ASC Pyrin Domain Self-associates and Binds NLRP3 Protein Using Equivalent Binding Interfaces

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Javier Oroz, Susana Barrera-Vilarmau, Carlos Alfonso, Germán Rivas, and Eva de Alba

From the Centro de Investigaciones Biológicas, Departments of Chemical and Physical Biology and Cellular and Molecular Biology, and Centro Nacional de Biotecnología, Consejo Superior de Investigaciones Científicas, Ramiro de Maeta, 9 Madrid-28040, Spain, the German Center for Neurodegenerative Diseases (DZNE), Max Planck Institute for Biophysical Chemistry, Am Fassberg 11, Göttingen-37077, Germany, and the Health Sciences Research Institute, University of California at Merced, Merced, California 95343

Death domain superfamily members typically act as adaptors mediating in the assembly of supramolecular complexes with critical apoptosis and inflammation functions. These modular proteins consist of death domains, death effector domains, caspase recruitment domains, and pyrin domains (PYD). Despite the high structural similarity among them, only homotypic interactions participate in complex formation, suggesting that subtle factors differentiate each interaction type. It is thus critical to identify these factors as an essential step toward the understanding of the molecular basis of apoptosis and inflammation. The proteins apoptosis-associated speck-like protein containing a CARD (ASC) and NLRP3 play key roles in the regulation of apoptosis and inflammation through self-association and protein-protein interactions mediated by their PYDs. To better understand the molecular basis of their function, we have characterized ASC and NLRP3 PYD self-association and their intermolecular interaction by solution NMR spectroscopy and analytical ultracentrifugation. We found that ASC self-associates and binds NLRP3 PYD through equivalent protein regions, with higher binding affinity for the latter. These regions are located at opposite sides of the protein allowing multimeric complex formation previously shown in ASC PYD fibril assemblies. We show that NLRP3 PYD coexists in solution as a monomer and highly populated large-order oligomerized species. Despite this, we determined its monomeric three-dimensional solution structure by NMR and characterized its binding to ASC PYD. Using our novel structural data, we propose molecular models of ASC-ASC and ASC-NLRP3 PYD early supramolecular complexes, providing new insights into the molecular mechanisms of inflammasome and apoptosis signaling.

Proteins of the death domain superfamily mediate in apoptotic, innate immune, and inflammatory responses mainly through homotypic interactions (1). Specifically, they are responsible for the assembly of large signaling complexes in which kinases and caspases are activated (1–4). Despite the lack of sequence convergence within the superfamily, its members are characterized by adopting the so-called death fold, consisting of a globular fold with six α-helices arranged in a greek-key topology (1, 3). However, some structural differences between the subfamilies are observed within this common fold, including different charge distributions throughout the protein surface, diverse locations of hydrophobic patches, and the different lengths of certain α-helices. These differences are thought to lead to homotypic binding specificity (5), albeit certain heterotypic interactions are also known with proposed inhibitory roles in apoptotic and inflammatory pathways (1, 6).

Detailed structural information on complexes formed by several members of the death domain superfamily is available. Some examples are the death domain interactions in the PID-Dosome (7), the MyDDosome (8), the Fas-FADD (9, 10), and the Pelle-Tube death domain complexes (11), among others. In addition, structural studies on CARD-CARD interactions have been reported for the Caenorhabditis elegans apoptosis (12), the procaspase-9-Apaf-1 (13), and CED-4-CED-9 complexes (14), together with the death effector domain-containing protein of the MC159 tandem complex (15). In contrast, little information is currently available on the interactions established by PYD domains (6, 16–19). This kind of information is key for designing strategies targeted to modulate complex formation and thus develop therapeutic applications in the context of inflammatory and apoptosis-related disorders (2, 20, 21).

Only three different types of death domain interactions responsible for the formation of multimeric complexes have
been characterized thus far (18, 22). The type I interaction defines contacts between helices 1 and 4 of one death domain and helices 2 and 3 of the other; the type II interaction occurs when helix 4 and the loop connecting helices 4 and 5 of one domain interact with the loop between helices 5 and 6 of the other; and finally, the type III interaction includes contacts between helix 3 of one domain and the loops linking helices 1 and 2 and helices 3 and 4 of the other domain (3). It is noteworthy that these binding interfaces do not overlap, and thus it is possible for a single death domain to establish multiple interactions (7, 18). It is assumed that all death domain subfamilies are capable of forming these three interaction types. However, important differences present in each type are believed to be responsible for the homotypic specificity (18).

The apoptosis-associated speck-like protein containing a CARD (ASC) and NLRP3 (also known as cryopyrin) are two members of the PYD subfamily that play crucial roles in inflammasome activation. Upon the onset of danger signals, ASC and NLRP3 activate the inflammasome leading to procaspase-1 activation and further proteolytic activation of the pleiotropic inflammatory cytokines IL-1β and IL-18 (23). This process will ultimately prompt NF-κB activation thus inducing proinflammatory responses (1, 2, 24–27). ASC contains a PYD and a CARD domain (28, 29), whereas NLRP3 carries a PYD, a nucleotide-binding oligomerization domain, and a leucine-rich repeat domain (1). It is assumed that both proteins interact through their PYD domains, whereas ASC and caspase-1 binding would be mediated by a CARD-CARD interaction, subsequently leading to a highly multimeric complex formation, which is essential for the proteolytic activation of the initiator caspases (30, 31).

The PYD domain is characterized at the structural level by a large loop connecting helices 2 and 3, with the latter being particularly short (6, 17, 32–38). Despite the abundant structural information reported for PYD domains (the three-dimensional structures of 10 different PYD domains have been deposited in the PDB) (6, 17, 29, 32–38), our understanding on the ASC-NLRP3 PYD homotypic interaction is very limited. Several models for this interaction have been proposed based on the NLRP3 crystallographic structure and mutational data (4, 16), in addition to the recent semi-atomic structure characterization of ASC PYD mature fibril assemblies by cryo-EM and solid-state NMR (18, 39).

