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Tuning Protein Autoinhibition by Domain Destabilization

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Abstract

Activation of many multi-domain signaling proteins requires rearrangement of autoinhibitory interdomain interactions that occlude activator binding sites. In one model for activation, the major inactive conformation exists in equilibrium with activated-like conformations that can be stabilized by ligand binding or post-translational modifications. The molecular basis for this model is established for the archetypal signaling adapter protein Crk-II by measuring the thermodynamics and kinetics of the equilibrium between autoinhibited and activated-like states using fluorescence and NMR spectroscopies, together with segmental isotopic labeling *via* expressed protein ligation. The results demonstrate that intramolecular domain-domain interactions both stabilize the autoinhibited state and induce the activated-like conformation. A combination of favorable interdomain interactions and unfavorable intradomain structural changes fine-tunes the population of the activated-like conformation and allows facile response to activators. This mechanism suggests a general strategy for optimization of autoinhibitory interactions of multi-domain proteins.

Many autoinhibited signaling proteins are responsive to small changes in the populations of cognate binding partners^{1–5}. The dynamic equilibrium between autoinhibited and activated-like conformations of signaling proteins in the absence of activators is believed to play a key role in achieving the seemingly contradictory tasks of preventing uncontrolled activation and enabling prompt response to cellular stimuli^{6,7}. This model implies that the interdomain interface responsible for autoinhibition must be sufficiently stable to prevent constitutive activation, but sufficiently labile to allow facile kinetic exchange between autoinhibited and activation-competent states.

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Author Contributions J.H.C. designed and performed all experiments, analyzed the data, and wrote paper. V.M. and M.V.P. contributed to synthesis of the ligands and preparation of the segmentally labeled protein. D.P.R., T.W.M., and A.G.P. designed the experiments, analyzed the data, and wrote paper.

The conformation of a protein in complex with a target molecule must, in the absence of the target, have a free energy equal to or greater than that of the experimentally observed apo structure. Consequently, the binding affinity for the target must be sufficient to overcome the energetic penalty associated with forming higher energy holo conformations⁸. The thermodynamics and kinetics of such conformational changes are critical for understanding the physical principles controlling binding affinity; however, detection of low populations of high energy states in the absence of binding partners represents a considerable experimental challenge.

A full description of the molecular mechanisms responsible for controlling the dynamic equilibrium between autoinhibited (closed) and activated-like (open) states requires comparison of the structures and thermodynamics of individual domains in isolation and in the context of full-length molecules, as well as knowledge of the kinetics of interconversion between the various populated conformational states. Moreover, unique spectroscopic probes are needed for studying the individual domains within the full-length multi-domain protein^{9,10}. Consequently, complete descriptions are not yet available of the molecular basis for control of autoinhibition of multi-domain signaling proteins.

Archetypical signaling adaptor proteins of the Crk family are comprised of Src homology 2 (SH2) and 3 (SH3) domains and have been implicated in many cellular processes including cell motility, proliferation and adhesion^{11–14} (Fig. 1a). Crk proteins mediate diverse protein-protein interactions, through their SH2 and SH3 domains, in downstream transduction of growth and differentiation signals¹¹. Crk-II consists of an N-terminal SH2 domain and two SH3 domains (SH2-nSH3-cSH3). The binding of the N-terminal SH3 (nSH3) domain to C3G, a guanine-nucleotide exchange factor for small molecular weight GTPase¹⁵, Abl¹⁶ and DOCK180¹⁷ contributes to the integrin signaling¹⁸ and the regulation of fibroblast migration¹¹. The biological activity of the nSH3 domain of Crk-II is negatively regulated by the C-terminal SH3 (cSH3) domain *via* intramolecular domain-domain interactions^{11,19}. The solution structure of Crk-II shows that the cSH3 domain stabilizes a closed autoinhibited state of Crk-II in which the ligand binding site of the nSH3 domain is blocked by interdomain interactions with the SH2 domain; however, the active site of the nSH3 domain is not directly occluded by the cSH3 domain¹⁹ (Fig. 1b). Alternative splicing of the *c-crk* gene yields Crk-I (SH2-nSH3), which lacks the C-terminal SH3 domain and is constitutively active, causing transformation of hematopoietic and fibroblast cells^{12,20}. The NMR solution structure of Crk-I indicates that the active site of the nSH3 domain is exposed to solvent¹⁹, consistent with the inability of the SH2 domain alone to exert autoinhibitory control of nSH3 activity⁹. Various models have been proposed for control of autoinhibition of Crk-II^{19,21,22}; however, a complete biophysical analysis of this process, which necessarily involves analysis of the full-length protein and individual domains, has not been reported. Herein, we present a thermodynamic, kinetic and structural analysis that reveals a critical role for selective domain destabilization in tuning the responsiveness of Crk-II to activation.

