**Crystal Structure of a Complex of NOD1 CARD and Ubiquitin**

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**Abstract**

The Caspase Recruitment Domain (CARD) from the innate immune receptor NOD1 was crystallized with Ubiquitin (Ub). NOD1 CARD was present as a helix-swapped homodimer similar to other structures of NOD1 CARD, and Ub monomers formed a homodimer similar in conformation to Lys48-linked di-Ub. The interaction between NOD1 CARD and Ub in the crystal was mediated by novel binding sites on each molecule. Comparisons of these sites to previously identified interaction surfaces on both molecules were made along with discussion of their potential functional significance.

**Introduction**

NOD1 is an innate immune receptor that recognizes γ-D-glutamyl-meso-diaminopimelic acid (ε-DAP), a dipeptide present in peptidoglycan of bacteria, and activates inflammation and autophagy in response to bacterial infection [1,2]. Loss of NOD1 function increases susceptibility to severe systemic infections and is associated with a variety of inflammatory and autoimmune syndromes in humans [3–8]. NOD1 is composed of three domains: an N-terminal Caspase Recruitment Domain (CARD), a central nucleotide-binding oligomerization domain and a C-terminal ligand-binding domain [9]. Upon ligand binding, NOD1 oligomerizes into a signaling scaffold, which in turn recruits downstream effector molecules including kinases and ubiquitin ligases [10].

The N-terminal CARD of NOD1 has been shown to coordinate interactions with its various effectors. CARDs are part of a larger family of protein interaction motifs known as Death Domains (DD). In addition to CARDs, the DD superfamily includes Death Domains, Death Effector Domains and Pyrin Domains. These domains are typically around 100 amino acid residues in length and share a common six-helical bundle structural fold. They are involved in almost all aspects of immune and apoptotic signaling, and are found on various receptors, such as NOD1, as well as downstream signaling molecules such as RIP2, the effector kinase for NOD1 [11]. The majority of these domains have yet to be characterized structurally, and there are even fewer data describing how DDs complex with effector proteins. Like other DD subfamilies, CARDs were thought to engage almost exclusively in homotypic interactions with other CARDs [11]. Recently, however, Ub has been found to bind the CARDs of the innate immune receptors NOD1, NOD2, RIG-1 and MDA5 and regulate their activity [12–14].

Ub functions when covalently attached to substrate proteins as either a single mono-Ub or as a poly-Ub chain where distal Ubs are linked by their C-termini to one of seven lysine residue sidechains or the N-terminus of a proximal Ub moiety. Lys48-linked poly-Ub targets substrates to the proteasome for degradation, and this process plays an important role in immune signaling through destruction of 1κB, which is required to release the cytosol-bound NF-κB transcription factor to activate genes involved in the immune response [15]. In contrast, linear Met1-linked or Lys63-linked poly-Ub chains form platforms assembling Ub-binding kinases and other effectors that activate signaling pathways [reviewed by [16–19]]. In some pathways, Lys63-linked poly-Ub chains alone are sufficient to nucleate effector complexes and activate signaling [20].

The interactions between Ub and its effectors are complex and only a relatively small proportion of them have been characterized structurally. Here we report the crystal structure of a complex of NOD1 CARD and Ub. NOD1 CARD is present as a domain-swapped homodimer, similar to previously reported structures [21,22], while two Ub monomers interact in a conformation similar to that of Lys48-linked di-Ub [23]. We discuss the implications of this structure with regard to how NOD1 and other Ub-binding proteins may recognize Ub in vivo.

**Results**

Both subunits of the NOD1 CARD dimer interact with a single Ub moiety

The asymmetric unit of the crystal structure contains one NOD1 CARD and one Ub, while symmetry-related molecules reveal that both NOD1 CARD and Ub form homodimers (Figure 1A). As observed in two previous crystal structures, NOD1 CARD formed a homodimer that involved swapping of the sixth...
helix of each subunit [21,22]. Consistent with the previous structure determined by Coussens et al., the two Cys59 sidechains at the dimer interface in our structure remained reduced (Figure 1B). Superposition of the NOD1 CARD dimer determined in this structure with those previously determined shows close similarity with RMSD values of 1.21 Å and 1.79 Å over the Cα chain for 2NSN [22] and 2NZ7 [21], respectively.