We report here the three-dimensional solution structure of NLRP3 PYD and a detailed study of ASC PYD self-association and its interaction with NLRP3 by NMR and analytical ultracentrifugation (AUC) using different conditions of pH and salinity. All in all, our data indicate that the ASC-ASC and ASC-NLRP3 complexes share the same binding interface. However, we observed higher affinity for the heteromeric interaction. Furthermore, using the available atomic structures and our structural results on the interactions, we have built models of the supramolecular complexes for both ASC-ASC an ASC-NLRP3 PYDs, which provide molecular insights into the first possible stages of inflammasome formation. In addition, our data could be used to target ASC and NLRP3 proteins in the design of therapeutic agents for the treatment of diseases associated with inflammasome disjunction.

### Results and Discussion

ASC PYD self-associates through Two Opposing Binding Interfaces with Similar Affinities—ASC self-association is an essential step in the formation of macromolecular assemblies like the pyroptosome and the inflammasome (40), which have critical roles in cell death and inflammation. NMR or x-ray crystallographic studies are necessary to characterize these interactions at the atomic level. However, oligomerization poses significant challenges for both techniques. It is well known that ASC PYD solubility is too low at neutral pH for conducting NMR experiments (29, 32). Therefore, previous NMR attempts to characterize ASC PYD interactions used an L25A mutant that is soluble at the typical NMR concentrations (27, 37). Unfortunately, this mutation has been shown to diminish the binding capabilities of the protein (see below). Remarkably, recent advances in cryo-EM imaging and solid-state NMR enabled the reconstitution at quasi-atomic resolution of ASC PYD inside a fibril, providing novel structural details into the homo-oligomerization mechanism of ASC (18, 39). However, these groundbreaking structural reconstitutions only provide detailed characterization on mature fibrils, leaving the initial stages of aggregation still unrevealed.

To improve our limited understanding on PYD death domain interaction, we have monitored ASC PYD self-association by NMR under conditions that help increase protein solubility. The chemical shift of NMR signals in 1H-15N HMQC spectra of native 15N-labeled ASC PYD varies upon the addition of unlabeled ASC PYD at different concentrations (Fig. 1A) until reaching a plateau (Fig. 2A), indicating that the protein is self-associating under these conditions. Once the titration reaches an end, the resulting spectrum overlies that recorded at high 15N-labeled ASC PYD concentration (Fig. 1A). The presence of a single set of signals during titration indicates that ASC PYD self-association is fast on the NMR chemical shift time scale. Residues with the largest chemical shift changes are shown in Fig. 1B and appear highlighted in the structure of ASC PYD (PDB code 2KN6 (29)) in Fig. 1D. Our results indicate that self-association is mediated by two interacting surfaces, one involving helices H1, H4, and H5 N terminus and the other composed of helices H2, H3, and H5 C terminus (Fig. 1D).

In general terms, this finding agrees with previous mutational studies on ASC PYD oligomerization (16, 41). However, several mutations of solvent-exposed residues that abolished ASC PYD self-association do not show, according to our data, the largest variations in chemical shift upon interaction (like Glu-13, Lys-21, and Asp-48, Fig. 1B). This discrepancy could be explained if the absence of the interaction results from an overall destabilization of the protein structure caused by the mutations, rather than from a perturbation of the interaction interface in the complex. Moreover, in one mutational study, the Thr-63A mutant does not seem to significantly affect ASC self-association (16). However, given that Thr-63 is located at the boundaries of the H1, H4, and N-terminal H5 interface (Fig. 1D), it is likely that its contribution to the interaction is minor.

Electrostatic interactions have been traditionally considered as carrying most of the weight with PYD-PYD complex formation (16, 18, 37, 41). As a result, information on the importance...
of apolar or hydrophobic interactions is scarce. Our NMR titration data show several hydrophobic residues with some of the largest chemical shift perturbations, including Ala-4, Leu-25, Val-30, Leu-44, Leu-45, and Leu-68. Among them, Leu-25, Val-30, and Leu-45 are sufficiently exposed to participate in side chain-side chain interactions. Interestingly, these residues are all located in the interface formed by H2 and H3. Our results thus indicate that hydrophobic interactions play a critical role in one of the two binding surfaces, in agreement with mutational studies pointing to an important function for Leu-25 and Leu-45 in ASC PYD filament formation (41). On the contrary, the V30A mutant can still form filaments, probably because Val-30 is located at the boundaries of H2-H3 interface, whereas Leu-25 and Leu-45 show a central position in this interface (Fig. 1D). Besides, a significant number of the residues showing large chemical shift perturbations have some degree of polarity (Thr-16, Ser-46, Thr-63, Tyr-64, Met-76, and His-90), which indicates that polar interactions are playing an important role in self-association.

However, binding affinity data on ASC PYD self-association is so far unknown, probably due to the previously mentioned aggregation propensity of ASC PYD at neutral pH (29). NMR is an optimum technique to obtain binding affinity constants that fall within the micromolar to millimolar range. We have selected some of the residues with the largest chemical shift variations to obtain apparent values of the ASC PYD dissociation constant (Fig. 2A). Our results indicate that both surfaces in ASC PYD (H1, H4, N-term H5 and H2, H3, C-term H5) self-associate with similar affinity showing $K_D$ values in the micromolar range ($\sim 40$ to $\sim 100 \mu M$).
Solution Structure and Binding Properties of NLRP3 PYD

