2015

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Structural Studies of AAV2 Rep68 Reveal a Partially Structured Linker and Compact Domain Conformation

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Supporting Information

ABSTRACT: Adeno-associated virus (AAV) nonstructural proteins Rep78 and Rep68 carry out all DNA transactions that regulate the AAV life cycle. They share two multifunctional domains: an N-terminal origin binding/nicking domain (OBD) from the HUH superfamily and a SF3 helicase domain. A short linker of ~20 amino acids that is critical for oligomerization and function connects the two domains. Although X-ray structures of the AAV5 OBD and AAV2 helicase domains have been determined, information about the full-length protein and linker conformation is not known. This article presents the solution structure of AAV2 Rep68 using small-angle X-ray scattering (SAXS). We first determined the X-ray structures of the minimal AAV2 Rep68 OBD and of the OBD with the linker region. These X-ray structures reveal novel features that include a long C-terminal α -helix that protrudes from the core of the protein at a 45° angle and a partially structured linker. SAXS studies corroborate that the linker is not extended, and we show that a proline residue in the linker is critical for Rep68 oligomerization and function. SAXS-based rigid-body modeling of Rep68 confirms these observations, showing a compact arrangement of the two domains in which they acquire a conformation that positions key residues in all domains on one face of the protein, poised to interact with DNA.

The nonstructural Rep proteins from adeno-associated virus (AAV) are multifunctional proteins with specialized domains equipped to handle the complex interactions with DNA during the AAV life cycle.^{1,2} AAV has a single-stranded DNA genome of ~4.7 kb containing two major open reading frames (ORFs) flanked by inverted terminal repeats (ITRs). The ITRs form T-shaped hairpin structures and contain the cis-regulatory functions required for replication, transcriptional regulation, and possibly site-specific integration.^{3–9} The stem of the hairpin forms a double-stranded region containing a Rep binding site (RBS) made up of several 5'-GCTC-3' repeats.¹⁰ A terminal resolution site (trs) located upstream of the RBS is the site of a strand- and site-specific endonuclease reaction required to complete the replication of the AAV genome.^{3,11,12} Despite its limited genome size, AAV generates eight different polypeptide chains. The right ORF regulated by the P₄₀ promoter produces three capsid proteins (VP1–3) and an assembly activating protein (AAP).^{13–18} The left ORF uses two different promoters, and alternative splicing, to generate four nonstructural proteins: two large Rep proteins (Rep78 and Rep68) transcribed from the p5 promoter and two small Rep proteins (Rep52 and Rep40) regulated through the p19 promoter.^{14,19–21} Thus, Rep40 is equivalent to the helicase domain of Rep68. Most of the biochemical activities required for transcriptional regulation,^{22,23} DNA replication^{24,25} and site-specific integration^{26–28}

are carried out by the large Rep78/Rep68 proteins, whereas the small Reps are thought to be important for DNA packaging into preformed capsids.^{29,30}

The wide spectrum of functions performed by Rep78/Rep68 is a direct reflection of the multifunctionality of their domains. The N-terminal origin binding domain (OBD) displays site- and strand-specific endonuclease activity and recognizes the GCTC repeats.^{24,31–34} Structurally, the OBD is a member of the HUH endonuclease family specialized in the cleavage and rejoining of single-stranded DNA substrates (ssDNA), which occurs during transposition, bacterial conjugation, rolling-circle replication of bacterial DNA and bacteriophages, and in the replication of small eukaryotic viruses.³⁵ This family is characterized by two signature motifs that participate in the nuclease reaction: the HUH motif, which consists of two histidine (H) residues separated by a large hydrophobic residue (U), and the Y motif, with either one or two tyrosine (Y) residues. Structurally, the common feature of HUH endonucleases is a five-stranded antiparallel β -sheet surrounded by α -helices, where the HUH motif is found in one of the strands.

Received: June 3, 2015

Revised: August 20, 2015

Published: August 28, 2015

Figure 1. Structure of AAV2 OBD. (A) Domain structure of AAV2 Rep68 protein: OBD is shown in green, Rep40 (SF3 helicase domain), in blue, and linker and C-terminal tail, in gray. The Rep40 protein spans residues 225–536 of Rep68. (B) Topology diagram of AAV2 OBD. (C) Ribbon diagram of the OBD structure. The α -helices are light green, and β -strands are dark green. Secondary structure elements are labeled with α -helices A–F, and β -sheets are numbered 1–5. The DNA binding loop L_{DB} with missing residues in the structure is shown as a dotted line. (D) A view of the structure rotated by 90° clockwise. α -Helix F is shown protruding from the core of the structure at an angle of almost 45°.