During refinement, we noticed a large volume of electron density immediately adjacent to the sidechain of Cys59 on NOD1 CARD that could not be accounted for by a sulphhydril alone (Figure 1D). Given that NOD1 CARD was maintained in buffer containing 5 mM DTT from its initial purification, the likelihood of oxidative modification of the sulphhydril of Cys59 was quite low. DTT has been observed to form adducts with Cys sidechains [24], but in this case the density we observed was not sufficient to account for that particular addition. The buffer in the crystallization condition contained cacodylate, which has been shown to form arsenic (As) adducts with proteins [25–27]. In the presence of DTT, pentavalent cacodylate can be reduced to a highly reactive trivalent dimethyl-As-DTT intermediate. This intermediate reacts with surface exposed Cys residues on proteins, leading to addition of dimethyl-arsenic moieties to the sidechain sulphhydrils [28,29]. Thus, the components of the crystallization buffer strongly suggested the possibility of chemical modification of NOD1 CARD with dimethyl-As. Subsequent refinement of the model with a dimethyl-As moiety on Cys59 resulted in a good fit with the observed electron density (Figure 1B), and we found no evidence that either of the other two cysteine sidechains (Cys39 or Cys61) were similarly modified.

Ub forms a homodimer similar in conformation to Lys48-linked Di-Ub

Two Ubs were found in the unit cell as a homodimer in a conformation nearly identical to the structure of covalently linked Lys48 poly-Ub [23] despite the lack of any covalent bonds enforcing a particular conformation (Figure 2A). Previous studies have shown that pH can be a critical factor in determining which conformations Lys48-linked poly-Ub chains adopt, with acidic conditions favoring an open state and more neutral environments favoring a closed state, characterized by the hydrophobic patch of each Ub (comprised of Ile44, Val70 and Leu8) facing each other [23,30]. Additionally, the Ub dimer in our structure demonstrates the chelation of a multivalent anion by the sidechains of Arg42 and Arg72 in a fashion also observed for Lys48 linked poly-Ub [23] (Figure 2B,C). Since the proportion of basic residues to acidic ones in the interface is relatively high, chelation of a large anion through dimerization may serve to mitigate this electrostatic disparity. At lower pH, protonation of the few acidic residues, as well as anionic ligands, would be expected to disrupt this chelation through charge repulsion and favor an open state of the linked Ub moieties, explaining the pH-dependence of the conformational change.

Our crystals formed at a pH of ~8.6, significantly higher than the neutral pH values previously shown to favor inter-Ub interactions relative to acidic environments for Lys48-linked chains. Nonetheless, this closed conformation of Ub has not been observed in the absence of conformational constraint imposed by Lys48-Gly76 covalent bonds between Ub subunits, and it was a remarkable finding in our crystal structure. A recent non-covalent dimeric structure of Ub has been described which chelates cadmium [31]. However, the Ub homodimer in that structure does not match any other known poly-Ub conformation, and the effects of cadmium chelation may explain this idiosyncrasy.

NOD1 CARD binds a unique interface on Ub

The two common binding sites on Ub are the canonical hydrophilic pocket recognized by the majority of UBDs (which is partially buried in the Ub dimer interface in the closed Lys48-linked conformation – see Figure 1, Ile 44 black spheres) and the C-terminus, which is critical in the process of conjugation by ligases and cleavage by deubiquitinating enzymes. In our structure, NOD1 binds a different surface encompassing about 500 Å² that is centered on Phe4 (Figure 3A,B). Here, Thr66 is completely buried in the interface, and Lys63 and Glu64 make considerable contributions to binding NOD1 CARD. The sidechain of Glu64 on Ub, for instance, participates in six hydrogen bonds with four different residues on NOD1 CARD and also forms an edgewise intramolecular interaction with the aromatic ring of Phe4. This latter interaction is seen variably in other Ub structures (e.g. UEV domain of Vps23/TSG101:Ub (1UZX), CUE domain of Cue2:Ub (1OTR) and Lys48-linked tetra Ub (206V) display similar contacts, while CUE domain of Vps9p:Ub (1P3Q), mono-Ub (1UBQ) and ZnF UBP:Ub (2G45) do not), and it slightly alters the topology of the β-sheet on Ub by extending the β4/β5 loop at the expense of the fifth β-strand. In our structure, this may serve to better position Lys63 and Glu64 at the interface, as well as to orient the Phe4 aromatic sidechain for interaction with the modified Cys59 sidechain in the cleft between the second and third helices of NOD1 CARD.