FIGURE 2. Dissociation constant \( (K_d) \) values indicate preference of ASC PYD for NLRP3 versus self-association. Changes in chemical shift versus protein concentration for the ASC-ASC (A) and the ASC-NLRP3 (B) interaction. \( K_d \) values for the ASC-ASC interaction are as follows: Arg-5 (98 ± 14 \( \mu \text{M (r = 0.99) \); Asp-10 (43 ± 8 \( \mu \text{M (r = 0.98) \); Thr-16 (41 ± 8 \( \mu \text{M (r = 0.98) \); Glu-19 (63 ± 12 \( \mu \text{M (r = 0.98) \); Asp-54 (38 ± 10 \( \mu \text{M (r = 0.97) \); Thr-63 (106 ± 11 \( \mu \text{M (r = 0.99) \); Glu-67 (65 ± 25 \( \mu \text{M (r = 0.93) \); Asp-75 (74 ± 11 \( \mu \text{M (r = 0.99) \); and His-90 (56 ± 15 \( \mu \text{M (r = 0.96) \). \( K_d \) values for the ASC-NLRP3 interaction are as follows: Arg-5 (42 ± 17 \( \mu \text{M (r = 0.98) \); Asp-10 (37 ± 36 \( \mu \text{M (r = 0.92) \); Thr-16 (3 ± 1 \( \mu \text{M (r = 0.99) \); Glu-19 (2 ± 3 \( \mu \text{M (r = 0.96) \); Asp-54 (8 ± 7 \( \mu \text{M (r = 0.95) \); Thr-63 (101 ± 58 \( \mu \text{M (r = 0.97) \); Glu-67 (55 ± 20 \( \mu \text{M (r = 0.99) \; Asp-75 (95 ± 85 \( \mu \text{M (r = 0.94) \; and His-90 (71 ± 47 \( \mu \text{M (r = 0.96) \). Note that the fittings show poor quality for ASC-NLRP3 because of the lower ratio reached (1:3).

However, because the observed chemical shift perturbations are an average of the species resulting from the self-association process, they do not provide the association state of ASC. To gain insight into ASC PYD aggregation state, AUC experiments were performed. The protein appears mainly monomeric in the range of concentrations tested (50–750 \( \mu \text{M}, \text{Fig. 3A}) however, AUC data indicate a tendency of the protein to oligomerize upon increasing concentrations, in agreement with the NMR observations. Nevertheless, the increase in the association state is very small precluding the determination of the oligomeric state formed.

Solution Structure and Oligomerization Properties of NLRP3 PYD—The conformational behavior and self-associating capabilities of NLRP3 PYD are not very well understood. Previous work using multianaly light scattering coupled to size exclusion chromatography indicates that NLRP3 PYD, although mainly monomeric, can form dimers in solution at pH 5 and 150 mM NaCl (36). However, the effect of protein concentration was not analyzed. Here, we have used sedimentation velocity experiments to study the effect of salt and protein concentration on NLRP3 PYD aggregation (Figs. 3B and 4). We found that in the absence of NaCl, NLRP3 PYD is mainly monomeric at low concentrations (peak at ~1.5 S; Fig. 3B). At higher protein concentrations, the population of monomeric species decreases, and several higher order oligomerized species are observed. In the presence of NaCl, the population of monomeric species drastically decreases even at the lower protein concentration (Fig. 4), which indicates that the presence of salt favors self-association and could point to an important role of hydrophobic interactions in the NLRP3 PYD aggregation process.

Results from the sedimentation velocity experiments agree with the solution behavior of NLRP3 PYD observed by NMR. At 0.2 mM protein concentration (pH 3.6, 303 K) in the absence of salt, the NMR \( ^1\text{H}-\text{H}^{15}\text{N} \) HSQC spectra of NLRP3 PYD shows at least two sets of signals (Fig. 5A) as follows: one set clearly corresponds to the monomer according to the NMR signal line-width, and the other set results from soluble aggregated species. The presence of different sets of signals indicates that monomer-oligomer formation is slow on the NMR chemical shift time scale. It is noteworthy that on the basis of the sedimentation velocity experiments (Fig. 3B), less than 20% of mono-
meric species are present at 0.2 mM protein concentration, rendering the NMR study with significant challenges. Moreover, the intensity of monomeric NMR signals gradually decreases with time (Fig. 5B), and signals disappear at higher concentration (Fig. 5C). Temperature increase (Fig. 5D) and the presence of NaCl (Fig. 5E) also promote aggregation, in line with sedimentation data (Fig. 4).

Despite the mentioned challenges, the solution structure of NLRP3 PYD was calculated (Fig. 6). It consists of the typical six-helix bundle motif, with a long loop connecting helices 2 and 3, with the latter being the shortest out of the six helices. The backbone r.m.s.d. of the solution ensemble of conformers, excluding and including loops, is 0.49 and 0.71 Å, respectively. These results indicate the high convergence of the NMR data. MolProbity structural check indicates a 65th percentile of clash score and a 5.13% of poor rotamers. PROCHECK-NMR program (42) shows that the NMR ensemble has 91.9% of the residues in the most favored regions of the Ramachandran plot, 7.8% in additionally allowed regions, 0.3% in generously allowed regions, and 0% in disallowed regions (supplemental Table S1).

Overall, the NLRP3 PYD structure is similar to other PYD domains (Fig. 7A), although by superimposing the different structures, we found that it is more similar to NLRP4 and NLRP10 than to NLRP1, NLRP7, or NLRP12 (6, 32, 35, 37, 38). Strikingly, those NLRPs showing higher structural similarities to NLRP3 (namely NLRP4 and NLRP10) contain an opposing inhibitory role of the inflammatory response, whereas those showing higher structural divergence contain a similar regulatory role (43). This points to the relevance of slight structural divergences determining the specific functions of NRLPs.

Moreover, the crystallographic (36) and solution structures of NLRP3 PYD share a high degree of resemblance, with a backbone r.m.s.d. value of 1.66 Å (Fig. 7B). The arrangement of helices 2 and 3 (both involved in the dimer interface of the
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**FIGURE 7. Comparison of NLRP3 PYD with other pyrin domains.** A, superposition of the known three-dimensional solution structures of NLRP pyrin domains to NLRP3 PYD represented as ribbon diagrams (PDB codes: NLRP3, 2NAQ; NLRP1, 1PN5; NLRP4, 4EWI; NLRP7, 2KM6; NLRP10, 2DO9; and NLRP12, 2I6A). Blue (NLRP3), magenta (NLRP1), ivory (NLRP4), cyan (NLRP7), orange (NLRP10), and pink (NLRP12) are shown. Helices are numbered. Superimposed residues belong to helices 1, 2, and 4–6. Helix 3 (absent in NLRP1, NLRP10, and NLRP12) and the loop connecting helices 2 and 3 were excluded because of significant conformational variability. The r.m.s.d. values between NLRP3 and the other PYDs are as follows: NLRP4, 3.9 Å; NLRP10, 3.4 Å; NLRP1, 4.9 Å; NLRP7, 5.2 Å; and NLRP12, 5.9 Å. B, superposition of the crystallographic dimeric structure of NLRP3 PYD (green, PDB code, 3QF2; Ref. 36) onto the monomeric NMR structure (blue). The image at the right side shows a detailed view of the dimeric interface, including side chain orientation. The image was displayed with MOLMOL (65).