The position of the Y motif can vary: in the relaxase subgroup, it is located N-terminally, whereas it is found at the C-terminus in the Rep subclass.^{36–38}

The helicase domain found in all Rep proteins is a representative of the SF3 family helicase with 3' to 5' unwinding activity.^{38–43} The ATPase core is a modified version of the AAA⁺ domain with additional features that include an N-terminal helical bundle and a β -hairpin that is involved in DNA binding during translocation.³⁸ Consequently, the large Rep proteins can interact with DNA in two different modes:

one is mediated by the OBD and recognizes DNA in a sequence-specific manner, whereas the SF3 helicase domain interacts with DNA nonspecifically.⁴⁴ Furthermore, the large Rep proteins show a complex and dynamic oligomerization behavior, which may add an additional level of regulation to Rep interactions with DNA.⁴⁵ How the two domains cooperate with each other in order to interact with DNA and how they are arranged in the context of Rep78/Rep68 is not known. Moreover, studies have established that the linker region plays a critical role in the oligomerization and function of AAV Rep

proteins; therefore, it is likely that this region acquires a particular secondary structure and may not simply be extended.^{46,47} To answer these questions, we performed X-ray crystallography and small angle X-ray scattering (SAXS) studies on the AAV2 Rep68 functional domains and full-length protein. Our results show that the OBD has a long extended helix that includes part of the linker, which is not extended, but is tilted in a way that brings the two domains into a compact configuration. SAXS studies on a monomeric version of full-length Rep68 confirm this observation and show for the first time the overall domain architecture of a full-length AAV Rep protein.

MATERIALS AND METHODS

Protein Expression and Purification. The DNA region encoding amino acids 1–208 (OBD) and 1–224 (OBD plus linker) from adeno-associated virus type 2 (AAV2) (GeneBank protein_id = AF043303.1) was cloned into pET15b (Novagen) using restriction sites *NdeI* and *XhoI*. The residue C151 was mutated to serine, as it was found to produce disulfide bonds and inhibit crystallization, and Rep-C151S was shown to be fully functional in supporting the AAV life cycle.⁴⁵ The OBD constructs were overexpressed in *Escherichia coli* strain BL21 pLysS at 37 °C in Luria–Bertani (LB) broth. IPTG (isopropyl- β -D-thiogalactopyranoside) was added to a final concentration of 1 mM when an OD of 0.6 was reached. Cells were harvested after 5 h and stored at –80 °C. The cell pellets were resuspended in binding buffer (20 mM Tris-HCl, 500 mM NaCl, 10 mM imidazole, 10% glycerol, 1 mM TCEP, pH 7.9) and lysed by sonication. The OBD was purified with a Ni-NTA column (Qiagen) using step gradients of 10 and 30 mM imidazole to wash nonspecific proteins binding to the column and then eluted with 100 mM imidazole. Protein was loaded onto a HiLoad desalting column (GE) to change into thrombin buffer (25 mM Tris-HCl, 200 mM NaCl, 10% glycerol pH 8.0). His-tag was cut by addition of thrombin (1 unit/mg) and removed by passing through a Ni-NTA column. The untagged OBD was collected from the flow through, concentrated, and further purified by gel filtration on a HiLoad 16/60 Superdex 75 column (GE Healthcare) previously equilibrated with gel filtration (GF) buffer (25 mM Tris-HCl, 200 mM NaCl, 1 mM TCEP, pH 7.5). The protein was concentrated to ~40 mg/mL using Millipore Centricon (10 kDa cutoff). AAV2 Rep40 was purified as described elsewhere.³⁸ AAV-2 Rep68_{wt} (1–536) and Rep68Y224 Δ were expressed in *E. coli* strain BL21 pLysS at 18 °C as described in earlier reports.⁴⁶ In brief, histidine-tagged Rep68 was purified in a Ni-NTA column, and after PreScission protease cleavage of the His-tag, the protein was purified on a HiLoad Superdex 200 16/60 column (GE Healthcare). OBDL was purified using the same procedure.

Crystallization, X-ray Data Collection, and Structure Determination. Crystallization was carried out using the hanging-drop method with commercially available screening kits at 4 °C. Crystals grew after 2 to 3 days in 50 mM cacodylate, pH 6.5, 80 mM sodium acetate, 15 mM magnesium acetate, and 8–10% isopropanol. Crystals were cryoprotected in reservoir buffer and supplemented with 20% MPD before flash freezing them in liquid nitrogen. The crystals diffracted to 2.3 Å and belonged to space group $P2_12_12$ with unit cell dimensions $a = 186.4$ Å, $b = 154.4$ Å, $c = 38.8$ Å. Diffraction data was collected at the National Synchrotron Light Source (NSLS) at Brookhaven National Laboratory beamline X6a. The data were processed with the program HKL2000,⁴⁸ and the structure was solved by molecular replacement using the

Table 1. Data Collection and Refinement Statistics

	OBD (1–208)	OBDL (1–224)
space group	$P2_12_12$	$P2_1$
Cell Dimensions		
a, b, c (Å)	186.4, 154.4, 38.8	75.6, 178.7, 130.4
α, β, γ (deg)	90, 90, 90	90, 91.7, 90
wavelength (Å)	0.9792	1.54
resolution (Å)	30–2.30 (2.34–2.30)	30–2.6 (2.69–2.6)
no. of measured	566 473	426 030
no. of unique	50 960	105 074
data coverage (%) ^a	99.8 (100)	99.1 (98.9)
$R_{\text{merge}}(\%)^{a,b}$	0.099 (0.402)	0.075 (0.311)
I/σ^a	17.7 (8.6)	12.4 (4.1)
refinement statistics		
resolution range	30.0–2.3	29.8–2.6
reflections	50 960	105 007
$R_{\text{work}}(\%)^c$	22.5	19.1
$R_{\text{free}}(\%)^d$	24.2	23.8
non-hydrogen atoms	5024	17677
protein	4704	17493
metal	2	11
water	314	173
average B-factors (Å ²)		
protein	50.0	59.2
metal	49.2	88.4
water	46.4	47.3
RMS Deviations		
bonds (Å)	0.02	0.003
angles (Å)	1.4	1.53
Ramachandran Plot Quality		
most favored (%)	99.0	97.0
additional allowed (%)	1.0	3.0
generously allowed (%)	0	0.0
disallowed (%)	0	0

