

Competition between Intradomain and Interdomain Interactions: A Buried Salt Bridge Is Essential for Villin Headpiece Folding and Actin Binding

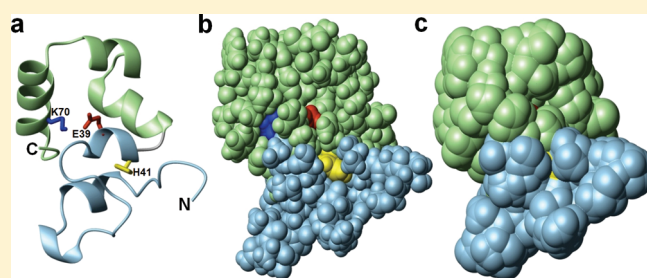
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ABSTRACT: Villin-type headpiece domains are ~70 residue motifs that reside at the C-terminus of a variety of actin-associated proteins. Villin headpiece (HP67) is a commonly used model system for both experimental and computational studies of protein folding. HP67 is made up of two subdomains that form a tightly packed interface. The isolated C-terminal subdomain of HP67 (HP35) is one of the smallest autonomously folding proteins known. The N-terminal subdomain requires the presence of the C-terminal subdomain to fold. In the structure of HP67, a conserved salt bridge connects N- and C-terminal subdomains. This buried salt bridge between residues E39 and K70 is unusual in a small protein domain. We used mutational analysis, monitored by CD and NMR, and functional assays to determine the role of this buried salt bridge. First, the two residues in the salt bridge were replaced with strictly hydrophobic amino acids, E39M/K70M. Second, the two residues in the salt bridge were swapped, E39K/K70E. Any change from the wild-type salt bridge residues results in unfolding of the N-terminal subdomain, even when the mutations were made in a stabilized variant of HP67. The C-terminal subdomain remains folded in all mutants and is stabilized by some of the mutations. Using actin sedimentation assays, we find that a folded N-terminal domain is essential for specific actin binding. Therefore, the buried salt bridge is required for the specific folding of the N-terminal domain which confers actin-binding activity to villin-type headpiece domains, even though the residues required for this specific interaction destabilize the C-terminal subdomain.



Villin is an actin bundling protein found in the brush border microvilli located at the apical surface of the cells that compose the gastrointestinal and renal absorptive epithelium. It contains two actin binding sites, one of which is contained in the C-terminal “headpiece” domain.^{1,2} Headpiece domains are structurally and functionally independent modular domains.³ They are compact motifs containing ~70 residues. Villin headpiece (HP67) is 67 residues in length and contains two subdomains, an N-terminal subdomain (P10–H41) and a C-terminal subdomain (L42–F76), that folds independently from the N-terminal subdomain.³ Residues responsible for actin binding are found in both subdomains.^{4–7} Buried within the continuous hydrophobic core between the N- and C-terminal domains of HP67 is a salt bridge between E39 and K70 (Figure 1). This buried salt bridge is a highly conserved feature among essentially all known headpiece domain sequences. The presence of a buried polar interaction is unusual for a protein of this small size. It has been shown that as the size of a protein decreases, the probability of a charged group being buried in the hydrophobic core also decreases.⁸

Despite its small size and continuous hydrophobic core, *in vitro* studies have shown this small protein exhibits multistate

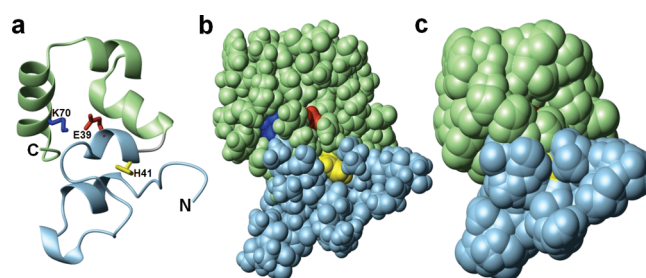


Figure 1. Buried salt bridge in HP67. (a) Ribbon representation of the crystal structure of HP67 (1YU5). The C-terminal subdomain is colored in light green, and the N-terminal subdomain is colored in light blue. Side chains of the buried salt bridge residues, glutamate 39 and lysine 70, are highlighted in red and blue, respectively. The buried histidine residue in the N-terminal subdomain is shown in yellow. (b) Space-filling representation using van der Waals radii. (c) Solvent accessible surface showing burial of E39, H41, and K70.