crystallographic structure) and helix 4 shows the largest differences between both crystal and solution structures. In contrast, side chain orientation of residues in close contact within the dimer interface of the crystallographic structure is very similar both in the NMR monomeric structure and the crystallographic dimer (Fig. 7B). This is a striking result as side chain intermolecular interactions likely contribute to the formation of the dimer, and thus side chain orientation is expected to be different in the monomeric solution structure. Besides, several confronted residues in the dimeric crystallographic interface are charged and therefore should induce repulsive interactions (Fig. 7B). In addition, as mentioned above, NLRP3 PYD is mainly monomeric at the concentrations used for crystallization (36). Therefore, the evidence indicates that the dimeric structure could result from a crystal-packing artifact, as suggested previously (36, 37).

ASC PYD Interacts with NLRP3 PYD through the Same Binding Sites Involved in Its Self-association Albeit with Higher Affinity—After studying the oligomerization capabilities and conformational behavior of NLRP3 in solution, we attempted to analyze a possible interaction between ASC PYD and NLRP3 PYD in solution. For this purpose, $^1$H–$^{15}$N HMQC spectra of a $^{15}$N-enriched ASC PYD sample were recorded upon the addition of increasing amounts of unlabeled NLRP3 PYD. Chemical shift changes of ASC NMR signals show binding of both PYDs (Fig. 1, A, C, and E), indicating that ASC PYD self-association and its interaction with NLRP3 PYD are mediated by almost identical binding regions. This evidence agrees with a previous study showing that upon mutation of critical residues in the ASC PYD binding interfaces described, both ASC:ASC and ASC-NLRP3 PYD interactions were identical diminished (16). Moreover, the influence of hydrophobic interactions in ASC-NLRP3 PYD binding is again highlighted in residues Leu-25, Val-30, and Leu-45 (Fig. 1C).

In this case, it was also possible to measure the dissociation constants from the NMR titration experiments, which fall within the micromolar range (Fig. 2B). However, the determination of these dissociation constants is somewhat less accurate than for the ASC-ASC PYD binding because NLRP3 increased the tendency for aggregation, which precluded reaching the plateau. Nevertheless, several accurate values were obtained from the change in chemical shifts of Arg-5, Thr-16, Glu-19, Asp-54, and Glu-67 (Fig. 2B). The average value for these residues is 22 $\mu$M, which indicates that the affinity of the ASC-NLRP3 PYD complex is slightly higher. A comparable dissociation constant was reported for the ASC-POP1 PYD interaction ($\sim$ 4 $\mu$M; 27). However, no analysis on the oligomeric state of ASC PYD was shown in that study.

**ASC PYD Self-association and Interaction with NLRP3 Are Enhanced by Increasing the pH and NaCl Concentration of the Milieu**—Because we have already shown both by NMR and AUC that the presence of salt enhances NLRP3 PYD oligomerization, we also analyzed its potential effect on ASC PYD self-association and its interaction with NLRP3 PYD. In the presence of 100 mM NaCl, the $^1$H–$^{15}$N HMQC spectrum of ASC shows characteristic patterns corresponding to the ASC:ASC interaction, as if the protein had already oligomerized (Fig. 8A). Analogously, the first spectra obtained during the titrations in the present of salt are more similar to the spectra obtained at the end of the titration with no salt, indicating that the complex is formed even at lower protein concentrations under these conditions.

The effect of pH in the ASC:ASC and ASC-NLRP3 PYD interactions was also studied. NMR $^1$H–$^{15}$N HMQC spectra of labeled ASC PYD at pH 5.8 were acquired upon the addition of unlabeled ASC and NLRP3 PYD. Even though significant precipitation occurred, changes in chemical shift values were observed (Fig. 8, B–D). The obtained data indicate that the binding regions at higher pH are very similar to those found at more acidic pH.

**ASC PYD L25A Mutation Hinders ASC Interaction through the H2-H3 Face**—The L25A mutant has been typically used in ASC PYD interaction studies because of its enhanced solubility (27, 37). In agreement with previously reported data (16), the $K_D$ values that we obtained from the NMR titration studies of ASC L25A PYD with itself indicate a reduced tendency to self-associate (Fig. 9). The average $K_D$ value for self-association is 165 $\mu$M compared with the average value obtained for the wild type of 69 $\mu$M (Fig. 2A). However, the binding interfaces remain H1, H4, and N-terminal H5 and H2, H3, and C-terminal H5 (where the L25A substitution is located). Interestingly, the
NMR analysis for the L25A mutant shows that the mutation induces structural changes in key residues located in the interface formed by helices H2, H3, and H5 C terminus (particularly in the vicinity of Leu-25 and Leu-45, Fig. 10) and around residue Thr-16, also positioned between the two interfaces. Remarkably, it was proposed that residue Lys-24 is critical for stabilizing H3, and therefore L25A could show a destabilized H3 and hence reduced binding capabilities (17).

In addition, these structural effects on the H2-H3 interface provide a possible explanation for previous studies reporting that the L25A mutant interacts with the PYD-only protein POP1 exclusively through the H1-H4 interface of ASC PYD (27). Interestingly, the relevance of this interface for complex formation was discussed recently regarding ASC-NLRP10 PYD interaction (38). Moreover the L25A mutant was also used to test the interaction between ASC and NLRP4 PYDs with negative results (37). More importantly, the L25A mutant was shown unable to form fibrils (41), likely because the interaction through H2-H3 interface is diminished and therefore only dimers interacting through H1-H4 interface can be formed with no possibility of further oligomerization. Besides, the area where Leu-25 is located was shown to be critical for the stability of H3 in the protein AIM2 PYD, which likely has an effect in the protein interaction capabilities (17). All the evidence supports the importance of using WT ASC PYD rather than solubility-enhanced variants for interaction studies.