Results

Thermodynamics and kinetics of a dynamic equilibrium of Crk-II

Crk-II exists as an equilibrium mixture of the autoinhibited state and the open state, in which the binding site of the nSH3 domain is exposed to solvent (Fig. 1c). Target proteins can bind to the nSH3 domain of Crk-II only when the protein adopts an open state. The free energy of the closed autoinhibited state relative to the open activated-like state of Crk-II was determined from the binding affinity of a C3G-derived²³ ligand to the nSH3 domain in Crk-II and Crk-I. The binding affinity (K_{int}) of the ligand to the nSH3 domain in the open state was estimated using Crk-I (**Methods**). The free energy difference, $G^\circ = G^\circ_{open} - G^\circ_{closed} = 1.5 \pm 0.1 \text{ kcal mol}^{-1}$, corresponds to closed and open state populations of $p_{closed} = 92.6\%$ and $p_{open} = 7.4\%$, respectively, for Crk-II (Table 1 and Supplementary Fig. 1).

The kinetics of ligand binding to the nSH3 domain in the context of full-length Crk-II were estimated from an NMR-detected titration experiment using uniformly ^{15}N -labeled Crk-II and fluorescein-labeled C3G-derived ligand. Two ^{15}N resonances in the nSH3 domain, W169 ($\omega_N = \omega_{bound} - \omega_{free} = 790 \text{ s}^{-1}$) and A172 ($\omega_N = 165 \text{ s}^{-1}$), undergo slow and fast chemical exchange, respectively, relative to the NMR chemical shift timescale, upon addition of ligand (Fig. 1d). Therefore, the chemical exchange rate constant is $165 \text{ s}^{-1} < k_{ex} < 790 \text{ s}^{-1}$, corresponding to a bimolecular on-rate constant of $5.8 \times 10^9 \text{ M}^{-1} \text{ s}^{-1} < k_{on} < 2.8 \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}$ for ligand binding to the nSH3 domain in the open state of Crk-II. The observed K_D is 1.9 orders of magnitude lower for fluorescein-labeled C3G ligand than for the unlabeled ligand²⁴. Assuming that k_{off} is unaffected by the presence of the label, $6.5 \times 10^7 \text{ M}^{-1} \text{ s}^{-1} < k_{on} < 3.1 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ for an unlabeled ligand is estimated (**Methods**). These results are in excellent agreement with measured on-rates for ligand binding to isolated SH3 domains²⁵ and thus support use of Crk-I to estimate the intrinsic ligand binding affinity, K_{int} , for the open state of full-length Crk-II (Fig. 1c and Table 1).

Collectively, these results demonstrate that regulation of the interconversion between the open and closed autoinhibited conformations plays a central role in controlling Crk-II function. These data also show that the stability of the interface with the cSH3 domain limits the activation rate of Crk-II, because autoinhibition must be relieved to populate the open state.

Intra- and inter-domain interactions in the closed state

Considerable basal levels (~7%) of the open states of Crk-II persist, even though a large solvent accessible surface area (~1360 Å²) is buried between the cSH3 and other domains in the structure of autoinhibited Crk-II (Fig. 1a). We quantified the energetics of the interdomain interactions by comparing the stability of the cSH3 domain in isolation and in full-length Crk-II. The unfolded cSH3 domain cannot bind to the other domains of Crk-II; consequently, unfolding and dissociation of the cSH3 domain are thermodynamically linked. The effect of ligand binding on global unfolding, by monitoring the change in T_m or H/D-exchange, has been reported for other isolated SH3 domains^{26,27}.