^aValues for the outermost shells are given in parentheses. ^b $R_{\text{merge}} = \sum |I - \langle I \rangle| / \sum I$, where I is the integrated intensity of a given reflection. ^c $R_{\text{work}} = \sum ||F_o| - |F_c|| / \sum |F_o|$. ^dFor R_{free} calculations, 5% of data was excluded from refinement.

program PHENIX. We used the structure of the AAV5 OBD as a search model (PDB ID: 1M55). Model building was carried out using PHENIX,⁴⁹ and manual building, using the program COOT.⁵⁰ OBDL data was collected using our X-ray home source that consists of a Rigaku Micromax 007 X-ray generator and a Raxis IV⁺ area detector.

Sedimentation Velocity. Sedimentation velocity experiments were carried out using a Beckman Optima XL-I analytical ultracentrifuge (Beckman Coulter Inc.) equipped with an eight-position AN-60Ti rotor. Rep protein samples were loaded in the cells, using, in all cases, the GF buffer. Samples in double sector cells were centrifuged at 25 000 rpm. In all experiments, temperature was kept at 20 °C. Sedimentation profiles were recorded using UV absorption (280 nm) and interference scanning optics. For analysis of the data, the program Sedfit was used to calculate sedimentation coefficient distribution profiles using the Lamm equation.⁵¹

Small-Angle X-ray Scattering (SAXS). Data were collected at three different concentrations that produce a single homogeneous population, as determined by sedimentation velocity studies. Synchrotron SAXS measurements were performed at

Figure 2. Superposition of OBD molecules in the asymmetric unit. (A) Ribbon diagram of the superposition of the three molecules in the asymmetric unit. Molecule A is shown in green; molecule B, blue; and molecule C, red. The three loops L_{DB} have different conformations and are shown as dotted connections. (B) Ribbon diagram of the dimer in the asymmetric unit formed by molecules A and B. The small dimer interface occurs through residues in α -helix F. (C) Details of the residues involved in the oligomeric interaction with Ser197, Val196, and Leu193 from molecule A interacting with His192, Leu193, and Val196 from molecule B. (D) Sedimentation velocity profiles of Rep68_{wt}, Rep68-L193AV196A, Rep68-H192A, and Rep68-S197A. Experiments were carried out at 20 °C and 25 000 rpm on a Beckman XL-I analytical ultracentrifuge. Scans were collected every 2 min using absorbance at 280 nm. Data was analyzed using the program Sedfit.⁵¹ (E) Helicase assay of Rep68 mutants. Fluorescent-labeled DNA molecule has a 3' single-stranded tail and an 18 bp region (first lane). Upon ATP addition, helicase activity is shown as the 18 nucleotide fluorescein-labeled ssDNA is displaced (second lane). Rep68, L193AV196A, H192A, and S197 proteins all unwind DNA. K340H mutant is ATPase-negative and does not unwind DNA. Last lane shows the DNA substrate after heating at 100 °C for 5 min. (F) Comparison of the production of rAAV2-GFP infectious particles in the presence of wt or LV mutant. Various volumes of supernatant (in μ l, x axis) from 293T cells producing rAAV2-GFP were added to HeLa cells, and the percentage of GFP-positive infected cells was determined by FACS analysis, as described in [Materials and Methods](#). Data is presented as average $\pm$ standard deviation from three independent experiments.

Brookhaven National Laboratories at beamline X9 and at the Advanced Light Source at the Lawrence Berkeley National Laboratory at the SYBILS beamline. Three different concentrations of each sample were prepared and measured (1, 2, and 3 mg/mL). All data were processed with the package ATSAS.⁵² Buffer subtraction was carried out using beamline-specific software. Radii of gyration (R_g) were evaluated using the Guinier approximation, $sR_g < 1.3$. Distance distribution functions and maximum diameters D_{max} were calculated using the program

GNOM.⁵³ SAXS molecular envelopes were calculated using the programs DAMMIN and GASBOR.^{54,55} Conformational flexibility of the linker and C-terminal tail was analyzed with the program EOM.⁵⁶ We used the structures of Rep40 (PDB ID: 1s9h) and of OBDL and connect the two domains with a flexible linker (residues 215–224) to generate an atomic model of AAV2 Rep68. Using this initial model, we performed rigid-body and molecular dynamics using BILBOMD to generate the best Rep68 model that fits the SAXS data.⁵⁷

DNA Helicase Assay. The helicase assay was based on a modification of the strand-displacement assay described elsewhere.⁵⁸ The DNA substrate (28:18) consists of 3' tail of 10 nucleotides adjacent to 18 bp. The top strand has been labeled with fluorescein (F) at the 5' end. All reactions were performed in a final volume of 50 μ L in a buffer containing 25 mM HEPES, 50 mM NaCl, pH. 7.0. For the reaction, 1 μ M protein was mixed with 0.5 μ M ds F-DNA (28:18). The reaction was started by addition of 5 mM ATP-MgCl₂ and 2.5 μ M trap ssDNA. Reaction was incubated at 25 °C for 1 h. EDTA was used to stop the reaction at a final concentration of 20 mM. Aliquots of 10 μ L were loaded in a 12% bis-acrylamide gel (30%) (19:1) using 6 $\times$ loading dye (0.25 xylene cyanol FF, 30% glycerol). For densitometry and analysis of the bands, a Gel Doc EZ Imager was used, using the automatic lane and band detection tool. Lane background subtraction, white illumination, and an activation time of 300 s were used for the analysis.