**NLRP3 Binding to ASC**—In spite of the high tendency of NLRP3 PYD to aggregate, we were able to perform NMR titrations adding unlabeled ASC and to monitor the regions of the protein affected by the interaction. NLRP3-1H-15N HMQC spectra show severe signal overlap in the amide central region (Fig. 5), precluding the analysis of residues with chemical shift differences versus residue number of ASC PYD upon binding for the ASC-ASC and ASC-NLRP3 interaction at pH 5.8 (both at a ratio of 1:5 protein/protein concentration).

In the case of the ASC-NLRP3 interaction adding unlabeled ASC and to monitor the regions of the exposed residues undergoing above-threshold changes in chemical shift upon titration of up to 76% of the total number of signals (Fig. 11), precluding the analysis of residues with chemical shift differences adding unlabeled ASC and to monitor the regions of the exposed residues undergoing above-threshold changes in chemical shift upon titration of up to 76% of the total number of signals.

**FIGURE 9. ASC L25A mutant shows reduced binding affinity.** A and B, mapping of the exposed residues undergoing above-threshold changes in δ for ASC L25A upon self-binding (A) and NLRP3-binding (B). C, KD values for the ASC L25A-ASC L25A interaction are as follows: Arg-5 (120 ± 2 μM (r = 0.95)); Leu-9 (143 ± 61 μM (r = 0.92)); Thr-16 (126 ± 55 μM (r = 0.92)); Glu-19 (174 ± 76 μM (r = 0.92)); Thr-63 (375 ± 264 μM (r = 0.87)); Glu-67 (216 ± 49 μM (r = 0.98)); His-90 (214 ± 115 μM (r = 0.90)). D, KD values for the ASC L25A-NLRP3 PYD interaction are as follows: Arg-5 (23 ± 20 μM (r = 0.93)); Leu-9 (49 ± 32 μM (r = 0.96)); Thr-16 (3 ± 5 μM (r = 0.91)); Glu-19 (4 ± 7 μM (r = 0.85)); Thr-63 (51 ± 32 μM (r = 0.96)); Glu-67 (7 ± 6 μM (r = 0.94)); His-90 (21 ± 17 μM (r = 0.94)).
α-helix (residues Thr-4, Gly-35, Ile-37, Phe-59, Gly-61, Glu-63, Trp-66, Ala-67, Val-70, Trp-71, Ala-74, Glu-89, and Lys-91). Interestingly, signal shifts observed upon titration with ASC coincide in direction with those observed when a higher concentration of NLRP3 is used (Fig. 11A), indicating that NLRP3 self-associates using identical interfaces as those used for its interaction with ASC PYD (as it occurs in ASC PYD). In general, these interfaces coincide with those in pyrin involved in binding to ASC PYD (44). As in the case of PYD ASC, the presence of opposing interfaces for the interaction enables the formation of oligomeric ASC-NLRP3 PYD heteromolecular complexes, which is the basis for inflammasome molecular activation (Fig. 12) (1–4, 18).

**Supramolecular Complex Models of ASC:ASC and ASC:NLRP3 Oligomers**—Using the NMR data on the specific residues critical for interaction, we generated several models of the dimeric complexes formed by docking the interfaces observed. Specifically, for ASC:ASC PYD pair we only obtained continuous oligomeric structures using as unity the complex formed by ASC PYD H1-H4 with ASC PYD H2-H3 (Fig. 13). In turn, this second PYD domain would still contain an accessible H1-H4 interface, which we used as a template for the structural alignment of a second ASC:ASC pair. Given the distribution of charged residues in the domain, we can clearly observe differently charged interfaces that could promote polymerization (Fig. 14A).

In this way, the structure obtained is an intertwined helical fibril, which somehow resembles the mature ASC PYD fibril described recently by cryo-electron microscopy (18). However, these mature fibrils contain a helicoidal arrangement of PYD domains with six repeats per turn, with much larger dimensions than the resulting fibril obtained based on our NMR data. Interestingly, the authors also show that the fibrils grow significantly fast from protofibrils with small dimensions to larger assemblies. The mature fibrils are stabilized by three types of interactions between the PYD subunits with the type I as dominant and the only one found within each strand, whereas type II and III are highly variable and are found between the strands stabilizing the mature fibril. In our experimental conditions, we only observed the type I of interaction (Figs. 1, B and D) (13). Besides, the structure of the ASC PYD domain inside the mature fibrils is perturbed (18), with significant conformational changes on H3, which the authors attribute to conformational rearrangements. Along these lines, the type I interaction characterized here is slightly different from the one described in the mature fibril. We observed the following three critical electrostatic clusters in our dimer: Glu-18 and Glu-19 from H2 contact with Arg-5 from H1; Lys-21 from H2 interacts with Asp-6 and Asp-10 from H1; and Lys-22 and Lys-26 from H2 interact with...
Asp-47 from H4 (Fig. 14A). In contrast, the electrostatic clusters found in the type I interaction observed in the mature fibril were as follows: Lys-21 from H2 interacts with Asp-48 and Asp-51 from H3-H4; Lys-22 and Arg-41 from H2-H3 interact with Asp-54 from H4; and Lys-26 from H2 contacts with Arg-3 and Arg-5 from H1 (18). In general lines, these differences result from the rotation of one protomer relative to the other. Remarkably, in the protofibril proposed here, the residues responsible for the type II and III interactions found in the mature fibril remain accessible, and therefore, it would still be possible to pack protofibrils laterally to form the mature fibril.

Nonetheless, whether the so-called protofibril we describe here represents relevant initial steps of fibrillization is still to be investigated. However, it is worth mentioning that among the 12 residues mutated in the study with effects in fibril formation (18), seven participate in the dimer interface shown here (Glu-13, Lys-21, Lys-22, Lys-26, Arg-41, Leu-50, and Asp-51). Another ASC-ASC PYD homodimer model has been proposed (16). In this case, the arrangement of the protomers is significantly different from what our NMR data indicate (Fig. 14B). The differences might arise from the low number of restraints used in the mentioned study to drive the docking calculations.