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unfolding. First, the less stable N-terminal subdomain unfolds, followed by the thermostable C-terminal subdomain.⁹ Indeed, the C-terminal subdomain is stable as an isolated 35-residue peptide (HP35).¹⁰ Linking the two subdomains within the continuous hydrophobic core is the highly conserved buried salt bridge between E39 in the N-terminal subdomain and K70 in the C-terminal subdomain.

Previous studies have shown that histidine 41, which is buried in the hydrophobic core of the N-terminal subdomain, acts as a pH-dependent unfolding switch.^{9,11,12} Protonation of H41 results in unfolding of the N-terminal subdomain while the C-terminal subdomain remains folded. Mutation of H41 to tyrosine (H41Y) eliminates the sequential pH-dependent unfolding of the N-terminal subdomain and stabilizes the C-terminal subdomain.⁹ Despite the increase in overall stability, the H41Y mutant still exhibits multistate unfolding, with unfolding of the N-terminal subdomain preceding the C-terminal subdomain.⁹

Salt bridges in proteins can be energetically favorable or unfavorable.^{13–18} If the salt bridge is primarily solvent accessible, its contribution to stability may be small. If the salt bridge is largely inaccessible, its contribution can be large.^{19,20} In a statistical analysis of salt bridges in proteins, Sarakatsannis and Duan have shown that the most frequent percentage solvent accessible surface area (PSASA) for salt bridges is greater than 20% with a maximum at 32%.²¹ Using their criteria, the salt bridge in HP67 is much more buried than the average salt bridge with only 2.3% PSASA. Another trend noted by these authors and others¹⁷ is that, by far, the most common separation for partners involved in salt bridges was four residues (e.g., one turn of α -helix), where as the residue separation in HP67 is 31. Thus, while most salt bridges in proteins are very local in the sequence, whether buried or exposed, the salt bridge in HP67 involves residues widely separated in the primary sequence.

In addition to their role in protein stabilization, salt bridges can provide specificity for a unique fold.^{8,9,13} There are a limited number of orientations for favorable charged residue packing. In terms of achieving energetically equivalent structures, packing of hydrophobic residues is generally less stringent. Structural specificity provided by a salt bridge can result in the proper formation of a discrete protein fold resulting in the proper ligand binding surface.^{13,15,22,23}

To investigate the role of the highly conserved buried salt bridge in HP67, we created a series of mutants that targeted the E39–K70 interaction. These mutants include replacement of both of residues of the salt bridge with flexible, hydrophobic methionine residues, single residue replacements, and a double mutant which swapped the partners of the salt bridge. We used NMR to assay the structure, circular dichroism spectroscopy (CD) to measure the stability, and actin sedimentation assays to determine the effect on the biological function of these mutants. We find that the wild-type salt bridge is absolutely essential for folding of the N-terminal subdomain, and no mutations were tolerated. The salt bridge is not essential for the C-terminal subdomain fold. Sedimentation assays reveal that none of the mutants exhibited significant actin binding activity. Thus, proper folding of the N-terminal subdomain is essential for activity. Furthermore, the headpiece–actin binding energy is insufficient to fold the N-terminal subdomain.

EXPERIMENTAL PROCEDURES

Cloning, Expression, and Purification of Proteins. The HP67- and H41Y-containing pET24a vectors used for cloning

and expression of the salt bridge mutants have been previously described.^{3,11} They encode the last 67 residues of villin headpiece (HP67) corresponding to residues 760–826 of chicken villin. All salt bridge mutants were made with the QuikChange mutagenesis kit (Stratagene). Pairs of primers were designed to mutate: E39M, E39K, K70M, and K70E. All mutations were confirmed by DNA sequencing. All constructs were expressed in *Escherichia coli* BL21(DE3) cells (Novagen) as described.³ Labeling with ¹⁵N isotope was accomplished as described by Marley et al.²⁴ Protein concentrations were determined by absorbance at 280 nm using extinction coefficient of 5690 M^{−1} cm^{−1} (6990 M^{−1} cm^{−1} for the H41Y constructs).²⁵ Supravillin headpiece was a gift from Jeffrey W. Brown.²⁶

