HELIX-STRAIGHTENING AS AN ACTIVATION MECHANISM IN THE GELSONIN SUPERFAMILY OF ACTIN REGULATORY PROTEINS*

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Running head: Helix-straightening in the gelsolin superfamily

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Villin and gelsolin consist of six homologous domains of the gelsolin/cofilin fold (V1 through V6, and G1 through G6, respectively). Villin differs from gelsolin in possessing at its C-terminus an unrelated seventh domain, the villin headpiece. Here we present the crystal structure of villin domain V6 in an environment in which intact villin would be inactive: absence of bound Ca2+ or phosphorylation. The structure of V6 more closely resembles that of the activated form of G6, which contains one bound Ca2+, rather than that of the calcium ion-free form of G6 within intact inactive gelsolin. Strikingly apparent is that the long helix in V6 is straight, as found in the activated form of G6, as opposed to the kinked version in inactive gelsolin. Molecular dynamics calculations suggest that the preferable conformation for this helix in the isolated G6 domain is also straight in the absence of Ca2+ and other gelsolin domains. However, the G6 helix bends in intact calcium ion-free gelsolin in order to allow interaction with G2 and G4. We suggest that a similar situation exists in villin. Within the intact protein, a bent V6 helix when triggered by Ca2+, straightens and helps push apart adjacent domains to expose actin-binding sites within the protein. The sixth domain in this superfamily of proteins serves as a keystone that locks together a compact ensemble of domains in an inactive state. Perturbing the keystone initiates reorganization of the structure to reveal previously buried actin-binding sites.

Actin is crucial to such processes as cell movement, cell division and apoptosis, which are regulated by numerous actin-binding proteins, including gelsolin, Arp2/3 and profilin [reviewed in (1)]. Gelsolin, the most potent actin filament-severing protein known, can bind to, sever, cap and nucleate actin filaments in a calcium, pH, ATP and phospholipid-dependent manner [reviewed in (2)]. Villin, found in microvilli of absorptive epithelium, is a second member of the gelsolin family of actin-binding proteins. In addition to standard gelsolin-type activities, villin is able to bundle actin filaments, and is subject to regulation by tyrosine phosphorylation, as well as by Ca2+ and phosphatidylinositol 4,5-bisphosphate [reviewed in (3)]. Many comparisons have been made between gelsolin and villin. The two share 50% amino acid sequence identity and show similar proteolytic cleavage patterns (4). Both contain six similarly-folded domains, but villin possesses a seventh domain at its C-terminus, the headpiece domain (HP), which folds into a compact structure that introduces a second F-actin binding site into the protein. Recent studies indicate that villin uses the HP actin-binding site through a "hinge mechanism" (6).

Biochemical and structural studies have revealed eight Ca2+-binding sites of two types in gelsolin [reviewed in (7)]. Each of the six domains contains a complete and evolutionarily conserved site, termed type 2, while G1 and G4 provide partial Ca2+ coordination at interfaces with actin through sites termed type 1. Sequential mutagenesis of these sites in villin has identified six functional Ca2+-binding sites (8): two major...
sites, one each of type 1 and type 2, in V1, plus four type 2 sites in V2-V6. The type 1 site in V1 regulates F-actin capping and severing activities, while the lower affinity type 2 site in V1 only affects severing (9). The other four sites are involved in stabilizing villin conformation, but do not directly influence actin-severing activity. NMR studies of a fragment of villin that consists of V6 and the HP domain have implicated V6 residues Asn647, Asp648 and Glu670 in binding Ca$^{2+}$ (10). These experiments also revealed the first 80 residues of V6 to undergo significant conformational change as a result of Ca$^{2+}$-binding.

Nanomolar to micromolar concentrations of free Ca$^{2+}$ govern the actin-binding activities of gelsolin. In contrast, micromolar and millimolar concentrations of calcium ions, respectively, are required for villin to exhibit capping and severing. However, after tyrosine phosphorylation, villin can sever actin filaments even at nanomolar Ca$^{2+}$ concentrations (11). Furthermore, while the actin-severing ability of the N-terminal half of villin is calcium-dependent, that by the N-terminal half of gelsolin is not. In contrast, the binding of G-actin by the C-terminal half of both villin and gelsolin requires Ca$^{2+}$. Creation of hybrid proteins demonstrated that the domains of villin and gelsolin are not interchangeable (12).

