mean-squared deviation (rmsd) of 1.7 Å across all alpha-carbon atoms (Figure S1A available online). In common with other RING domains, FANCL contains the helical element involved in E2 recognition (Figure 1B) (Deshaies and Joazeiro, 2009). In complex with FANCL, Ube2T adopts a typical UBC-fold comprising a four-stranded β-meander, flanked by an N-terminal helix and two C-terminal helices (Figure 1C). In order to determine whether significant conformational changes occur in Ube2T upon RING binding, we superimposed the bound Ube2T in our structure to unbound Ube2T (Protein Data Bank [PDB] ID code 1YH2) (Sheng et al., 2012). The two structures align with an rmsd of 0.67 Å across all alpha-carbon atoms, indicating that no major structural rearrangements occur upon complex formation (Figure S1B).

The interface between the RING domain and Ube2T buries a total surface area of ~700 Å². In common with other E3-E2 structures, the interface consists of a conserved hydrophobic interface between Pro62, Phe63, and Pro100 of Ube2T and Ile309, Trp341, and Pro360 of FANCL (Figure 1D), as observed in other RING-E2 structures (Bentley et al., 2011; Dou et al., 2012; Plechanovová et al., 2012; Pruneda et al., 2012; Yin et al., 2009; Zheng et al., 2000). However, the hydrophobic surface of FANCL is extended by Tyr311, which is involved in pi stacking between
Arg6 and Arg9 (Figure 1D). Further analysis of the interface reveals an extensive electrostatic and hydrogen bonding network between residues Ser5, Arg6, Arg9, Arg60, Arg99, Ser101, and Asn103 of Ube2T and Asp306, Tyr311, Glu340, and Ser363 of FANCL, with additional main-chain interactions with Ile309, Cys310, and Tyr361 of FANCL (Figure 1D).

Structure-based alignments reveal conservation of the residues attributable to the hydrophobic interface across RING domains and E2s (Figure 2). Based on our observations, we hypothesized that not only the conserved hydrophobic residues Ile309 and Trp341, but also the FANCL-specific Tyr311 revealed by our structure, are important for the FANCL-Ube2T interaction.

To test this hypothesis, we purified single RING-point mutants Ile309Ala, Tyr311Ala, and Trp341Ala. In contrast to wild-type FANCL, Ser5Arg-Ube2T, Arg60Glu, and Arg99Ser/Ser101Arg-Ube2T further anchors Tyr311 of FANCL into position. Ser5 of Ube2T must be attributed to the electrostatic and hydrogen bonding network surrounding a conserved hydrophobic interaction with Ube2T by Tyr311, the corresponding Ube2T-Ser5Arg-Ube2T interaction by mass spectrometry (Figure 3B). Indeed, FANCL exclusively formed a complex with Ube2T, as confirmed by SDS-PAGE analysis of collected fractions and protein identification by mass spectrometry (Figure 3B).

It is clear from our results that FANCL preferentially selects Ube2T. Although FANCL extends its hydrophobic surface for interaction with Ube2T by Tyr311, the corresponding Ube2T-interacting residues Arg6 and Arg9 are conserved in some of the E2s we have tested for FANCL binding and function (e.g., Ube2K, Figure 2A). Therefore, the selectivity of FANCL for Ube2T must be attributed to the electrostatic and hydrogen bonding interactions, which are highly variable among the E2s (Figure 2A). In order to test this hypothesis, we incubated purified mutants of Ube2T Ser5Arg, Arg60Glu, and Arg99Ser/Ser101Arg with the wild-type FANCL RING domain and assessed interaction by size-exclusion chromatography. Only the Arg60Glu mutant of Ube2T is unable to bind the FANCL RING domain (Figure 4A). Consistent with the binding profile of Ube2T mutants, Ser5Arg-Ube2T, Arg99Ser/Ser101Arg-Ube2T, and wild-type Ube2T all support FANCL-dependent monoubiquitination of FANCD2 (Figure 4B). By contrast, Arg60Glu-Ube2T is unable to either bind FANCL or facilitate FANCD2 monoubiquitination (lane 8, Figure 4B). We therefore conclude that the positive selector in Ube2T for FANCL is Arg60, which forms a salt bridge with Glu340 of FANCL (Figure 1D) and is required for FANCL-Ube2T-mediated monoubiquitination of FANCD2.