F-actin was purified from chicken pectoral muscle using standard procedures and stored dialyzing against F-buffer (10 mM TRIS, 1 mM MgCl₂, 100 mM NaCl, 0.1 mM ATP, 0.2 mM DTT, 3 mM NaN₃, 0.1 mM CaCl₂, pH 8.0) at 4 °C.^{7,27}

Nuclear Magnetic Resonance (NMR) Spectroscopy. All NMR experiments were conducted on a Bruker DMX 500 MHz spectrometer. All samples were prepared in 10% D₂O containing 10 mM sodium phosphate at pH 7, uncorrected for the effects of D₂O. Chemical shifts were referenced to trimethylsilylpropionic acid. One-dimensional NMR spectra were acquired using Watergate pulse sequences to suppress water.²⁸ For ¹⁵N-heteronuclear single quantum coherence (HSQC) spectra, protein concentrations were ~1 mM, and all spectra were taken at 25 °C with 128 t1 increments of 2048 data points. The data were processed using NMRPipe²⁹ and analyzed in NMRView.³⁰

Circular Dichroism (CD) Spectroscopy and Unfolding Experiments. CD spectra were acquired on an Aviv 62DS spectrometer equipped with a Peltier temperature controller and automated titration apparatus. A 10 mm path length cell was used for all CD experiments. Wavelength spectra are the average of three scans of 20 μ M protein in 10 mM phosphate buffer, pH 7, from 200 to 250 nm with an averaging time of 20 s at each wavelength in 1 nm steps at 25 °C. For thermal, GuHCl, and urea denaturation experiments the CD signal was monitored at 222 nm. Thermal denaturation was carried out from 20 to 98 °C, with 2 deg increments, and protein concentrations of 5 μ M in 10 mM phosphate buffer, pH 7.0. The first derivative of the smoothed thermal unfolding data was used to determine the T_m value.

GuHCl and urea unfolding experiments were performed with a titrator unit interfaced with the spectrometer at 25 °C and protein concentrations of 10 μ M in 10 mM phosphate buffer at pH 7.0. Urea unfolding was done in the same buffer at pH 6.5. Concentrations of urea and GuHCl were determined from the refractive index of the solution.³¹ Concentrations of urea and GuHCl increased from 0 to 9.8 M and 0 to 8.5 M in 0.2 M increments, respectively. To determine thermodynamic parameters ΔG° , C_m, and m values, chemical denaturation curves were fit as described by Santoro and Bolen³² with Origin 6.0 (Microcal Software, Inc., Northampton, MA).

Actin Sedimentation Assays. Actin sedimentation assays were performed at 4 °C as previously described with 15 μ M F-actin and 0–200 μ M headpiece domain.^{7,12} After incubation for 1 h in F-buffer (10 mM TRIS, 1 mM MgCl₂, 100 mM NaCl, 0.1 mM ATP, 0.2 mM DTT, 3 mM NaN₃, 0.1 mM CaCl₂, pH 8.0), the binding reactions (50 μ L) were spun at 100000g for 1 h and the supernatants were decanted. Pellets were washed with 48 μ L of F-actin buffer and then resuspended in 48 μ L of 7% acetic acid. Samples were stored at −20 °C. The bound protein

concentrations in the actin pellets were determined by UV quantitation during reverse phase high-performance liquid chromatography (HPLC).³³ A C18 column (Vydac) 4.6 × 200 mm was run at 1 mL/min in water–acetonitrile gradients with 0.1% trifluoroacetic acid and monitored at 220 nm. A standard curve was constructed by running samples of known concentrations of headpiece. Binding data were fit with Origin 6.0 as previously described.⁷ Three parameters were obtained by fitting: (1) equilibrium dissociation constant (K_D), (2) maximal binding (B_{max}), (3) nonspecific (NS) binding. The data points used in fitting were the average of three separate sedimentation experiments.