Additional amphipathic residues from the first and second β-strands of Ub participate in the interactions, most notably the sidechains of Glu2, Thr14 and Glu16. Furthermore, the involvement of Lys6 extends the hydrophilic patch toward the traditional hydrophobic pocket, being in close proximity to the functionally important Leu8 residue. The remainder of the interface participates in relatively weak van der Waals interactions with NOD1 CARD. Interestingly, this surface of Ub overlaps with residues (e.g. Phe4) previously found to be functionally important for endocytosis [32], as well as with interaction sites used by other Ub-binding domains, including the UEV domain of TSG101/Vps23 [33], the NZF domain of HOIL-1L [34] and NEMO [35]; the first of these uses this patch to bind mono-Ub, while the latter two use the I44 hydrophobic patch simultaneously to bind linear poly-Ub (Figure 3C,D,E).

Ubiquitin binds a novel site on NOD1 CARD

The Ub interface on NOD1 CARD is localized primarily to an apical region of the molecule where the second, third and fourth helices converge (Figure 1, Figure 4A,B). The interaction is anchored in large part by the ζ1/ζ2 loop region and the fourth helix of NOD1 CARD, with the sidechains of Asn36, Thr37 and Glu64 hydrogen bonding with the sidechain of Glu64 from Ub. Glu64 of NOD1 CARD also hydrogen bonds with the functionally significant Lys63 sidechain of Ub. A second hub in the interface is centered on the third and fourth helices and the intervening loop region of NOD1 CARD, which bind to Thr66 of Ub. Two noteworthy contributions to the NOD1 CARD:Ub interaction arise from the third helix. First, Glu56 of NOD1 CARD is involved in a salt bridge with Lys6 of Ub, which is the only salt bridge found at the NOD1 CARD:Ub interface. Perhaps most interesting is the interaction between the sidechain of Phe4 of Ub and the chemically modified sidechain of Cys59 on NOD1 CARD (Figure 4).

Overall, the NOD1 CARD:Ub interface in our structure is small, and thus the affinity for the interaction would be expected to be weak. Indeed, in previously reported NMR solution studies of the interaction of NOD1 CARD with Ub, titrations of 15N-labeled NOD1 CARD with poly-Ub did not reveal an interaction surface.
Figure 1. Cartoon view of the NOD1 CARD-Ub complex. A) NOD1 CARD and Ub crystallized as a dimer of dimers. NOD1 CARD A (cyan) and Ub C (mauve taupe) are in the asymmetric unit. Symmetry-related subunits NOD1 CARD B (blue) and Ub D (yellow) complete the full complex. The NOD1 CARD homodimer demonstrates swapping of the sixth helices, similar to other reported NOD1 CARD dimers (2NSN and 2NZ7). The ubiquitin dimer chelates a phosphate anion (shown as spheres), similar to chelation of sulfate seen in another structure of Lys48-linked tetra-Ub (2O6V). Phe4 of Ub (green spheres) and the modified Cys59 of NOD1 CARD (orange spheres) are key residues mediating the NOD1 CARD-Ub interaction. The Lys48 residues of both Ub molecules are shown as magenta spheres. The N- and C- termini are highlighted as blue and red spheres, respectively. B) Close-up view of the dimethyl-arsenic adduct on NOD1 CARD. The Cys59 sidechain of NOD1 CARD (cyan) is shown in isolation. The adduct is within ~3.5 Å of the aromatic Phe4 sidechain on Ub (mauve taupe), also shown in isolation. Electron density contoured at 2.0 sigma and carved at 2.3 Å shows the Cys59 sidechain is modified (left), while the nearby solvent-exposed Cys61 is not.