The dearth of specific E3-E2 structures has hampered the understanding of how E3s select their E2s. Our structure of FANCL-Ube2T reveals a specific extensive electrostatic and hydrogen bonding network surrounding a conserved hydrophobic interaction. In particular, Tyr311 of FANCL, which is a highly variable residue in other E3s, acts like a key in a lock, pi stacking between Arg6 and Arg9 of Ube2T. Additionally the nonconserved Asn103 of Ube2T further anchors Tyr311 of FANCL into position. Ser5 of Ube2T acts as a negative selectivity factor: as in other E2s it is a
Figure 2. Structural Comparison of the FANCL RING Domain-Ube2T Complex with Other RING-E2 Complexes

(A) A structure-based sequence alignment of E2s. PDB ID codes of E2s, as listed in the figure: 1FBV, 3RPG, 4AP4, 4AUQ, 3RZ3, 2YB6, 3K9O, 3H8K, 2Z5D, 2F4W, 3HCT, and 3BZH.

(B) A structure-based sequence alignment of RING and Ubox domains. Ubox domains are highlighted by a cyan box. PDB ID codes used of RING and Ubox domains, as listed in the figure: 1FBV, 4F52, 4AUQ, 3HCT, 2C2V, 3LIZ, 3RPG, 4AP4, 4EPO, 2YHO, 2Y43, 4KBL, and 4K7D. Residues shaded in red to yellow colors indicate conserved residues, where red corresponds to strict conservation. Gray bars indicate zinc coordinating atoms. Green circles highlight residues involved in the hydrophobic interface between FANCL and Ube2T. Purple circles denote residues involved in hydrogen bonding and electrostatic interactions in the FANCL-Ube2T interface.

(C) Superpositions of the FANCL RING-Ube2T complex (colored pink and blue, respectively), with c-cbl-Ube2L3 complex (left) shaded gray (PDB ID code 1FBV), idol-Ube2D1 complex (middle) shaded gray (PDB ID code 2YHO), and ring1b-Ube2D3 complex (right) shaded gray (PDB ID code 3RPG). Numbered residues are the same as the FANCL RING-Ube2T complex, with dashed lines showing interactions.
much larger, bulkier residue and is typically arginine or lysine. These residues would result in loss of the hydrogen bond that occurs between Ser5 of Ube2T and the main chain oxygen of Cys310 of FANCL and potentially clash with FANCL residues Cys310 or Ala312 and/or the main chain of Gln314. Importantly, we have identified Arg60 of Ube2T as the positive selector for FANCL (Figures 1D, 2A, and 4). In other E2s, the equivalent position frequently has the opposite charge (Glu in Ube2L3, Asp in the Ube2D family, and Glu in Ube2B). Ube2E1 and Ube2E2 have 97% sequence identity and share common E3s but also have distinct E3 partners. The residue equivalent to Arg60 in Ube2T is predicted to be important for this distinction in a recent proteome-scale modeling study (Kar et al., 2012). Our study provides experimental support for the importance of this position in a more divergent E2. Arg60 of Ube2T forms a salt bridge with Glu340 of FANCL (Figure 1D). In other E3s, the equivalent residue is poorly conserved (Figure 2B).

Our structural and biochemical data can be used to refine algorithms for predicting E3-E2 pairs and provide a druggable platform for the development of chemotherapeutics.

**EXPERIMENTAL PROCEDURES**

**Protein Expression and Purification**

To generate the RING-linker-Ube2T fusion construct, we cloned human FANCL RING domain (residues 289–375) from synthetic human FANCL DNA (GeneArt) codon-optimized for Escherichia coli expression and inserted N terminally to Ube2T encoded in a vector containing a 6xHis-Smt3 tag, by restriction-free cloning (RF) (van den Ent and Löwe, 2006). A 14 amino acid (TGSTGSTETGYTQG) linker was inserted between the C-terminal of the RING domain and N-terminal of Ube2T by Phusion site-directed mutagenesis. Human Ube2T, Mouse Rbx1, and Xenopus tropicalis FANCL were cloned from I.M.A.G.E clones (Geneservice) into a vector containing a 6x His-Smt3 tag by RF methods. Human FANCL RING domain (residues 289–375) was cloned from the synthetic human FANCL DNA as described above. Human Ube2D3 (UbcH5c), Ube2L3 (UbcH7), Ube2K, Ube2B, and Ube2R1 were cloned from I.M.A.G.E clones (Geneservice) and inserted into the pDEST17 (Invitrogen) and pET RSF vectors containing an N-terminal 6xHis-Smt3 tag, by restriction-free cloning (RF) (van den Ent and Löwe, 2006). A 14 amino acid (TGSTGSTETGYTQG) linker was inserted between the C-terminal of the RING domain and N-terminal of Ube2T (Pellegrini et al., 2002) by Phusion site-directed mutagenesis. Human Ube2T, Mouse Rbx1, and Xenopus tropicalis FANCL were cloned from I.M.A.G.E clones (Geneservice) into a vector containing a 6xHis-Smt3 tag by RF methods. The Human FANCL RING domain (residues 289–375) was cloned from the synthetic human FANCL DNA as described above. Human Ube2D3 (UbcH5c), Ube2L3 (UbcH7), Ube2K, Ube2B, and Ube2R1 were cloned from I.M.A.G.E clones (Geneservice) and inserted into the pDEST17 (Invitrogen) and pET RSF vectors containing an N-terminal 6xHis tag and a TEV cleavage site. Ube2H was purchased as a synthetic gene in expression vector pJ441 containing a 6xHis tag and a TEV cleavage site from DNA 2.0. Proteins were expressed in E. coli BL21 cells (Invitrogen) or Rosetta (DE3) cells (Millipore) in the case of X. tropicalis FANCL, Ube2K, and Ube2R1. Cells were cultured in Lysogeny broth (LB) supplemented with antibiotics and 0.5 mM ZnCl2 for proteins with a RING