RESULTS

To investigate the role of the highly conserved buried salt bridge in villin headpiece (Figure 1), we made a series of point mutations at positions E39 and K70. To determine whether the

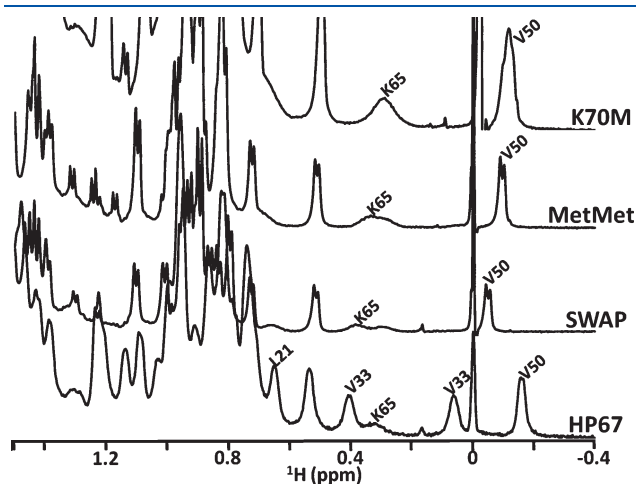


Figure 2. Upfield region of the ^1H 500 MHz 1D NMR spectra of HP67, SWAP, MetMet, and K70M. The methyl resonance of V50 and the $\text{H}\beta$ of K65 in the C-terminal subdomain and the methyl groups of V33 and L21 in the N-terminal subdomain are labeled. Spectra were recorded at 25 °C in 10% D_2O , 10 mM sodium phosphate buffer at pH 7.0.

buried salt bridge could be replaced by a hydrophobic interaction without introducing much steric hindrance, we created E39M/K70M, or “MetMet”, a double mutant. To determine whether the salt bridge could still be formed in the opposite orientation, we created E39K/K70E, termed “SWAP”. In the process of making the double mutants, all possible single mutants were also constructed (i.e., E39K, E39M, K70E, K70M).

NMR Indicates That the Salt Bridge Is Essential for the N-Terminal Subdomain Fold. 1D proton NMR spectra were obtained to determine the effect of double mutations on the headpiece fold. The upfield region of the ^1H 500 MHz NMR spectrum of HP67 contains well-resolved resonances from both the N- and C-terminal subdomains (Figure 2). The methyl groups of L21 and V33 from the N-terminal subdomain and V50 as well as the $\text{H}\beta$ of K65 from the C-terminal subdomain act as probes for folded structure within the two individual domains. In all of the salt bridge mutant spectra there is a loss of the L21 and V33 methyl peaks, indicating that the N-terminal subdomain is unfolded. The retention of the V50 and K65 peaks indicates the C-terminal subdomain remains folded. The small changes in chemical shift of the V50 and K65 peaks are due to localized changes in conformation within the C-terminal subdomain and are consistent with the chemical shifts observed in the isolated HP35 domain.

NMR Indicates That the C-Terminal Subdomain Retains the Native Fold. To further define the folded and unfolded regions within the salt bridge mutants, we mapped the chemical shift changes of backbone amide protons relative to HP67 in the ^1H – ^{15}N HSQC spectra. Figure 3 shows the HSQC spectra of SWAP and MetMet (red) overlaid with HP67 (black). The peak labels indicate the assignments for HP67. Unlike the spectrum of HP67, the spectra of the mutants contain more resonances than expected from their sequences, indicating some regions are slowly exchanging between two states. In addition, there are more peaks in the region corresponding to random coil chemical shifts (8–9 ppm) in the mutants than in HP67. What is clear from these spectra is that the well-dispersed peaks from residues in the N-terminal of HP67 are absent in the spectra of salt bridge mutants (e.g., L13, E14, L19, V20, L21, A25, E27, L29, G32, D34, R37, L42, S43). In contrast, most of the disperse peaks