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Figure 2. Mono-Ub in our structure formed dimers similar in conformation to other structures of K48-linked di-Ub. A) Ribbon representation of an alignment of K48-linked diubiquitin structures. The proximal di-Ub (2O6V-1, green) and distal di-Ub (2O6V-2, magenta) from tetra-Ub and di-Ub from 1AAR (blue) were aligned to the Ub dimer structure reported in this study (Ub C, mauve taupe and Ub D, yellow). Main chain RMSDs for this alignment were 1.309, 0.892 and 0.737 Å, respectively. B) Loop representation of the Ub dimer from our structure shown chelating a phosphate anion. This chelation is mediated mainly by Arg42 and Arg72 from each subunit. C) Loop representation of the proximal di-Ub moiety from K48-linked tetra-Ub (2O6V-1). In this case, Arg42 and Arg72 mediate chelation of a sulfate anion. In both structures, the Arg residues, which are located at the periphery of the hydrophobic pocket and often participate in interactions with UBDs, are engaged instead through these chelation interactions. This may serve as a mechanism by which UBDs discriminate between K48-linked poly-Ub and other forms of poly-Ub. K48 of the distal Ub moiety and the N-terminus (NT) of the proximal Ub moiety of each dimer are shown, demonstrating the close proximity in the dimer relative to the covalently linked chain.
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Figure 3. Identified UBD binding sites in Ub. Ub is colored in mauve taupe. A) Cartoon representation of Ub. Residues mediating interaction with NOD1 CARD are shown as sticks and are colored red. The canonical hydrophobic pocket residues are shown as sticks and colored black. B) Surface views of the NOD1 CARD interaction site on Ub. C) Surface views of the Phe4 hydrophobic patch (F4 patch) on Ub alone (green) and merged with the NOD1 CARD site (yellow). D) Surface views of one site on Ub recognized by the NZF domain of HOIL-1L alone (green) and merged with the NOD1 CARD site (yellow). This domain specifically recognizes linear chains of Ub by binding to the hydrophobic pocket of the distal Ub and to an area surrounding Phe4 of the proximal Ub of a linear di-Ub moiety. Residues on the proximal Ub interacting within 3.6 Å of the NZF domain in a crystal structure of its complex with linear di-Ub (PDB ID: 3B08) are shown. E) Surface views of a second site on Ub recognized by the Vps23 UEV domain (Vps23 site) alone (green) and merged with the NOD1 CARD site (yellow). The Vps23 UEV domain binds both the hydrophobic pocket and this second site on Ub. Residues on Ub interacting within 3.6 Å of the UEV domain in a crystal structure of its complex with mono-Ub (PDB ID: 1UZX) are shown.

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similar to that observed in our structure, even at millimolar concentrations [14]. The chemical shift perturbations on labeled NOD1 CARD caused by poly-Ub were relatively weak, however, which necessitated rather stringent cutoffs that may have masked more minor interactions. A notable overlap in Ub-binding sites between our structure and the previous study centers on Tyr88 of NOD1 CARD, which was reported as the most perturbed residue in the presence of poly-Ub and which blocked Ub-binding when mutated. While almost the entirety of the NOD1 CARD:Ub interaction in our structure is mediated by the NOD1 CARD A and Ub C moieties in the asymmetric unit (Figure 1), the sidechain of Tyr88 from the opposing dimer subunit (NOD1 CARD B) also interacts with Ub C (Figure 4A,B,C). Its sidechain hydroxyl initiates a hydrogen bond network that intersects sequentially through the sidechains of Glu16 and Gln2 of Ub to terminate at the ε-NH2 of Gln38 on the opposite NOD1 CARD subunit. Although this interaction only buries an additional 75 Å2 of surface area, this represents a 15% increase over the roughly 500 Å2 at the primary interface.