The interaction between ASC and NLRP3 PYDs has remained elusive in in vitro studies despite much evidence of its occurrence in vivo (27). However, NMR data clearly indicate the presence of such interaction under our experimental conditions. Based on the titration results, we modeled by docking four different types of binding between ASC and NLRP3 PYDs that agree with the interaction interfaces characterized on monomeric PYDs (Figs. 1 and 11) as follows: ASC H1-H4/
stabilizing the ASC-NLRP3 ring (namely Glu-15, Lys-23, Glu-64, and Asp-82).

Similarly to what was previously found in PYD fibrils (18, 39), the topology of the ring (Fig. 15B) makes the CARD domains of ASC molecules accessible for further interaction given the flexibility of the linker between ASC PYD and CARD domains (29). This flexibility could in turn serve as a mechanism for oligomerization into ring structures (45, 48). Interestingly, several mutations associated with hereditary periodic auto-inflammatory syndromes were found in the PYD of pyrin protein (49, 50). The residues involved in some of these disease-related mutations (corresponding to Tyr-11, Val-18, Lys-24, Arg-41, and Asp-90 of NLRP3) are located in the ASC-NLRP3 interaction interfaces described here. Thus, it is reasonable to speculate that those mutations could hamper the formation of the ring shown here.

In our study we also show that the PYD-PYD dimer interaction is low affinity. This feature, typical within the death domain superfamily (3, 45), might allow the formation of flexible oligomeric structures that would recruit more components to form robust multimeric ensembles. Although ASC-NLRP3 PYD interaction is of higher affinity, we cannot discard simultaneous self-association and heteromeric interaction. Thus, different combinations would result in different supramolecular complexes formed. Indeed, an auto-inhibition mechanism is proposed to occur in NLRP3 through oligomerization by its leucine-rich repeat domain (18), and thus the oligomerization properties might be different from the ones described here. Therefore, it is of paramount importance to describe in detail the supramolecular structure of the oligomeric rings triggering NLRP3 inflammasome activation.

All in all, the study reported herein expands our understanding of the role of PYD domains in NLRP3 inflammasome formation. Our results likely represent the initial stages of the oligomerization process in the activation cascade and provide insight into the molecular basis underlying this labyrinthine mechanism. Such insight is in fact of critical importance to target specific oligomeric structures or oligomerization steps in NLRP3 inflammasome formation in the therapeutic treatment of a wide range of diseases (23, 25, 51). Along this line, our results anticipate that NMR spectroscopy can be used as a powerful, robust technique for the detailed analysis of antagonists of inflammasome complex formation.

### Experimental Procedures

**Protein Cloning, Expression, and Purification—Human**

ASC PYD (residues 1–90; UniProtKB/Swiss-Prot accession number Q9ULZ3), ASC PYD L25A mutant, and NLRP3 PYD cDNAs (residues 3–93; UniProtKB/Swiss-Prot accession number Q96P20) were cloned into pET15b expression vectors (Novagen). ASC PYDs contained an N-terminal six-histidine tag and a thrombin cleavage site, whereas NLRP3 did not contain tags to avoid any possible perturbation in its solution structure.

Proteins were expressed in C41(DE3) _Escherichia coli_ strain (52). For unlabeled protein production, cells were grown in LB media, whereas isotopic labeling was achieved in M9 minimal media, whereas isotopic labeling was achieved in M9 minimal
media supplemented with $^{15}$NH$_4$Cl or $^{13}$C-glucose as the sole nitrogen and carbon sources, respectively. Overexpression of the recombinant proteins was induced during 4 h at 37 °C by the addition of 1 mM isopropyl $\beta$-D-1-thiogalactopyranoside to the respective cultures after an $A_{600}$ of 0.8 was reached. Bacterial pellets were resuspended in 50 mM Tris-HCl (pH 8.0), and cells were lysed by sonication. The soluble fraction was removed by centrifugation at 48,000 $\times$ g for 30 min. Inclusion bodies were then washed twice in the aforementioned buffer. The resulting pellet was then dissolved in the presence of 6 M GdnHCl and centrifuged at 48,000 $\times$ g for 1 h.

ASC PYD was purified by Ni$^{2+}$-affinity chromatography. Specifically, pellets were dissolved in 20 mM Tris-HCl, 500 mM NaCl, 5 mM imidazole, 6 M GdnHCl (pH 7.9) (binding buffer). His Bind resin (Novagen) was charged according to the manufacturer’s instructions. Protein was attached to the resin using binding buffer and washed in the presence of 20 mM imidazole. Elution was achieved by using binding buffer in the presence of

![Figure 15](https://example.com/figure15.png)

**FIGURE 15. Proposed ASC-NLRP3 PYD oligomeric complex.** A, four different types of interaction between ASC and NLRP3 PYDs are possible based on our NMR data as follows: ASC H1-H4/NLRP3 H5; ASC H1-H4/NLRP3 H1-H4; ASC H2-H3/NLRP3 H5; and ASC H2-H3/NLRP3 H1-H4. ASC PYD and NLRP3 PYD are shown as ribbon diagrams and colored in green and purple, respectively. The residues highlighted as stick representation in the structures were used to drive the specific dockings. In each pair, a green circle represents ASC PYD, and a purple square represents NLRP3 PYD, and the numbers indicate the corresponding type of interaction. B, hexameric ring modeled using as template the resulting docking pairs of the different dimeric units shown in A. ASC PYD, ASC CARD, and ASC linker and NLRP3 PYD are colored in green, blue, gray, and purple, respectively. The C termini of NLRP3 PYDs placed in the ring point to the bottom of the image shown on the right. The structures were structurally aligned and represented with PyMOL (73).
1 M imidazole. Immediately after elution, the protein was subjected to several dialysis steps against H$_2$O (pH 4.0) to slow protein self-association processes. The sample was subsequently purified by reverse phase chromatography using a prepacked C8 column (Higgins Analytical). Binding buffer consisted of 5% acetonitrile, 0.1% trifluoroacetic acid (TFA), and elution buffer was 95% acetonitrile, 0.1% TFA. As described previously and contrary to WT ASC PYD (35, 37), ASC PYD L25A mutant remains in the soluble fraction and thus could be purified in non-denaturing conditions. A single Ni$^{2+}$-affinity chromatography purification step was performed for this protein.