Abundant X-ray crystallographic structural information exists for gelsolin, including the calcium ion-free (Ca-free), inactive structure of the intact protein (13), the activated N- and C-terminal halves, each in a bimolecular complex with actin (7,14), and the activated C-terminal half on its own (15,16). Structural data for intact villin is unavailable and is limited to fragments V1 (17), solved using NMR methods, and the HP domain, solved by NMR and X-ray crystallography (18,19). NMR experiments also indicate that HP is connected to V6 by a 40-residue disordered linker. As a result, HP has been proposed to bind actin independently of the remainder of the protein (10).

In this report, we present the structure of Ca-free, isolated villin V6, which exhibits a typical gelsolin domain fold. The long helix in V6 in this structure is straight, unlike the corresponding helix in G6 of intact Ca-free gelsolin, which is bent, and only straightens on calcium activation of the intact protein. Hence, V6 appears to be in an active conformation in the absence of Ca$^{2+}$. Molecular dynamics simulations indicate that the preferred state of the long helix is also straight for isolated G6 in the absence of Ca$^{2+}$. Furthermore they suggest a bistable mechanism of helix conformational change regulated by the presence of the remaining domains, by calcium ions and by other interactants. We therefore propose a mechanism for the gelsolin-family proteins, whereby Ca$^{2+}$ triggers the straightening of the domain 6 helix in the native conformation of the inactive proteins in order to propagate more widespread conformational changes.

**Experimental Procedures**

**Protein production** – Villin domains V4-V6 with an N-terminal 8-histidine tag was expressed in E. coli from the expression vector, pSY5, and purified through sequential Ni$^{2+}$-affinity and gel filtration chromatographies (see supplemental information file). Actin was purified from rabbit skeletal muscle by a modified method of Spudich and Watt (20). Villin V4-V6 in 10 mM Tris-HCl, 150 mM NaCl and 5 mM CaCl$_2$, pH 7.5, was added to actin in 2 mM Tris-HCl, 0.2 mM CaCl$_2$, 0.2 mM ATP, 1 mM DTT, pH 7.6-7.8 (buffer A), to a molar ratio of 1:1. The resulting solution was held at 4 °C overnight prior to gel filtration (Bio-Rad Sephacryl S300; 90 x 2.5 cm), with elution by buffer A. The fractions containing the protein complex were pooled and concentrated to 10 mg/ml, as determined by UV absorbance at 280 nm (Perkin Elmer Lamda 4B) using a calculated absorption coefficient of 1.4 ml mg$^{-1}$ cm$^{-1}$.

**Crystallization and data collection** – Crystals were grown at 4 °C using the hanging drop vapor diffusion method. The protein solution (10 mg/ml) was mixed in a 1:1 ratio (v/v) with a reservoir solution containing 15% (w/v) PEG8000, 100 mM Na acetate buffer, pH 5.0. Crystals required 3-4 weeks of incubation to grow to a suitable quality for diffraction analysis. Prior to X-ray data collection, the crystals were transferred into a cryoprotectant solution, 25% (v/v) glycerol, 15% (w/v) PEG8000, 100 mM Na acetate buffer, pH 5.0, and flash frozen in liquid nitrogen.

Diffraction data initially were collected to a resolution of 3.0 Å using a Rigaku MM007HF rotating Cu-anode source with OSMIC VariMaxHR mirrors and a MAR345 image plate detector at the UBC Centre for Blood Research.
(Vancouver, Canada). Data were indexed, integrated and scaled using HKL2000 software (21). Subsequently, data were collected to a resolution of 2.0 Å using an ADSC Quantum-315 CCD detector on beamline BL31B1 at NSRRC (Taiwan, ROC; supplemental Table 1).

Structure solution and refinement— Structural analysis was initiated by molecular replacement using single domains G4 through G6 (from PDB ID 1P8X) as a model. An unambiguous solution showed that the diffraction originated solely from V6, with the V4 and V5 domains and actin not evident. The crystallographic asymmetric unit contains two V6 domains, which permits calculation of a solvent content of 55%. Refinement was carried out using the CCP4 suite of crystallographic programs (22).