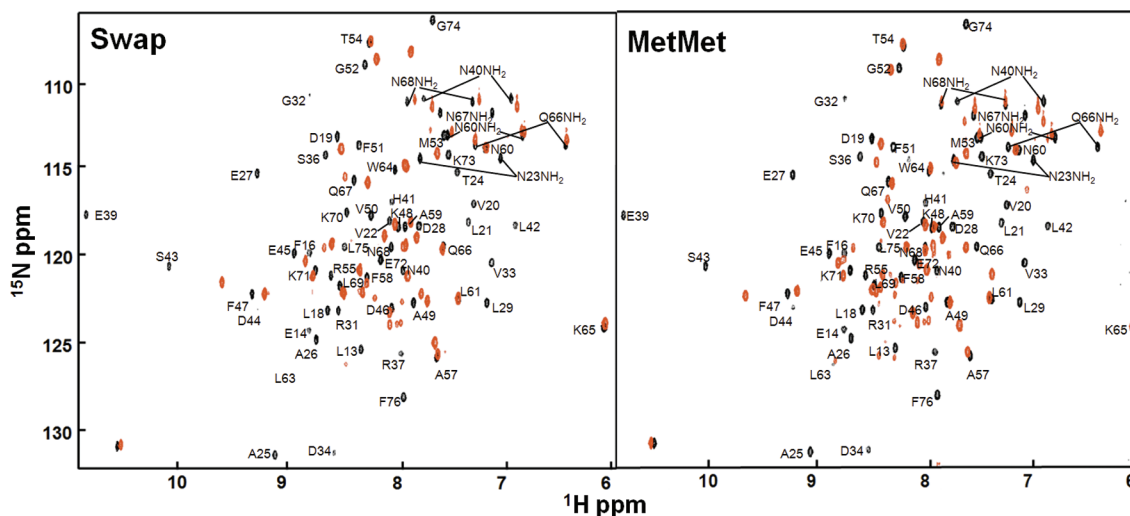


Figure 3. Comparison of the ^{15}N -HSQC spectra of Swap and MetMet with HP67. The spectrum of HP67 is shown in black, and the assignments are labeled.³ The spectra of SWAP and MetMet are shown in red on the left and right spectrum, respectively. The spectra were acquired at pH 7.0, 20 °C, in 10 mM phosphate buffer. The protein concentrations for both SWAP and MetMet are 1 mM.

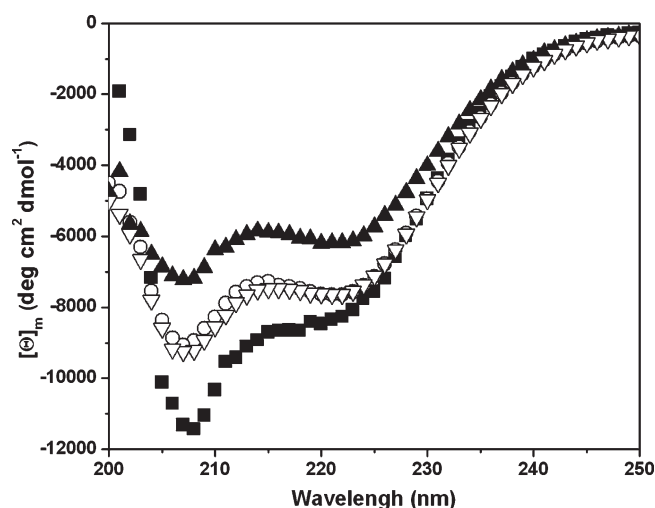


Figure 4. CD spectra of HP67 and salt bridge mutants. Far-UV scans of 10 μ M protein at 25 $^{\circ}$ C in 10 mM sodium phosphate buffer at pH 7.0 in a 10 mm path-length cell. HP67 (■), MetMet (▲), SWAP (○), and K70M (▽).

arising from residues within the C-terminal subdomain have a corresponding peak directly under, or very close to, those from HP67 in the spectra of the mutants (e.g., A49, A57, T54, L61, K65, Q66). The resonances from residues corresponding to the end of the C-terminal helix of HP67 are also shifted (e.g., K70, G74, F76). However, these shifts show a close correspondence to those seen in the isolated HP35 subdomain.¹⁰ These observations are consistent with the unfolding of the N-terminal domain while the C-terminal domain adopts the structure the HP35 subdomain.