NLRP3 PYD was purified using a double-reverse phase chromatography, the first one performed in denaturing conditions. C4 resin (Applied Separations) was equilibrated in 20 mM Tris-HCl, 500 mM NaCl, 6 M GdnHCl, 0.1% TFA. Inclusion bodies extract (resuspended in the same buffer used for C4 resin equilibration) was slowly bound to the resin in several cycles. Once the protein was bound, the column was re-equilibrated in 5% acetonitrile, 0.1% TFA using a very slight gradient, and protein was eluted in 60% isopropyl alcohol, 40% acetonitrile, 0.1% TFA. Different fractions of the chromatogram (with different purity) were pooled, lyophilized, and resuspended in 5% acetonitrile, 0.1% TFA prior to their subsequent purification in regular reverse phase chromatography using a C8 pre-packed column as described above.

After purification, all three proteins were >95% pure, as monitored by SDS-PAGE, and the correct molecular weight was calculated from MALDI-TOF mass spectrometry data. Proteins were lyophilized and stored for subsequent studies.

**NMR Spectroscopy for Binding Studies**—Unless otherwise stated, NMR experiments were acquired on a Bruker Avance 600 MHz spectrometer equipped with a triple-resonance cryogenic probe and on a Bruker Avance III 600 MHz spectrometer with a triple-resonance, three-axis gradient probe in 20 mM 600 MHz spectrometer equipped with a triple-resonance cryo-stated NMR experiments were acquired on a Bruker Avance spectrometer at CSIC - Centro de Investigaciones Biológicas on October 7, 2016.

**Solution Structure and Binding Properties of NLRP3 PYD**

Calculation of Dissociation Constants—The observed $\delta$ values in the NMR titration experiments are weighted average values of the free and bound forms of the protein because the exchange rate between both forms is fast on the NMR chemical shift time scale. The difference in $\Delta(\delta)$ was obtained as follows: $\Delta(\delta)_{\text{av}}$ = $\delta_{\text{bound}} - \delta_{\text{free}}$ (57). Those residues showing $\Delta(\delta)_{\text{av}}$ values higher than 1.5 S.D. upon ligand binding and with a total solvent exposure index larger than 30% as calculated by GETAREA (57) were defined as critical for the corresponding interaction. To obtain dissociation constants ($K_d$), the $\Delta(\delta)_{\text{av}}$ versus ligand protein concentration plots were fit to the equation as follows: $\Delta(\delta)_{\text{av}} = K_d + L + P_0 - (K_d + L + P_0 - 4P_0L)^{1/2}/2P_0$, where $P_0$ is protein concentration (50 $\mu$M), and $L$ is total ligand concentration (58). Data fitting was done with OriginPro 8 (OriginLab).

**NMR Spectroscopy for NLRP3 Structural Studies**—NMR samples were prepared at 0.1–0.2 mM NLRP3 PYD in 5 mM $d_{15}$-TCEP, 0.1 mM Na$_3$H$_5$, 5% D$_2$O (pH 3.6). NMR experiments were acquired at 303 K in a Bruker Avance 600 MHz spectrometer equipped with a triple-resonance cryogenic probe. Sequence backbone assignments were obtained from the following experiments: $^{1}$H-$^{15}$N HSQC, three-dimensional HNCA, and three-dimensional CBCA(CO)NH. Side chain assignments were obtained from three-dimensional HBHA(CO)NH, three-dimensional (HCCC)O)NH TOCSY (22.5-ms mixing time), and three-dimensional HCH-TOCSY (15-ms mixing time). NOE data were obtained from three-dimensional $^{13}$C-$^{1}$H-$^{1}$H NOESY (110-ms mixing time) and three-dimensional $^{13}$C-$^{1}$H-$^{1}$H NOESY (120-ms mixing time). Information on NMR experiments for protein structure determination can be found elsewhere (59, 60). All experiments were processed with NMRPipe (55) and analyzed with PIPP (61).

**NLRP3 Structure Calculation**—Peak intensities from NOESY experiments were translated into a continuous distribution of interproton distances. Distances involving methyl groups, aromatic ring protons, and non-stereospecifically assigned methylene protons were represented as a summation averaging, $\langle r^{-6}\rangle^{-1/6}$ (62). Errors of 25 and 40% of the distances were applied to obtain lower and upper distance limits. The lower limit of some distance restraints that appeared smaller in the resulting structures for protons belonging to sequential amino acids was increased to 50%. A total of 32 hydrogen bond distance restraints ($r_{\text{NH-O}} = 1.9–2.5$ Å and $r_{\text{N-O}} = 2.8–3.4$ Å) was defined according to the experimentally determined secondary structure of the protein. The TALOS+ program was used to obtain $\phi$ and $\psi$ restraints for those residues with statistically significant predictions, and the associated predicted errors were used. Structures were calculated with the program X-PLOR-NIH 2.16.0 (63). The structure was heated to 3000 K and cooled in 10,000 steps of 0.002 ps during simulated annealing. The final ensemble of 20 NMR structures was selected based on lowest energy and no restraint-violation criteria. The 20 lowest energy conformers have no distance restraint violations and no dihedral angle violations greater than 0.5 Å and 5°, respectively. Structure quality was assessed with PROCHECK-NMR (42) and MolProbity (64). Structures were

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**NMR Spectroscopy for Binding Studies**—NMR experiments were acquired at 0.1–0.2 mM NLRP3 PYD in 5 mM $d_{15}$-TCEP, 0.1 mM Na$_3$H$_5$, 5% D$_2$O (pH 3.6). NMR experiments were acquired at 303 K in a Bruker Avance 600 MHz spectrometer equipped with a triple-resonance cryogenic probe. Sequence backbone assignments were obtained from the following experiments: $^{1}$H-$^{15}$N HSQC, three-dimensional HNCA, and three-dimensional CBCA(CO)NH. Side chain assignments were obtained from three-dimensional HBHA(CO)NH, three-dimensional (HCCC)O)NH TOCSY (22.5-ms mixing time), and three-dimensional HCH-TOCSY (15-ms mixing time). NOE data were obtained from three-dimensional $^{13}$C-$^{1}$H-$^{1}$H NOESY (110-ms mixing time) and three-dimensional $^{13}$C-$^{1}$H-$^{1}$H NOESY (120-ms mixing time). Information on NMR experiments for protein structure determination can be found elsewhere (59, 60). All experiments were processed with NMRPipe (55) and analyzed with PIPP (61).