**Figure 3. Conserved Hydrophobic RING Residues Are Required for Ube2T Binding and FANCL Selects Solely Ube2T In Vitro**

(A) Size-exclusion chromatogram profiles of wild-type (WT) or mutant RING domains (green dashed line) and WT Ube2T (blue dotted line) overlaid with profiles from binding experiments in which WT Ube2T has been incubated with WT or mutant RING domains (pink line) and subjected to size-exclusion chromatography. Binding was assessed by complex formation, which is indicated by a peak shift to the left labeled complex. (B) Size-exclusion chromatogram of FANCL RING domain incubated with an E2 mix consisting of Ube2T, Ube2D3, and Ube2L3 (pink line). Chromatograms of Ube2T (blue dotted line) and the RING domain (green dashed line) are also overlaid. A peak shift to the left is observed, indicating complex formation. SDS-PAGE gel of the fractions collected from the size-exclusion experiment and stained with Coomassie Brilliant Blue. The E2 gel bands found in the shifted peak were assessed by mass spectrometry for protein identification and confirmed as exclusively Ube2T.

See also Figure S2.
domain, at 37°C. Once OD600 had reached 0.6, protein expression was induced by the addition of 250 µM isopropyl-1-thio-β-d-galactopyranoside (IPTG) (500 µM in the case of the E2s). Cells were cultured overnight at 16°C and harvested the following day by centrifugation. Harvested cells were lysed by sonication of 4x 10s bursts on ice, in buffer containing 0.5 M NaCl, 0.1 M Tris (pH 8), 0.02 M Imidazole, and 250 µM tris(carboxymethyl)phosphine (TCEP). Cell debris was removed by centrifugation at 32,000 g. Supernatants were added to equilibrated Ni-NTA Agarose (QIAGEN) and incubated on a roller for 1 hr at 4 °C. 6xHis-Smt3 tags were removed overnight at 4°C by Ulp1 protease at a w/w ratio of 1:15, Ulp1:protein. Xenopus tropicalis Ube2L3, Ube2K, Ube2B, Ube2R1, and Ube2H were eluted from agarose, concentrated, and applied to a Superdex 200 column. Purified His-UBE1 was then flash frozen and stored at −80°C.

**Xenopus laevis**

*Xenopus laevis* FANCD2 plasmid was a kind gift from P. Knipscheer and J. C. Walter. We modified it to contain a N-terminal FLAG tag and prepared FANCL plasmid as a kind gift from P. Knipscheer and K. Iwai (Sato et al., 2008). It was expressed in sf9 cells cultured at 27°C in sf-900 serum-free media (GIBCO) in sf9 cells cultured at 27°C in sf-900 serum-free media (GIBCO) supplemented with antibiotics. Cells were harvested 3 days postinfection and lysed by sonication, 2x 5 s bursts in buffer containing 0.05 M Tris (pH 8), 0.1 M NaCl, 0.5 mM PMSF, and 1 mM EDTA. Cell debris was removed by high-speed centrifugation, 32,000 x g, and supernatants were added to equilibrated Ni-NTA Agarose. The His-UBe1 was then eluted from the agarose, concentrated, and applied to a Superdex 200 column. Purified His-Ube1 was then flash frozen and stored at −80°C.