Molecular dynamics simulations— To monitor the relaxation of G6 isolated from the context of intact Ca-free gelsolin, five simulations were initiated from residues 640-731 (G6) excised from the inactive structure of gelsolin (PDB ID 1D0N) (13). Two control runs, meant to assure that the long helix does not bend spontaneously in the isolated domain, were initiated from residues 640-742 (G6) excised from the activated form of the C-terminal half of gelsolin (PDB ID 1H1V) (14). All simulations were performed using GROMACS v.3.3.3. The force field used was OPLS-AA, with time steps of 2 fs, at a temperature of 300 K, with Berendsen pressure coupling and a pressure of 1 bar. Water was simulated explicitly using the SPC model, and Na+ cations were added as counter ions, as needed to render the system electrostatically neutral. The water box size was set to 0.9 nm from the protein surface. Energy minimization lasted 6 ps, and frames were sampled every 100 ps.

RESULTS

Structural comparison of V6 with G6— Fig. 1A represents the structure of V6 determined at 2 Å resolution. Like gelsolin G6, V6 consists of a five-stranded β-sheet at its core sandwiched between a pair of α-helices, a long helix that runs roughly parallel and a short helix that lies perpendicular to the strands. There is no calcium ion bound to V6 in this structure. Since the amino acid sequence of V6 is approximately 50% identical with that of G6, the folding and functions of these two domains invite direct comparison. Of the six domains of gelsolin, the structure of V6 most closely resembles that of G6 in its calcium ion-bound (Ca-bound) form (Fig. 1C) (root-mean-square deviation of Cα positions, RMSD = 3.0 Å). The long helix in each of V6 and the Ca-bound form of G6 is in an extended state, in contrast to the kinked state observed in G6 in the context of its intact Ca-free parent protein (Fig. 1E). Comparison of the arrangement of the residues important in binding Ca2+ in G6 (Ser646, Asn647, Asp670 and Glu692; Fig. 1D) with that of the homologous residues from V6 (Ser625, Asn626, Asp648 and Glu670; Fig. 1B) emphasizes the structural similarity to Ca-bound G6. By contrast, in Ca-free G6, the positions and interactions of these residues are altered significantly (Fig. 1F). These comparisons suggest that the present structure of V6 isolated from the rest of the villin domains is that of an activated state despite the absence of bound Ca2+.

V6 interaction interfaces— G6 is the central domain in inactive Ca-free gelsolin, forming direct contacts with all other domains (13). G6 interacts most extensively with G2 and G4 (Fig. 2A). Of particular note is a continuous β-sheet between G4 and G6 that seals the G4-G6 latch, obscuring the actin-binding site on G4. While binding Ca2+ triggers disruption of this latch, it causes only subtle changes at the G6 interface with G4 (compare Figs. 2B & 2C). There is on the surface of V6 (Fig. 2D) a region sufficiently similar to that on Ca-free G6, as portrayed in Fig. 2B, to suggest that interaction with the edge of the core β-sheet of V4 may occur, as observed between G6 and G4. However, models for the loop that exits the domain 4 strand indicate small steric clashes with both Ca-bound G6 (Fig. 2C) and Ca-free V6 (Fig. 2D; supplemental Fig. 1), suggesting that binding of domain 6 to domain 4 would be unfavorable in these two conformations. Surface residues at this interface, including residues involved in the steric clash, show conservation within, yet divergence between, gelsolins and villins (supplemental Fig. 1). Taken together, these data suggest the manner in which V6 interacts with V4 in villin to be analogous to that in which G6 interacts with G4 in gelsolin, yet to display aspects of uniqueness. Most notably, Ca-free V6 in isolation adopts an
activated conformation.

In inactive gelsolin, electrostatic interactions between a negatively charged surface on G6, which includes Asp670, with Arg168 and Arg169 (in G2) (Figs. 2A & 3A) fasten the tail latch over the F-actin binding site of G2 (14,23). Coordination of a Ca\(^{2+}\) by G6 requires participation of the side chain of Asp670 (Fig. 2A & 3A) onto the structure of Ca-free G6 bound to G2 (Fig. 3A), coordinates excised from PDB ID 1D0N) allows the effects of conformational variation in domain 6 in gelsolin family members to be assessed with regard to interaction with domain 2. The AB loop within domain 6 adopts different conformations in the presence of the straight versus kinked version of its long helix (compare Figs. 1A & 1C with Fig. 1E). Movement of the AB loop to go from the Ca-free to the Ca-bound form of G6 is sterically incompatible with binding of G6 to G2 (compare Figs. 3A & 3B), and likely is a major factor in the separation of G6 and G2 during activation of gelsolin by Ca\(^{2+}\). Similarly, the AB loop in Ca-free isolated V6 shows steric clashes in the model presented in Fig. 3C, suggesting a similarity in the activation mechanisms for gelsolin and villin. During deactivation, movement of the AB loop may be needed, in combination with bending of the long helix, for V6 to adopt an inactive conformation within whole villin.