Discussion

Here we show the first crystal structure of the NOD1 CARD with Ub. The finding that Ub monomers crystallized as dimers in a similar conformation to Lys48-linked dimers is presented here as an interesting finding mostly because it reveals a new interface between Ub and NOD1 that we did not predict from our previous studies using NMR. Importantly, this structure does not necessarily support the idea that NOD1 CARD has specificity to Lys48-linked poly-Ub. Rather, the interaction between NOD1 and Ub can happen through multiple interfaces and that particular combinations may work together to mediate binding of different poly-Ub chain topologies. It is tempting to speculate that NOD1 interactions with different poly-Ub chains could result in distinct conformations of the NOD1:Ub complex that could mediate distinct biological effects. In general, UBDs that show specificity for particular poly-Ub chains are composed of two binding surfaces arranged so that two Ubs in a chain have simultaneous access to these sites. Often, this involves binding of one Ub via its Leu8, Ile44, V70 hydrophobic surface combined with binding to a second Ub moiety through a similar or distinct surface. Although this second surface of Ub can be quite small and have negligible affinity on its own, it can contribute substantially to the overall strength of binding in the context of a poly-Ub chain. The interaction we found in the crystal could represent such an auxiliary binding site and might provide a mechanism to explain how NOD1 CARD favors binding to Lys63 and linear poly-Ub. This model would also be consistent and complementary to our previous studies examining how the NOD1 CARD binds monomeric Ub in solution through surfaces in both the CARD and Ub that are distinct from those identified in the current crystal structure [14].

Our previous NMR studies mapped the major binding interface of NOD1 onto the canonical hydrophobic patch of Ub centered on Ile44, Val107 and Leu8, which in the current structure is used to mediate binding to another Ub molecule. Mutagenesis experiments in previous work showed that mutations of residues in NOD1 at this interface dramatically diminished binding of Ub, supporting the hypothesis that this previously determined binding mode was valid. Our previous data and the crystal structure presented here are best reconciled with Ub binding in two modes: a major mode which is required for measurable binding and which dominates in solution where monomeric NOD1 and monomeric Ub exist; and a minor mode, which was captured here in the crystal structure. These two interfaces could work together to impart better binding to poly-Ub or to guide specificity for binding poly-Ub chains linked via specific linkers.

The alternative interface we discover in the crystal structure could potentially mediate di-Ub binding even for Ubs linked by chains other than Lys48, such as Lys63 and linear linked Ub chains that are known to have important signal transduction roles in innate immunity. The recent structure of the HOIL-1L NZF domain bound to linear di-Ub [34].

Figure 5. Cartoon view of a model for how monomeric NOD1 CARD could bind linear di-Ub generated by the ZDOCK server (see Materials and Methods). The sidechains of Phe4 (purple) and Ile44 (black) from Ub and Cys59 (red) and Tyr88 (green) are shown as spheres. The combination of the previously identified NMR Ub-binding site (Tyr88:Ile44) with the new Ub-binding site observed in our crystal structure (Cys59:Phe4) may explain the mechanism by which NOD1 CARD preferentially binds linear and Lys63-linked poly-Ub. doi:10.1371/journal.pone.0104017.g005
domain in a complex with poly-Ub is instructive with regard to our proposed model for NOD1 binding. HOIL-1L NZF binds two surfaces on Ub, one centered on the Phe4 patch and the other on the Ile44 hydrophobic pocket [34]. These two small surface patches of ~400 Å² each combine to yield a much greater affinity for poly-Ub chains when the NZF domain engages both as it "wedges" between two Ub moieties in a linear poly-Ub chain. Perhaps CARDs similarly engage two Ub surfaces, which would explain how both NOD1 CARD and RIG-I CARD [13] prefer binding to poly-Ub chains. Such a model is shown in Figure 5. Here, docking experiments were used to rationalize both the binding surface on Ub described here and the ability of NOD1 to bind Ub or other Ub-interacting proteins remains for future studies.