Although Unfolded, the N-Terminal Subdomain Influences the Stability of the C-Terminal Subdomain. The C-terminal subdomain of HP67 contains most of the helical structure in HP67, and therefore the signal from the C-terminal subdomain dominates the CD spectrum. The far-UV CD spectra for all the salt bridge mutants exhibit significant helical content, similar to that observed for HP67, which is consistent with an independently folded C-terminal subdomain (Figure 4).

The thermal stability of the C-terminal subdomain was determined by monitoring the helical CD signal at 222 nm as a function of temperature. HP67 exhibits a thermal unfolding temperature (T_m) of 79.8 $^{\circ}$ C. Despite having unfolded N-terminal subdomains, both mutants containing the K70 M mutation (MetMet and K70M) exhibit increases in T_m compared to that of wild-type (93.6 and 83.5 $^{\circ}$ C, respectively). In contrast, SWAP exhibits a T_m of 67.5 $^{\circ}$ C, a decrease of 12.3 $^{\circ}$ C in T_m . The hierarchy of T_m values is MetMet > K70 M > HP67 > SWAP.

To determine the thermodynamic stability of the folded C-terminal subdomain of each construct, we performed chemical denaturation as a function of increasing guanidine hydrochloride (GuHCl, Figure 5) and urea (Supporting Information Figure S1). Helical content was monitored by the CD signal at 222 nm. As expected from their elevated T_m values, MetMet and K70M mutants both show an increase in stability relative to HP67. All constructs, except SWAP, show cooperative unfolding with similar m -values. In the SWAP mutant, unfolding cooperativity was slightly, but significantly, reduced.

The GuHCl unfolding curves were fit to obtain ΔG° , C_m , and m values (Table 1).^{31,34} SWAP was the only mutant which

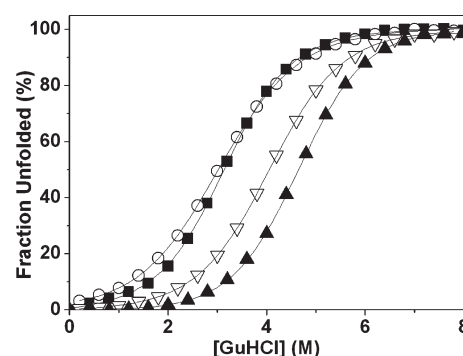


Figure 5. Guanidine hydrochloride denaturation of headpiece constructs. GuHCl-induced denaturation of HP67 (■), MetMet (▲), SWAP (○), and K70 M (▽). Experiments were performed with 10 μ M protein samples at 25 $^{\circ}$ C in 10 mM sodium phosphate buffer, pH 7.0. CD signal was monitored at 222 nm in a 10 mm cell. Data were best fit to the Santoro and Bolen equation³⁴ and then plotted as percent unfolded.

Table 1. Thermodynamic Data for the HP67 and H41Y Constructs (GuHCl)

	$\Delta G^{\circ}_{\text{GuHCl}}$ (kcal/mol)	$m_{\text{value,GuHCl}}$ (kcal/(mol M ⁻¹))	$C_{m,\text{GuHCl}}$ (mol)	T_m ($^{\circ}$ C)
HP67	2.6	0.8	3.4	79.8
SWAP	2.2	0.7	3.0	67.5
MetMet	4.0	0.9	4.6	93.6
K70M	3.1	0.8	4.0	83.5
H41Y	3.9	1.1	3.4	83.4
H41Y/SWAP	2.4	0.8	3.4	68.9
H41Y/MetMet	2.8	0.7	3.4	92.2
H41Y/K70M	3.5	0.8	4.4	84.4

exhibits a decrease in both ΔG° and C_m . The relative trends in ΔG° and C_m values for these mutants and HP67 are the same as for the thermal unfolding experiment: MetMet > K70M > HP67 > SWAP. Thus, as has been observed for the isolated HP35 subdomain, replacement of lysine 70 with methionine increases its stability.³⁵

A concern with using GuHCl unfolding data to calculate the stability of these salt bridge mutants is the potential ionic screening arising from the high concentrations of GuHCl. To address this concern, we performed the same experiments in urea (Supporting Information Figure S1). The results from the urea experiments are in qualitative agreement with those in GuHCl, and the stability of the constructs follows the same hierarchy: MetMet > K70 M > HP67 > SWAP. However, a complete thermodynamic analysis in urea is complicated due to the lack of unfolded baselines for most of these constructs.