In the G4-G6/actin complex (PDB ID 1H1V), there is interaction between the AB loop of G6 and an \(\alpha\)-helical section of peptide chain (residues 307 through 321) in actin subdomain 3 (Figs. 4A & 4B). This contact arises during the actin filament-severing process and contributes to disruption of actin-actin contacts during that process (14,23). Superimposition of the coordinates for Ca-free G6 from inactive intact gelsolin onto those for Ca-bound G6 in the activated structure of the G4-G6/actin complex (Fig. 4D) reveals how a transformation of G6 in this complex to a Ca-free state would disturb the interaction with actin. However, superimposition of the coordinates of Ca-free V6 onto those for Ca-bound G6 in G4-G6/actin suggests that contact between V6 and the actin helix could be maintained (Fig. 4C). From this, we infer that domain 6 from this family of proteins, once freed of interactions that prevent straightening of its long helix, may bind actin and induce severing at suboptimal calcium concentrations.

**DISCUSSION**

The data presented here are consistent with the actin filament-binding, severing and capping mechanisms of gelsolin and villin being very similar. The role of binding Ca\(^{2+}\), in the context of the whole protein, is to release interdomain contacts. One consequence of this is the straightening of otherwise bent \(\alpha\)-helices, which we speculate aids in opening latches that would otherwise block actin-binding surfaces.

The kink in the helix of the Ca-free G6 can be understood in terms of the context in which it was discovered, folded within the compact globular structure of intact Ca-free gelsolin (13)
The evidence above enforces predictions from sequence alignments that the structures of gelsolin and the V1-V6 body of villin are homologous. As such, the long helix of V6 within Ca-free villin may well be bent. During activation, the extended \( \beta \)-sheet shared between G4 and G6 and the interaction between G2 and G6 (and V4, V2 with V6) are torn apart. The energy cost to do so may be recouped from the binding of calcium ions (Fig. 1D), the formation of new contacts, such as between G5 and G6 (14,23), and by the straightening of the kinked helix of G6, and by analogy, V6.

While helix-straightening is shared among gelsolin family members during activation of their abilities to sever and cap actin filaments, differences among the proteins contribute to unique aspects of their functions. For example, activation of the villin headpiece domain to bundle actin filaments involves a hinged motion away from the gelsolin-like portion of the protein (6). How this is coupled with other stages of activation remains to be explained.

Millimolar calcium ion concentrations, such as those present in blood plasma, activate gelsolin to be extremely efficient at severing undesirable extracellular actin filaments. Such concentrations of free Ca\(^{2+}\) are not attainable intracellularly, yet gelsolin is able to regulate actin filaments in this environment. In addition, at low pH, gelsolin is able to sever actin in the complete absence of free Ca\(^{2+}\). Furthermore, villin has quite different calcium requirements for activation to those of gelsolin (8,9). The importance of Ca\(^{2+}\) in activation of members of this protein superfamily appears to vary with both protein identity and environmental setting, despite the high degree of conservation in Ca\(^{2+}\)-binding residues.

The structure of Ca-free V6 presented in this report and its similarity to that of Ca-bound G6 suggest an explanation for the diversity of activating conditions. V6 and G6 may adopt active conformations in the absence of Ca\(^{2+}\), when isolated from the other domains comprising their respective proteins, as a result of an inbuilt mechanism that involves their long helices. Any process, driven by Ca\(^{2+}\) or otherwise, that promotes release of domain 6 from the remaining domains could allow domain 6 to adopt an active conformation and permit the whole protein to interact with actin. We suggest that the long helix in domain 6 and, by analogy, the one in domain 3, within this superfamily of proteins can be triggered to straighten under a variety of circumstances.

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FOOTNOTES

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1The abbreviations used are: V1, villin domain 1; G1, gelsolin domain 1; HP, villin headpiece; Ca-free, calcium ion-free; Ca-bound, calcium ion-bound.