Materials and Methods

Protein purification

The CARD of human NOD1, from residues serine 16 to glycine 108, was produced in a pET-21a vector (Novagen). Amino acids leucine and glutamic acid were inserted between glycine 108 and the 6xHis-tag, due to the presence of an XhoI restriction site. The resulting plasmid was used to transform Escherichia coli strain BL21(DE3) (Invitrogen), which was grown at 310 K in LB medium, supplemented with ampicillin (100 μg/mL). At mid-log phase (OD₆₀₀ = 0.5), overexpression of NOD1 CARD was induced with 1 mM isopropyl-β-thiogalactoside (Research Products International Corp.). The culture was harvested 4 hours after induction by centrifugation for 30 min at 7280 g and 277 K. The cells were resuspended in 45 mL of buffer A (50 mM sodium phosphate, 150 mM NaCl, and 5 mM imidazole, pH 7.0) per 1 L of culture, and lysed using a French press at 16,000 psi. The lysate was spun for 45 minutes at 15,000 g and 277 K to separate the soluble and insoluble fractions. The resulting supernatant was separated from the pellet and mixed with TALON metal affinity resin (BD Biosciences) pre-equilibrated in buffer A. The mixture was allowed to incubate on a rotating platform at 277 K for 2 hours, followed by addition of the slurry to a gravity flow column. The supernatant was allowed to flow through and the resin was washed with 30 batch volumes of buffer A. The NOD1 CARD protein was eluted in 50 mM sodium phosphate, 150 mM NaCl, and 500 mM imidazole, pH 7.0. Eluted protein was concentrated and further purified by gel filtration over a preparative Superdex 75 column (GE Healthcare, Piscataway, NJ) equilibrated with GF buffer (25 mM sodium phosphate, 25 mM NaCl and 5 mM DTT, pH 7.0). Two sequential rounds

An additional aspect of this interface is that it includes two residues of NOD1 (Asn36 and Gln56) that are highly conserved across evolution, supporting the idea that it may be used for important protein interactions such as that of Ub described here [36]. Moreover, mutation of Gln56 compromises the ability of NOD1 to activate its downstream effector kinase RIP2 [37], through a yet unknown mechanism. This interface also includes Ser65 of Ub, which was recently found to undergo phosphorylation by PINK to create a form of Ub that could act allosterically to activate the Parkin Ub ligase [38–40]. Thus, it may be that the binding surface on Ub described here could operate for other Ub interactions and have the capacity to be regulated.

As in any crystal structure, there exists the possibility that the unique milieu of the crystal itself produced an artificial interaction. Although this binding interaction was not definitively observed in our prior NMR experiments, it was not definitively excluded either, given the low signal to noise of experiments performed in the range of low binding affinities expected for this interaction. Bosanac et al [41] reported a crystal structure of the zinc-finger domain of the protein A20 bound to three separate mono-Ubs with each site having progressively lower binding affinities. Our structure may represent even further progression to beyond the limit of detection by NMR or traditional pull-downs, where the majority of the binding events in solution represent the dominant interaction previously identified. Additionally, the exact binding mode and stoichiometry of protein complexes is not always in agreement between rigorous studies by NMR vs crystallography, as exemplified by the studies of the NEMO:linear di-Ub interaction [35,42].

The biological relevance of this new Ub binding surface and the ability of NOD1 to bind Ub or other Ub-interacting proteins remains for future studies.

Table 1. X-ray data collection and refinement statistics.

| Data Collection                         | Value            |
|-----------------------------------------|------------------|
| Wavelength (Å)                          | 1.000            |
| Space group                             | P4₁2₂2           |
| Cell dimensions (Å)                     | a = b = 61.69, c = 86.87 |
| Cell Angles (°)                         | α = β = 90       |
| Resolution range (Å)                    | 38.98 – 2.90     |
| Total reflections                       | 54090            |
| Unique reflections                      | 4048 (395)       |
| Completeness (%)                        | 99.88 (100.0)    |
| Multiplicity                            | 13.35 (13.92)    |
| Rmerge                                  | 0.05 (0.21)      |
| Rfree                                   | 0.270            |
| R.M.S.D.                                |                  |
| Bond lengths (Å)                        | 0.003            |
| Bond angles (°)                         | 0.704            |
| No. protein atoms                       |                  |
| Chain A                                 | 751              |
| Chain B                                 | 582              |
| No. ligand atoms                        | 5                |
| Average B factor (Å²)                   | 57.40            |
| Ramachandran favored (%)                | 98               |
| Ramachandran outliers (%)               | 1.2              |

Values for the highest resolution shell are given in parentheses.