The Conserved Buried Salt Bridge Is Essential for Specific Actin Binding. To test whether these salt bridge mutations affected the actin binding activity of villin headpiece, we measured the apparent actin–headpiece affinity of SWAP, MetMet, and K70M using a sedimentation assay. Non-actin binding Supervillin headpiece (SVHP) was used as a negative control.^{12,36} The binding constant (K_D), maximum of amount of protein specifically bound (B_{max}), and nonspecific binding (NS) were determined by fitting the binding curves as described previously¹² (Figure 6).

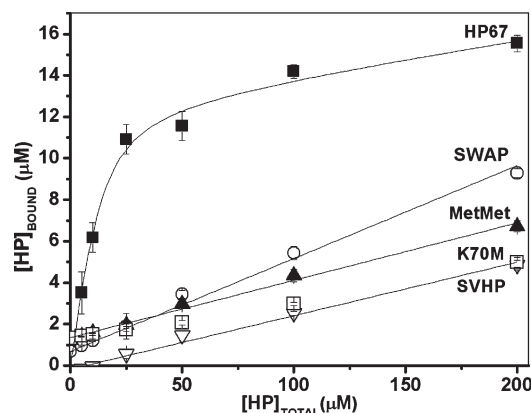


Figure 6. F-actin sedimentation assay. The concentration of the headpiece constructs cosedimenting with F-actin ($[HP]_{\text{bound}}$) is plotted against total headpiece construct concentration ($[HP]_{\text{total}}$) in each reaction. The data points are the average of the three separate experiments. Error bars indicate the standard deviations of the three experiments. The curves and lines are best fit to the average data points. For HP67 (■), a saturating binding equation, which includes nonspecific binding, was used. For MetMet (▲), SWAP (○), K70M (▽), and SVHP (□) the data are best fit to a linear equation corresponding to nonspecific binding of 3%, 4%, 2%, and 2%, respectively.

Under the conditions of this assay HP67 has a K_D of 5 μM , consistent with previous results.^{7,12} The B_{max} value (15 μM) is the same as the actin monomer concentration used in the reaction and is consistent with a 1:1 protein:actin stoichiometry. Unlike the wild-type sequence, mutant headpiece constructs bound only weakly to actin. Mutants accumulated in the pellet fractions in an essentially linear fashion as a function of concentration, without displaying signs of saturation. MetMet, SWAP, and K70M binding data gave poor fits for specific binding. When fit to a straight line, nonspecific binding was determined to be 3%, 4%, and 2% for MetMet, SWAP, and K70M, respectively. Note that the wild-type HP67 domain has a similar nonspecific value of 1% yet exhibited saturation of binding to F-actin.⁷ SVHP had a 2% nonspecific binding in this experiment. Therefore, the salt bridge mutants exhibited similar nonspecific binding values compared to a protein previously known to not bind F-actin. Thus, the correctly folded N-terminal subdomain is essential for specific F-actin binding. Furthermore, the lack of specific binding by any salt bridge mutant indicates that the binding energy of the headpiece–actin interface is insufficient to induce proper folding of the N-terminal subdomain.

Salt Bridge Mutants Remain Unfolded in the More Stable HP67/H41Y Background. In an attempt to induce folding of the N-terminal domain of the salt bridge mutants, the mutations were also made in the background of a stabilized mutant of HP67, H41Y (HP67/H41Y). H41 is highly conserved in headpiece domains; no other substitutions have been found at this site except tyrosine. On the basis of hydrogen exchange experiments, H41Y has increased stability of ~ 0.7 kcal/mol.^{9,11,37} The thermal unfolding (T_m) values for the salt bridge mutants in the H41Y background follow the same hierarchy as found for the wild-type HP67 background (Table 1). The ΔG° , m value, and C_m determined by GuHCl unfolding follow a slightly different pattern (Supporting Information Figure S2). H41Y exhibits the highest ΔG° value followed by K70M/H41Y, MetMet/H41Y, and then SWAP/H41Y. Thus, the increased stability of H41Y background

cannot compensate for the destabilizing effect of the salt bridge mutations to allow proper folding of the N-terminal subdomain.