2The atomic coordinates for the crystal structure of villin domain V6 have been deposited in the RCSB Protein Data Bank with PDB ID 3FG7.

3The on-line version of this article (available at http://www.jbc.org) contains supplemental material in a file named supp.pdf.

FIGURE LEGENDS

Fig. 1. Structural comparison of villin V6 (Ca-free) with gelsolin G6 (Ca-bound and Ca-free). A. Schematic representation of isolated Ca-free V6. B. Key residues that are involved in the putative V6 calcium-binding site, viewed from below with respect to Fig. 1A. C & D. The Ca-bound form of gelsolin G6, displaying a straight helix, taken from the Ca-bound form of G4-G6 (PDB ID 1P8X). E & F. The Ca-free form of G6 revealing a kinked helix and a translocated AB loop, taken from the structure of whole plasma gelsolin (PDB ID 1D0N). The Ca²⁺-binding residues are dislocated in the absence of Ca²⁺ in comparison to Fig. 1D. A hydrogen bond between Asn647 and Arg702 links the bending of the helix to movement of the AB loop. Protein representations were generated here, and in the figures that follow, using PYMOL (http://pymol.sourceforge.net/).

Fig. 2. Comparison of the domain 4-interaction surfaces of G6 and V6. A. Representation of Ca-free gelsolin showing the central position of G6 within the inactive structure. The black sphere pinpoints the unoccupied Ca²⁺-binding site within G6. B. Surface charge representation of G6 from Fig. 2A, rotated
approximately 90° around the vertical axis, showing the interaction surface with G4 via a common β-sheet. C. Ca-bound G6, and D. Ca-free V6 superimposed on G6 in Fig. 2B. These models show minor steric clashes with the G4 loop that follows the β-sheet interaction in Fig. 2B, while the alternating charge distribution at the β-strand edge is maintained.

Fig. 3. Comparison of the domain 2-interaction surfaces of G6 and V6. A. Arg168 and Arg169, from G2, bind to Asp670 and a negatively charged surface on G6 in the Ca-free form of gelsolin. The AB-loop is distant from G2 in this conformation. B. Ca⁡²⁺-binding by G6 involves direct coordination to Asp670. Superposition of Ca-bound G6 on Fig. 3A reveals that the AB-loop moves to clash with G2 in this model. C. Superposition of Ca-free V6 on Fig. 3A creates a model that demonstrates that Asp648 is similarly placed to activate villin via Ca⁡²⁺-binding or to secure the Ca-free structure through binding to V2. In this model the AB-loop clashes with domain 2 and, as such, would have to move to bind V2.

Fig. 4. Domain 6 interactions with actin. A. The structure of G4-G6/actin (PDB ID 1H1V) shows that the actin helix comprising residues 307-321 contacts the AB-loop of G6. B. Surface charge representation of Ca-bound G6 showing the interaction with actin residues 307-321. Ca-bound G6 is oriented as in Fig. 1C. C. Superposition of Ca-free V6 onto Fig. 4B produces a model that suggests that V6 might retain the ability to interact with these actin residues. D. Similar modeling of the transformation of G6 to a Ca-free state disturbs that interaction.

Fig. 5. The straightening of the G6 helix in isolation. A. Superposition of backbone configurations in 10 ns of a molecular dynamics trajectory in which the initial structure was the active form of G6 (from PDB ID 1H1V). B. Superposition of backbone configurations in the last 10 ns (after relaxation to the straightened helix conformation) of a molecular dynamics trajectory, shown in C, in which the initial structure was the inactive form of G6 (from PDB ID 1D0N). C. Fit of the long helix to the initial, bent conformation (green), and to the straightened helix (from PDB ID 1H1V) (red), over a 15 ns molecular dynamics simulation that started with the conformation from PDB ID 1D0N. As a control, the fit to the straight helix (from PDB ID 1H1V) is also shown for a simulation starting from the same structure (blue). In this trajectory, produced in the absence of other gelsolin domains and Ca⁡²⁺, the long helix relaxes toward the crystallographic structure of activated G6 (from PDB ID 1H1V) in less than 10 ns.
Figure 1

A

N

C

AB loop

B

Glu670

Asn667

Asp648

Ser625

Asp680

Asn626

C

N

C

Glu692

Glu689

Asp670

Ser646

D

Arg702

Asn647

E

Glu692

Glu689

Asn647

Asp670

Ser646

F
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