Fluorescent Anisotropy Binding Assays. Binding assays were performed using 5 nM fluorescein labeled 41-mer DNA containing the Rep binding site (TGGCGGCGGTTGGG-GCTCGGCGCTCGCTCGCTCGCTGGGCG). Rep68 at different concentrations was mixed with DNA at a final volume of 300 μ L using the following buffer: 25 mM HEPES (pH 7.0), 200 mM NaCl, 1 mM TCEP. Fluorescence readings were taken on a PC1 fluorimeter (ISS, Inc.) with excitation and emission filters at 492 and 528 respectively. Tubes were equilibrated at 20 °C for 20 min before measurement. Each anisotropy point is the average of 10 measurements. Anisotropy is calculated as the ratio of the difference between vertical and horizontal emission intensities over the total normalized intensity. The fraction of DNA bound (B) was calculated using eq 1

$$B = ([A]_x - [A]_{\text{DNA}}) / ([A]_{\text{FINAL}} - [A]_{\text{DNA}}) \quad (1)$$

where $[A]_x$ represents the anisotropy measured at protein concentration x , $[A]_{\text{DNA}}$ is the anisotropy of free fluorescence DNA, and $[A]_{\text{FINAL}}$ is the anisotropy at saturation. Data was fit to a single binding site model using the program PRISM6 (GraphPad). Each experiment was done in triplicate.

AAV Infectious Particles Assay. 293T cells were transfected with three plasmids using polyethylenimine (PEI): an AAV2 ITR-containing plasmid encoding a CAG-controlled GFP gene (pTRUF11), a helper plasmid expressing AAV2 Rep (wt or L193A-V196A cloned from the pHisRep68LV/15b) and Cap derived from pDG, and a third construct containing the adenovirus helper functions (HGTI plasmid).^{59,60} The presence of the L193A-V196A double mutation was confirmed by sequencing (Eurofins). After 72 h, the cell supernatant was harvested and increasing volumes of supernatant were used to infect HeLa cells. The percentage of GFP-positive HeLa cells was determined 48 h postinfection by FACS (FACSCanto, BD Biosciences).

RESULTS

We opted to structurally characterize full-length AAV2 Rep68 using small-angle X-ray scattering and rigid-body modeling. A diagram of the different domains and protein constructs used in this study is shown in Figure 1A. We first determine the crystal structure of its OBD including the region spanning the linker, followed by SAXS studies of Rep40 (SF3 helicase domain) to study the dynamic behavior of the C-terminal tail. Finally, to determine the relative orientation of its two functional domains, we determined the SAXS solution structure of Rep68.

Figure 3. Comparison of AAV2 and AAV5 OBD structures. (A) Ribbon diagram of the superposition of the AAV2 (green) and AAV5 (blue) OBD structures. The N and C termini are marked. Regions that have the largest rmsd are indicated by arrows. (B) Plot of rmsd differences against residue number between AAV2 and AAV5 OBD. (C) Solvent-accessible electrostatic surface comparison of AAV2 and AAV5 OBDs showing three views of 90° counterclockwise rotations. Positive regions are shown in blue, and negative, in red.

Structure of AAV-2 Rep OBD. The OBD construct (residues 1–208) was initially obtained from limited proteolysis experiments.⁶¹ The structure was solved by molecular replacement and refined to 2.3 Å resolution (Table 1). There are three molecules in the asymmetric unit: molecule A, spanning residues 1–138 and 143–206, molecule B, spanning residues 1–139 and 143–201, and molecule C, covering residues 1–131 and 141–191. The missing residues are all part of the DNA binding loop (L_{DB}). Molecules A and B form a dimer interacting through their C-terminal helix F, whereas molecule C does not make any close contacts. The OBD structure is characterized by a central five-stranded antiparallel β -sheet flanked by six α -helices, three at each side of the sheet (Figure 1B). The topology of the β -sheet follows a 4–1–3–2–5 order, with helices E and F on one side of the sheet and helices B–D positioned on the opposite side (Figure 1C,D). There are two long loops: the DNA binding loop L_{DB} connects β -strand 4 to α -helix E and loop L_{BC} connects α -helices B and C and protrudes underneath α -helix F (Figure 1C). L_{DB} was identified as being important in making specific contacts with the major groove of GCTC repeats and shows the highest degree of variation among all AAV Rep isoforms.^{31,62} This is a dynamic region, as shown by the lack of electron density in all three molecules found in the asymmetric unit. The C-terminal helix α F is long, spanning residues 182–206, and protrudes at an approximately 45° angle from the plane of the β -sheet

Figure 4. Structure of OBDL. (A) Ribbon diagram of the structure spanning residues 1–214. (B) Electron density of region 206–214 contoured at 0.9σ. The density fits most of the residues, but it suggests that the region is dynamic. (C) Superposition of OBD and OBDL structures illustrating the flexibility of helix F. (D) Comparison of the sedimentation velocity profiles between Rep68_{wt} (top) and Rep68-P214A (bottom). (E) Comparison of binding isotherms of Rep68_{wt} and Rep68-P214A. Fluorescent anisotropy binding assays were carried out with a 41-mer AAVS1 DNA labeled with fluorescein. Apparent dissociation constants were determined from triplicate experiments and fitted to a one binding site-specific binding model.