DISCUSSION

Buried salt bridges are unusual in small proteins, and the observation of a highly conserved and buried salt bridge in HP67 is unexpected.^{17,21} To test the constraints of the salt bridge, we replaced the two residues with hydrophobic interactions or swapped their polarity.

1D and 2D NMR data show that any mutation of this salt bridge results in unfolding of most of the N-terminal subdomain. Residues in the region linking the N- and C-terminal subdomains can still affect the thermostability of the C-terminal subdomain, as seen in the MetMet double mutant, which is more stable than the K70 M single mutant. Thus, it appears that when the buried E39 residue is replaced with methionine in the MetMet double mutant, the hydrophobic side chain can form stabilizing contacts with the C-terminal subdomain.

In this study, we found that any mutations to the residues participating in the buried salt bridge are not tolerated and result in partial unfolding of most of the N-terminal subdomain. Further, unfolding of the N-terminal subdomain disrupts the actin binding face. Thus, the buried salt bridge serves a vital role by specifying the actin-binding conformation of the N-terminal residues.

There appears to be a competition between the requirement for a highly specific salt bridge needed to ensure proper folding of the N-terminal subdomain, on the one hand, and the destabilization of the C-terminal subdomain by K70, on the other. The equilibrium unfolding of HP67 is known to be multistate. The first state is the canonical HP67 fold with both subdomains folded. In the second state, the N-terminal subdomain is unfolded while the C-terminal subdomain adopts the structure of isolated HP35, which contains slight alterations in the structure and dynamics, especially of the C-terminal residues. Another state, in which both domains are unfolded, is populated under more strongly denaturing conditions. Any mutation to the salt bridge favors the competition toward the second state. Thus, the presence of the native E39–K70 salt bridge stabilizes the N-terminal subdomain but destabilizes the C-terminal subdomain. This has some interesting consequences for the study of small domains derived from larger proteins. A commonly employed strategy to stabilize a domain is to use a sequence alignment to derive consensus residues at core positions and to then replace residues in the sequence of interest that deviate from the consensus. Such an approach would erroneously fail to predict that mutation of K70 stabilizes the isolated C-terminal subdomain, HP35. In the present case, K70 is conserved because its stabilizing intersubdomain interactions are more important for function than its destabilizing intrasubdomain interactions.

ASSOCIATED CONTENT

S Supporting Information. Denaturant unfolding of SWAP, MetMet, and K70M mutations in the more stable H41Y background is shown in Figures S1 (urea) and S2 (GuHCl). This material is available free of charge via the Internet at <http://pubs.acs.org>.

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ABBREVIATIONS

ATP, adenosine triphosphate; C_m , the midpoint of the denaturant-induced unfolding transition; CD, circular dichroism; E39K, glutamate 39 to lysine mutation in HP67; E39M, glutamate 39 to methionine mutation of HP67; ΔG° , apparent free energy of unfolding; D₂O, deuterium dioxide; DTT, dithiothreitol; F-actin, filamentous actin; GuHCl, guanidine hydrochloride; H41Y, mutant of HP67 replacing residue histidine 41 with tyrosine; HP35, residues 42–76 of the isolated C-terminal domain of HP67; HP67, 67 amino acid polypeptide encoding residues 10–76 of villin headpiece (residues 760–826 of intact chicken villin); HPLC, high-pressure liquid chromatography; HSQC, heteronuclear single quantum coherence; K70E, lysine 70 to glutamate mutation in HP67; K70M, lysine 70 to methionine mutation in HP67; MetMet, HP67 which contains two mutations, E39M and K70M; NS, nonspecific binding; NMR, nuclear magnetic resonance; PSASA, percent solvent accessible surface area; SVHP, Supravillin headpiece; SWAP, HP67 which contains two mutations, E39K and K70E; T_m , the midpoint of thermal unfolding; TRIS, tris(hydroxymethylamino)methane; UV, ultraviolet; WT, wild-type.

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