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NOD1-UQ Complex Structure

The biological relevance of this new Ub binding surface and the ability of NOD1 to bind Ub or other Ub-interacting proteins remains for future studies.
of gel filtration were performed to remove residual NOD1 CARD dimer and to ensure homogeneity. Ubiquitin was purified as previously described [43], with an additional step of gel filtration over a Superdex 75 column equilibrated in GF buffer.

Crystallization

Purified NOD1 CARD and Ub were separately concentrated to 2 mM, which was determined by absorbance at 290 nm using respective extinction coefficients of 9970 and 1280 M\(^{-1}\)cm\(^{-1}\). The separate proteins were mixed at a ratio of 1:1 (based on prior experiments suggesting 1:1 binding stoichiometry [14]) and incubated on ice for an hour to form complexes at a final concentration of 1 mM for each component. 2 mM complexes were created by two-fold concentration of 1 mM complexes after mixing. Complexes at concentrations less than 1 mM were made by diluting 1 mM complexes in appropriate volumes of GF buffer. Initial 96-well crystal screens utilizing the hanging drop/vapor diffusion method were set up for complexes at 2 mM or 1 mM using a TTP LabTech mosquito robot (Cambridge, MA) set to dispense 400 nL of protein solution plus 400 nL of well solution for each condition. Duplicate screens were set up and one each was left at either 277 or 291 K. Hits were obtained in PACT Suite (Qiagen) conditions \#29 and \#30. These contained 100 mM PCMB (sodium propionate, sodium cacodylate, bis-tris propane mixed at a molar ratio of 2:1:2 and pH adjusted with HCl – as described by the manufacturer) and 25% PEG 1500 at pH 8.0 or 9.0, respectively. No additional DTT was added to the drops. Crystals grew overnight at both 277 and 291 K, although growth was left at either 277 or 291 K. Hits were obtained in PACT Suite for each condition. Duplicate screens were set up and one each was left at either 277 or 291 K. Hits were obtained in PACT Suite for complexes at 2 mM or 1 mM incubated on ice for an hour to form complexes at a final concentration of 1 mM for each component. 2 mM complexes were made by diluting 1 mM complexes in appropriate volumes of GF buffer.

Data collection

Looped crystals were flash cooled in liquid \(N\)\(_2\). Several cryoprotectants were able to reduce ice formation, but also resulted in poorer diffraction. Ultimately, crystals used for data collection were flash frozen without additional cryoprotectant and screened for the best combination of minimal ice and maximal diffraction on a rotating anode (RU-H3R) Rigaku generator (1.54 Å) home X-ray source. Full datasets were collected on the best crystals on beamline 4.2.2 at ALS, Berkeley, California.

Data processing, structural calculations and refinement

Diffraction datasets were processed using d\(_{\text{TREK}}\) (Rigaku) [44]. Phases were determined by molecular replacement with NOD1 CARD (PDB ID 2NSN) and Ub (PDB ID 1UQB) as the targets using Phaser [45]. Refinement and model building were done using Phenix [46] and Coot [47], respectively. Processing statistics are summarized in Table 1. Figures were produced using Pymol [48].

Accession numbers

The structure for the NOD1 CARD:Ub complex was deposited in the Protein Data Bank under ID code 4JQW.

Docking calculations

To model the proposed similarity of the HOIL-1L NZF interaction with linear di-Ub to how NOD1 CARD might also bind linear poly-Ub, NOD1 CARD monomer (PDB ID: 2DBD) was paired with linear di-Ub from the HOIL-1L NZF:di-Ub crystal structure (PDB ID: 3B08) to predict possible complexes using the ZDOCK server [49,50]. Results were filtered by the criteria that Tyr88 of NOD1 CARD interacted with Le44 of the distal Ub moiety (NMR site) and Cys59 of NOD1 CARD interacted with Phe4 of the proximal Ub moiety (crystal site).

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Author Contributions

Conceived and designed the experiments: AVH RCP RS. Performed the experiments: AVH LG. Analyzed the data: AVH LG RS. Contributed reagents/materials/analysis tools: RCP LG. Wrote the paper: AVH LG RCP RS.

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