(Figure 1D). The residues at the end of helix F appear to be dynamic, as we could see only up to residue 201 in molecule B and to residue 191 in molecule C. Helix F is loosely packed against helix E and the loop connecting β5 to αF. The interactions that keep these three elements together are sparse, suggesting that helix F is flexible. Several dynamic regions in the AAV2 OBD can be observed by the superposition of the three molecules in the asymmetric unit (Figure 2A). The three molecules superimpose with an overall rmsd over 197 C_α of 0.62 Å. Two regions can be identified with larger than average values: region 1 includes residues 15–34 spanning helices A and B, and region 2 includes residues from L_{DB}. Not surprisingly, based on the AAV5 OBD–RBS structure, both of these regions are involved in DNA interactions: region 1 is involved in recognition of the ITR hairpin stem 2, and L_{DB} recognizes part of the GCTC repeat.³¹ Thus, folding and stabilization of these elements must occur upon DNA binding. Two of the OBD molecules in the asymmetric unit form a dimer through interactions with residues present in α-helix F, resembling a pseudocoiled coil (Figure 2B). These include L193, V196, H192, and S197 making hydrophobic and hydrogen-bond interactions (Figure 2C). This type of interface was also observed in the AAV5 OBD–RBS complex, which may suggest a functional role for this interface in the context of full-length Rep78/Rep68 proteins.³¹ To determine if these interactions are functionally relevant in Rep68 function and their effect on the

AAV life cycle, we examined the ability of several Rep68 mutants to oligomerize, to unwind dsDNA, and to support the production of infectious particles. Figure 2D shows that Rep68 mutants H192A, S197A, and L193AV196A have similar sedimentation velocity profiles as that of Rep68_{wt} forming the 13S peak that corresponds to heptameric/octameric ring species.⁴⁵ In addition, the mutants have similar unwinding activity as that of the wild-type protein (Figure 2E). Thus, the different mutants are functionally equivalent to Rep68_{wt}. This conclusion is supported by the ability of the Rep68-L193AV196A double mutant to produce infectious viral particles as efficiently as wild-type Rep68 (Figure 2F). These results show that the dimer interactions are generated during crystal packing and are not involved in Rep68 oligomerization and function.

Structural comparison of the AAV2 and AAV5 OBD.

The Rep proteins from the majority of AAV serotypes show an overall homology with the AAV2 OBD of 95% (Figure S1). In contrast, AAV5 Rep has only 59% identity over the N-terminal 224 residues. However, the AAV2 and AAV5 OBD structures superimposed well, with an overall rmsd of 0.94 Å for 191 aligned C_α carbons. As shown in Figure 3A,B, there are, nonetheless, three regions that show differences: region A (residues 12–15) includes the linker connecting β1 to αA, region B (74–77) spans the turn connecting β3 to αD, and region C (139–147) consists of L_{DB}. Among all AAV serotypes,

AAV5 and AAV8 have the largest sequence variations. For instance, AAV5 has a shorter L_{DB} loop than AAV2, whereas in AAV8, the loop is longer by 2 residues. In addition, the AAV5 protein has four additional residues after position 198. Moreover, sequence conservation in the αF region is low in AAV5, suggesting that this structure may indeed be shorter than in AAV2 (Figure S1). The overall electrostatic surface representations of the two proteins look similar, but subtle differences are visible. In both structures, one side of the surface is highly positively charged; however, in AAV2 OBD, the positive patch is wider and more intense. This is seen more explicitly if we compare the positive surface potential values between the two OBDs, with 2130.6 kcal/molq for AAV2 and 1720.4 kcal/molq for AAV5.⁶³ This area includes L_{DB} , helix D, and the N-terminal half of α -helix C. The former two interact with the GCTC repeats, whereas the latter is involved in the recognition of the RBE' hairpin in the ITRs.³¹ Helix F is mostly negatively charged, but in AAV2, there is a slightly positive patch located in the first three turns that is not visible in AAV5 (Figure 3C). Because the structure of the AAV5 construct includes up to residue 197 and the sequence homology around this region is less than 80%, it is not apparent whether helix F in AAV5 will be as extended as in AAV2. Whether these structural and sequence variations found in L_{DB} and the linker region account for the differences in nicking specificity observed for AAV2 and AAV5 Rep proteins remains to be determined and will require structures with DNA substrates along the nicking reaction pathway.^{64,65}