Human RING, Ube2T, and *Xenopus laevis* FANCD2 mutants were generated using site-directed mutagenesis and expressed as WT proteins.

**Crystallization and Structure Determination**

The RING-Ube2T fusion crystals were grown using a final concentration of 17.7 mg/ml by sitting drop vapor diffusion at 4°C in space group P4₃2₁2 with cell dimensions of a = 109.3 Å, b = 109.3 Å, c = 117.7 Å, α = 90°, β = 90°, and γ = 90°. Crystallization conditions were 1.6 M ammonium sulfate, 0.1 M NaCl, 0.1 M HEPES (pH 7.5). Crystals were cryo-protected with 20% glycerol and cryocooled in liquid nitrogen. Data were collected at Diamond Light Source on the microfocus I24 beamline at 0.96 Å wavelength. Data were processed using D*trek (Pflugrath, 1999), showing diffraction to 2.25 Å resolution. An estimated solvent content of 52.7% suggested two copies of the polypeptide chain in the asymmetric unit (ASU). Phases were generated by molecular replacement using the program Phaser (McCoy et al., 2007) with Ube2T (residues 1–154, PDB ID code 1YH2) and *Drosophila* FANCL (residues 312–371, 3K1L) as search models. The model was refined iteratively using phenix.refine (Cowsley and Cowtan, 2004, 2005) and manual model building using Coot (Emsley and Cowtan, 2004). Data were cut off to 2.4 Å resolution, and the last 400 images were omitted because of radiation damage. Omit maps were generated to check for model bias. The final model’s stereochemistry and geometry was checked with MolProbity (Davis et al., 2004) analysis (Table 1) and for favored regions of the Ramachandran plot (97.52%). Table 1 summarizes data collection and refinement statistics. The ASU contains four chains, two chains of Ube2T (A and B), and two chains of FANCL RING domain (X and Y). In both chains of Ube2T, electron density was not observed for the C-terminal (residues 154–197), and for chain B, the loop region from residue 27–32 was not modeled because of poor electron density. For chain Y of the RING domain, residues 352 and 353 were not modeled because of poor electron density. For chain Y of the RING domain, residues 352 and 353 were not modeled because of poor electron density. For chain Y of the RING domain, residues 352 and 353 were not modeled because of poor electron density. The interaction interface was calculated using PISA.

**Structural Analysis**

All structural analyses were carried out using PyMOL (Delano, 2002). Structural alignments were produced and manually adjusted using MegAlign software. The structural interface was calculated using PISA.
Analytical Size-Exclusion Chromatography

Interactions between RING domains and E2s were assessed by analytical size-exclusion chromatography, as described previously (Hodson et al., 2011). Briefly, interactions were incubated in buffer containing 0.1 M NaCl, 0.1 M Tris (pH 8), and 250 μM TCEP in a total volume of 500 μl and left on ice for 1 hr. Samples were loaded onto a Superdex 75 10/300 column (GE Healthcare), and 0.5 ml fractions were collected. Fractions were analyzed on 12% SDS-PAGE gels.

Native Gel Assays

Interactions between FANCL RING domain and E2s were assessed by native gel assays. Excess FANCL RING domain (~140 μM) was incubated with 80–100 μM of E2 and incubated in buffer containing 0.1 M NaCl, 0.1 M Tris (pH 8), and 250 μM TCEP in a total volume of 10 μl and left on ice for 10 min. A 10 μl of loading buffer containing 10% glycerol was added to the samples, and the samples were analyzed on 4%–12% native PAGE gels (Invitrogen). Gels were stained with SimplySafe Stain (Invitrogen).

FANCD2 Monoubiquitination Assay

The ability of FANCL to monoubiquitinate FANCD2 using different E2s was assessed by an in vitro monoubiquitination assay. Reaction volumes of 25 μl contained 17 nM His-UBE1, 0.64 μM HA-Ub (Boston Biochem), 0.5 μM E2, 1.86 mM MgCl₂, 0.5 mM DTT, and 2 mM ATP. Ube2D3, Ube2L6, and Ube2N/Ube2V1 were purchased from Boston Biochem. Reactions were left for 1.5 hr at room temperature. A 25 μl volume of LDS buffer (Invitrogen) containing BME was used to terminate reactions. Samples were loaded onto a 4%–12% SDS-PAGE gel and subjected to western blotting. Anti-HA antibody, raised against FLAG-FANCD2 or FLAG-FANCD2/Abcam, was used to probe for FLAG-FANCD2 monoubiquitination.

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