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analyzed with MOLMOL (65). Atomic coordinates were deposited in the Protein Data Bank with accession code 2NAQ.

Analytical Ultracentrifugation—AUC analysis of ASC PYD and NLRP3 PYD was performed in an XL-A analytical ultracentrifuge (Beckman-Coulter Inc.) with UV-visible optics detection system, using an An50Ti rotor and 3- or 12-mm double-sector centerpieces.

Sedimentation equilibrium experiments were performed on ASC PYD samples (50 –745 μM) resuspended in 20 mM glycine, 1 mM TCEP (pH 3.7) at several speeds (16,000 and 20,000 rpm) and the adequate wavelength (256–292 nm), using short columns (23 or 90 μL depending on the pathlength, 3 or 12 mm, respectively). After the equilibrium scans, a high speed centrifugation run (48,000 rpm) was performed to estimate the corresponding baseline offsets. Weight-average buoyant molecular mass of ASC PYD samples was determined by fitting data to the single species model using the SEDFIT program (66).

Non-ideality correction due to very high concentrations of protein was estimated according to Ref. 68. The actual weight-average molecular weight is related to the apparent weight-average molecular weight obtained experimentally by the simple Equation 1 (69),

\[ M_{w, a} = M_w \exp (-7.86 v W_{tot}) \]  
(Eq. 1)

where \( M_{w, a} \) is the experimentally measured apparent weight-average molecular weight, and \( v \) is the experimentally measured partial specific volume of the protein (we used the calculated \( v \) from the amino acid composition with SEDNTERP (67)).

Sedimentation velocity experiments were carried out at 48,000 rpm and 20 or 30 °C. Samples of ASC PYD ranged from 50 to 745 μM. Samples of NLRP3 PYD (9.8–243 μM) were resuspended in H₂O, 1 mM TCEP (pH 3.6). Sedimentation profiles were registered every 5–8 min at the adequate wavelength. The sedimentation coefficient distributions were calculated by least squares boundary modeling of sedimentation velocity data using the c(s) method (70), as implemented in the SEDNTERP program (67). These \( s \) values were corrected to standard conditions (water, 20 °C, and infinite dilution (71)) using the SEDNTERP program (67) to obtain the corresponding standard \( s \) values (\( s_{20,w} \)).

Molecular Modeling—Molecular docking was performed with the program HADDOCK (72) using both the ASC PYD domain of full-length ASC solution structure (PDB code 2KN6 (29)) and the NLRP3 PYD solution structure described in this work (PDB code 2NAQ) as templates. Those residues showing a \( \Delta D^{\text{min}} \) value of >1.5 S.D. upon ligand binding and with a total solvent exposure index of >30% as calculated by GETAREA (57) were selected as “active” residues to create a list of ambiguous interaction restraints used in the docking calculation. Specifically, the interfaces used for the docking calculations were those formed by \( \alpha \)-helices H1, H4, and the N-terminal end of H5 (residues Gly-2, Arg-3, Asp-6, Asp-10, Thr-16, Leu-50, Asp-54, Glu-62, Thr-63, Glu-67, and His-90) and by helices H2, H3, and the C-terminal end of H5 (residues Glu-19, Lys-22, Lys-24, Leu-25, Ser-29, Val-30, Leu-45, and Asp-75) from ASC PYD, whereas the interfaces formed by \( \alpha \)-helices H1 and H4 (residues Arg-10, Glu-13, Asp-14, Glu-16, Val-18, Lys-21, Asp-48, Val-50, and Asp-51) and by residues from H5 (Gly-61, Glu-63, Ala-67, and Trp-71) were selected from NLRP3 PYD. As a result, three different combinations of docking models were used for the ASC-ASC complex (ASC [H1,H4]-ASC [H2,H3]; ASC [H1,H4]-ASC [H1,H4]; and ASC [H2,H3]-ASC [H2,H3]) and four for the ASC-NLRP3 complex (ASC [H1,H4]-NLRP3 [H1,H4]; ASC [H2,H3]-NLRP3 [H1,H4]; ASC [H1,H4]-NLRP3 [H5]; ASC [H2,H3]-NLRP3 [H5]; ASC [H2,H3]-NLRP3 [H5]). HADDOCK default parameters were accepted for the docking protocol. No other restraints were incorporated.

The top seven HADDOCK clusters of structures out of 1,000 calculated were inspected to build the models. For the ASC-ASC interaction, only the supramolecular complexes resulting from the ASC [H1,H4]-ASC [H2,H3] pair are shown because they are the only ones conforming to a continuous oligomeric structure inducing no steric clashes, whereas for ASC-NLRP3, we show all the possible combinations of pairs following our docking models. The complexes were built by structural alignment on PyMOL using as templates consecutive units of the corresponding pair.

Author Contributions—J. O. produced and purified all proteins, conducted the great majority of NMR experiments, performed data analysis, and built all supramolecular models. J. O. and E. dA. conducted and analyzed NMR experiments, discussed results, and wrote the manuscript. E. dA. conceived the project, analyzed NMR data on NLRP3, and determined the NLRP3 three-dimensional structure. S. B.-V. produced and purified NLRP3 and analyzed preliminary NMR data on NLRP3. C. A. and G. R. conducted and analyzed the AUC experiments.

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ASC Pyrin Domain Self-associates and Binds NLRP3 Protein Using Equivalent Binding Interfaces

Javier Oroz, Susana Barrera-Vilarmau, Carlos Alfonso, Germán Rivas and Eva de Alba

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