Crystal Structure of OBD-Linker. We and others have described previously that the linker region connecting the OBD and the SF3 helicase domain of the large Rep proteins is pivotal for the assembly of multimeric Rep protein complexes.^{46,47} To better characterize this region, we expressed, purified, and crystallized a construct containing the OBD and the linker that we termed OBDL, spanning residues 1–224. OBDL crystals diffracted to 2.6 Å and belong to space group $P2_1$, with 11 molecules in the asymmetric unit. The electron density corresponding to the C-terminus of helix F and the linker region varies among the different molecules in the asymmetric unit; nevertheless, we were able to build a model of OBDL up to residue 214 (Figure 4). We did not detect any electron density from residues 215–224. The structure shows that helix F spans to residue 205 and is followed by a loop that makes a downward turn toward the core of the protein at proline 209 (Figure 4B). Superposition of OBD with OBDL shows that helix F is flexible and moves as a rigid body with respect to the core of the protein, pivoting around the loop that connects it with strand $\beta 5$. While the two OBD structures superimpose with an overall rmsd of ~ 0.8 Å, the two αF helices superimpose with an rmsd of only 1.3 Å and appear at a different angle with respect to the core of the protein, with a difference between them of $\sim 2.4^\circ$ (Figure 4C). The OBDL structure also implies that the linker is not in an extended conformation but, instead, has a particular structure that is constrained by the two prolines in the linker. This particular conformation may bring the two domains together in the context of full-length Rep68. To test the importance of the proline residues, we mutated proline 214 to alanine and measured the effect of this on Rep68's properties. Figure 4D shows that this mutation drastically alters the oligomeric behavior of Rep68 and affects its ability to form the 13S oligomer species (Figure 4D). Moreover, the mutation affects the ability of Rep68 to bind double-stranded DNA containing a Rep binding site (RBS). Figure 4E shows that the

Figure 5. SAXS analysis for OBD and OBDL. (A) SAXS scattering data from OBD (blue) and OBDL (red) in solution. Superimposed is the scattering calculated from the atomic models for OBD (■) and OBDL (△). (B) Normalized pair distribution functions $P(r)$ for OBD (black) and OBDL (red) calculated from the experimental scattering curves using the program PRIMUS. (C) Low-resolution envelopes of DAMMIN and GASBOR models for OBD (left) and OBDL (right) are superimposed with the X-ray structure of OBD and OBDL, respectively.

Rep68-P214A mutant binds DNA ~ 7 times less than wild-type protein. Taken together, our results show that the conformation of the linker is critical for the function of Rep68.

SAXS Studies of OBDL Validate the Conformation of the Linker. To gain further structural information about the linker domain and to validate the OBDL X-ray structure, we performed solution studies on OBD and OBDL. The concentration of NaCl in the buffer was kept at 0.5–1 M to prevent formation of oligomers induced by the presence of the linker.⁴⁶ The calculated sedimentation coefficient for both constructs is ~ 2.0 S (Figure S2). The small difference in S value suggests that the linker region is not in an extended conformation. However, the smaller sedimentation velocity coefficient of OBDL implies a slightly more elongated shape. We subjected both constructs to SAXS studies under conditions that generated single species, as determined by sedimentation velocity. Scattering profiles from three different concentrations were collected, and a final scattering curve was obtained by merging the best curves using the program Almerge.⁶⁶ The scattering profiles for OBD and OBDL are shown in Figure 5A. The merged data was used to calculate both the radius of gyration (R_g) and the $P(r)$ distribution function as described in the Material and Methods (Figure 5B). *Ab initio* models were produced using the programs DAMMIN and GASBOR. For each protein, 10 independent models were generated and averaged with Damaver. The final models from the two independent reconstructions were aligned using the program Supcomb, resulting in an overall normalized spatial discrepancy (NSD) value of 0.54, suggesting an excellent correlation between the two independent reconstructions.⁶⁷ The resulting GASBOR models for OBD and OBDL are shown

Table 2. Hydrodynamic Parameters

parameter	OBD (1–208)	OBDL (1–224)	Rep40 _{wt} (225–536)	Rep68Y224AΔ (1–490)
S (s) ^a	2.1	1.97	nd ^d	nd
R_g (nm) ^b	21.5 ± 0.3	23.2 ± 2.3	27.6 ± 3.8	38.9 ± 0.1
$D_{\max}$ (nm) ^c	68.2	74.4 (104.96)	98.8	123.8
χ	1.0 ^e	1.9 ^e	0.5 ^f	0.8 ^g

^aDetermined from sedimentation velocity data. ^bDetermined from Guinier analysis. ^cDetermined from $P(r)$ analysis. ^dnd, not determined.

^eExperimental/model data fit using FoXS.⁷⁴ ^fExperimental/model data fit using EOM. ^gExperimental/model data fit using CORAL.

Figure 6. SAXS analysis of Rep40/Rep68 C-terminal tail. (A) Shape reconstruction of Rep40_{wt} and the docked X-ray structure of Rep40. (B) Comparison of the R_g distribution of the initial EOM generated pool (black line) and the selected model ensemble (red line). (C) Scattering curve of Rep40_{wt} (black) and fit of the optimal EOM model ensemble (red). (D) Superposition of the ensemble generated model showing the preferred conformations of the C-terminal tails.

in Figure 5C with the superimposed X-ray structures of the corresponding constructs. The calculated R_g and $D_{\max}$ values are similar for both models (Table 2). The OBDL GASBOR *ab initio* model shows an envelope that resembles a round cylinder, suggesting that, under our experimental conditions, the region between 208 and 224 is not extended but may resemble the overall conformation seen in the OBDL crystal structure (Figure 5C, right panel). To further corroborate the hypothesis that the linker is neither extended nor highly flexible, we carried out molecular dynamics using BILBOMD to determine if an ensemble with multiple conformations of the linker described the SAXS scattering profile of OBDL.⁵⁷ The results show that inclusion of more than one conformation does not improve the fit to the experimental data significantly (data not shown).

Modeling Flexibility of C-Terminal Tail of Rep68. AAV Rep proteins Rep40 and Rep68 have a C-terminal tail of 46 residues that extends from the core of the helicase domain, as

determined by limited proteolysis experiments.³⁸ To determine the overall flexibility of the C-terminal tail, we carried out SAXS experiments of Rep40_{wt} at different concentrations, and data were analyzed as described previously. A GASBOR model of Rep40 clearly resembles the overall shape of the helicase domain and has a characteristic flat disk shape with an additional elongated density that is not represented in the X-ray structure, which should correspond to the C-terminal tail (Figure 6A). However, the dimensions are smaller than those expected from a fully extended C-terminal tail. To analyze the flexibility of the tail, we used EOM software to search for an optimal ensemble of conformations that will best fit the scattering data. This method generates a large pool of models (10 000) with random conformations and uses a genetic algorithm to select an optimized ensemble of configurations that best describes the SAXS scattering data (Figure 6B).^{56,68} The best ensemble consisted of four models that fit the experimental curve with a χ of 0.5 (Figure 6C). However, the

Figure 7. SAXS modeling of Rep68. (A) Fit of the experimental SAXS data (black circles) with the theoretical scattering profile obtained from CORAL model (red line) with a χ value of 0.8. (B) Two views of the GASBOR averaged molecular envelope for Rep68 Δ showing the approximate dimensions in angstroms. (C) Docking of OBDL and Rep40 atomic structures to the GASBOR *ab initio* envelope. (D) Porod–Debye plot of Rep68Y224A Δ SAXS data (black squares) supporting a compact protein with little flexibility. Red line represents the linear plateau. (E) Three views of the CORAL rigid-body final model of Rep68Y224A Δ .

narrowness of the ensemble R_g distribution suggests that a limited number of conformations are preferred, with one configuration in particular accounting for about 50% of the population. Interestingly, all of the selected models have the C-terminal oriented toward the opposite site of the presensor 1 β -hairpin (PS1 β H), a motif that is involved in DNA interactions during DNA translocation and unwinding (Figure 6D).^{30,69} Thus, our study shows that the C-terminal tail, although flexible, has a preferred set of dominant conformations.

SAXS-Based Structural Modeling of Rep68 Shows a Compact Conformation in Solution. We previously showed that Rep68 has complex dynamic oligomeric behavior in solution and is present as a mixture of multiple oligomers, including heptameric and octameric rings.⁴⁵ To obtain a homogeneous population of Rep68 monomers, we took advantage of the Y224A mutation that affects its tendency to form multiple oligomeric species.⁴⁶ The Rep68Y224A protein is present as a monomer at concentrations less than 5 mg/mL and under high

salt conditions. Moreover, we have determined that this mutation has helicase activity comparable to that of the wild-type protein and does not cause any significant structural change.⁷⁰ To facilitate model building and data interpretation, we used a truncated Rep68 (1–490) to eliminate the C-terminal tail residues.³⁸ SAXS data was collected at 2, 3, and 4 mg/mL and processed as described in the previous sections (Figure 7A). The GASBOR generated model produces an elongated prolate ellipsoid that is slightly curved with dimensions $140 \times 73 \times 37 \text{ \AA}^3$ (Figure 7B). Individual atomic structures of OBDL and Rep40 can be easily docked into the envelope; in particular, one end of the particle has the shape of a flat disk that resembles the GASBOR Rep40 *ab initio* model (Figure 7C). We performed rigid-body modeling using the X-ray structures of OBDL (1–214) and Rep40 (225–490) with CORAL.⁷¹ The domain boundaries define an unstructured linker spanning residues 215–224. However, secondary structure prediction suggests that the region between 220 and

Figure 8. Rep68 DNA interacting face. (A) Comparison of Rep68 model (left) and SV40-LTag (right). Colored equivalently are the OBD domain (green) and the helicase domain (blue). (B) (left) Rep68 structure showing the regions that interact with DNA colored in pink. Orientation of Rep68 is 90° counterclockwise from that in panel A. (right) Surface representation of Rep68 showing electropositive regions in blue and negative regions in red. View is in the same orientation as that on the left.

224 could form a α -helix extending into the first helix of the helicase domain.⁴⁶ Consequently, we generated multiple models that extended this helix at different positions. The Rep68 models fit the experimental data very well, with χ values in the range of 1.1–0.8. All models show that the two domains are positioned such that the long axis of the helicase domain is almost perpendicular to the OBD, as shown in Figure 7E. The best model was obtained by extending the N-terminal helicase helix to residue 223. Two main observations can be drawn from this model. First, the orientation of the two domains results in a Rep68 structure where the motifs that interact with DNA in both the OBD (Loop_{DB} and helix C) and helicase domain (β -hairpin 1) are on the same face of the protein. Second, the two domains are closely positioned, making an extended linker structure unlikely. To further support this conclusion, we assessed interdomain flexibility of Rep68 using two methodologies. First, a Porod–Debye plot of the scattering data shows a plateau that is a signature for a compact molecule.⁷² The data fits the linear region with a Porod coefficient of 4, again

consistent with a compact molecule (Figure 7D). In addition, we carried out BILBOMD and determined that a single model fits the data equally as well as with multiple conformations (Figure 7A).

DISCUSSION

Our results show new structural features that increase our knowledge of the architecture of AAV Rep proteins. The combined X-ray and SAXS studies show that the AAV2 Rep68 linker region is partially structured, with helix F of the OBD extending until residue 209 and protruding from the main core at a 45° angle. The presence of two proline residues (209 and 214) seems to impart a certain rigidity to the loop region, making a small turn before continuing toward the helicase domain (Figure 4). This conformation is important for Rep68 function, as mutation of one of the proline residues in the linker (P214) is sufficient to prevent proper oligomerization and DNA binding. In addition, part of the remaining linker may extend the first helix of the helical domain of Rep52/40.⁴⁶ Both

of these structural features bring the OBD and helicase domain closer to each other. However, although the linker is not extended, it allows for a certain degree of flexibility. Our rigid-body modeling generated different models with slightly different conformations of the two domains that fit the scattering data equally well (Figure S3). This suggests a certain degree of conformational flexibility between the two domains. Structurally, this flexibility originates from the linker, the OBD helix F, and the helical bundle of the helicase domain. Alignment of the OBD and OBDL structures shows that helix F can pivot and move relative to the main core of the domain (Figure 4C). Likewise, alignment of the three Rep40 molecules found in the asymmetric unit of the X-ray structure (PDB ID: 1s9h) shows the helical bundle at different positions relative to the AAA⁺ domain (data not shown). This flexibility is important to accommodate changes occurring upon DNA binding and oligomerization. Moreover, analysis of the flexibility of the C-terminal tail in Rep68 suggest that there is a preference in the conformations acquired by the C-terminal tail that positions it at the opposite end of the β -hairpin. This is important in the context of formation of Rep68 oligomeric rings because other conformations may produce steric clashes and inhibit their formation.

Finally, our Rep68 model closely resembles the configuration of the helicase and OBD in one of the molecules seen in the X-ray structure of the SV40 large T antigen (SV40-LTag) in complex with DNA.⁷³ This arrangement results in critical residues that interact with DNA being located on one face of the protein (Figure 8B). In our Rep68 model, these regions include loop L_{DB} and helix α_D in the OBD, the helicase domain (PS1 β residues K404 and K406), and linker residues R217 and K219 that have been shown to play a role in complex formation and DNA binding in AAV5.⁴⁷ In addition, the SV40-LTag structure shows that the helical bundle (Zn domain in SV40-LTag) interacts with DNA, docking into the major groove.⁷³ Our model shows that the position of the helical bundle is similar, and it is intriguing to postulate that the conformation of Rep68 is prealigned to interact with DNA in way similar to that of SV40-LTag (Figure 8A).

In conclusion, we have completed the first structural description of full-length AAV2 Rep68 protein, revealing the orientation and relative position of its functional domains. These observations provide new clues to explain its DNA binding mode; however, our model does not discard possible changes that should occur upon DNA binding and/or oligomerization. These questions require future structural studies of high-resolution structures of Rep68 oligomers alone and in complex with DNA.

■ ASSOCIATED CONTENT

● Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.biochem.5b00610.

Sequence and structural alignment of the AAV OBD serotypes, sedimentation velocity profiles of AAV2 OBD and OBDL, and superposition of different Rep68 models obtained by independent CORAL runs (PDF).

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Funding

This work was supported by NIH grant R01-GM092854 (C.R.E.) and UK MRC grant 1001764 (R.M.L.).

Author Contributions

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Notes

The authors declare no competing financial interest.

Accession Codes

PDB ID Codes: 4ZO0, 5DCX

■ ACKNOWLEDGMENTS

We would like to thank members of NSLS beamlines X6a (Vivian Stojanoff, Jean Jakoncic, and Edwin Lazo) and X9 (Lin Yang and Marc Allaire) and ALS SIBYLS beamline (Kevin Dyer and Kathryn Burnett) for help in data collection. We thank Vishaka Santosh for helpful comments on the preparation of this manuscript.

■ ABBREVIATIONS

AAV, adeno-associated virus; OBD, origin binding domain; OBDL, origin binding domain with linker; HUH, histidine–hydrophobic–histidine; DBD, DNA binding domain; SF3, superfamily 3 helicase; SAXS, small-angle X-ray scattering; ORF, open reading frame; ITR, inverted terminal repeat; RBS, Rep binding site; ssDNA, single-stranded DNA; AAA⁺, ATPases associated with various cellular activities; IPTG, isopropyl- β -D-thiogalactopyranoside; LB, Luria–Bertani; TCEP, tris(2-carboxyethyl)phosphine; NTA, nitrilotriacetic acid; MPD, 2-methyl-2,4-pentanediol; NSLS, National Synchrotron Light Source; R_g , radius of gyration; PEI, polyethylenimine; FACS, fluorescent activated cell sorting

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