NMR-detected backbone amide H/D-exchange experiments are routinely used to measure the unfolding free energy of small to intermediate sized proteins (<25 kDa); however, peak

overlap makes the application of the H/D-exchange experiment difficult for large (>30 kDa) proteins. We used expressed protein ligation (EPL)²⁸ to produce segmentally labeled full-length Crk-II, in which only the cSH3 domain is labeled with ¹⁵N (SH2¹⁴N-nSH3¹⁴N-cSH3¹⁵N) (Fig. 2a). Owing to the simplified NMR spectrum, all of the expected ¹H-¹⁵N resonances in the cSH3 domain are detected in segmentally labeled full-length Crk-II (~34 kDa) (Fig. 2b and Supplementary Fig. 2).

H/D-exchange rates were measured at 290 K for 20 residues (Fig. 2c), located in and around the hydrophobic core of the cSH3 domain (Fig. 2d). Strikingly, the protection factors are nearly equal or larger in the isolated cSH3 domain (average $G_{\text{ex}} = 5.58 \pm 0.52 \text{ kcal mol}^{-1}$) relative to the values measured for the cSH3 domain within full-length Crk-II (average $G_{\text{ex}} = 5.29 \pm 0.41 \text{ kcal mol}^{-1}$) (Fig. 2c). ¹H and ¹⁵N chemical shifts (Fig. 3a and 3b) and ¹⁵N transverse relaxation rate measurements (Fig. 3c) demonstrate that the similarity in the protection factors of the cSH3 domain in isolation and in the full-length Crk-II are not due to dissociation of the cSH3 domain from the other domains in Crk-II. These results indicate that the cSH3 domain does not experience any net favorable interdomain binding free energy, even though a large surface area is buried in the autoinhibited state.

We characterized the conformational properties of the linker region between the nSH3 and cSH3 domains, residues 224–237, in full-length Crk-II. Heteronuclear {¹H}-¹⁵N NOE values are < 0.6 for residues in the linker, compared with values >0.6 for residues in elements of secondary structure, in full-length Crk-II and are similar to values measured for the isolated cSH3 domain (Fig. 3d). We did not observe any resonance cross-peaks in the first NMR spectrum in H/D exchange experiments for residues in the linker region in full-length Crk-II, indicating that amide proton exchange protection factors are small. We note that segmentally labeled protein must be used to measure accurate NMR H/D exchange and spin relaxation parameters for the linker region owing to the serious spectral overlap in uniformly labeled Crk-II, which prevents quantitative peak intensity measurements (Supplementary Fig. 3). Thus, thermodynamically *unfavorable* interactions must occur simultaneously in the cSH3 domain, and not the linker, upon formation of the thermodynamically favorable domain-domain interface of the autoinhibited state.

Conformational change of Trp275 within the cSH3 domain

We investigated the structural changes in the cSH3 domain that may offset favorable interdomain interactions, and thereby allow facile activation of Crk-II. The backbone ¹⁵N and ¹H chemical shifts of the cSH3 domain in the context of autoinhibited Crk-II and activated Crk-II (Crk-II/C3G) are broadly similar (Fig. 3b), indicating that modest conformational changes around the interdomain interface of the cSH3 domain are sufficient to induce the open conformation of Crk-II. The ¹⁵N^{ε1} resonance for the indole ring of Trp275 of the cSH3 domain has the largest change upon activation (Fig. 3b and Supplementary Fig. 2). Trp275 is highly conserved in Crk-II (Supplementary Fig. 4), and is critical for stabilizing the cSH3 domain and the autoinhibited state of Crk-II^{19,29}. H/D exchange experiments show that side-chain indole ¹H^{ε1} of Trp275 is well protected from solvent in the isolated cSH3 domain (Supplementary Fig. 5a and 5c), but not in full-length Crk-II (Supplementary Fig. 5b and 5d). We performed fluorescence experiments to confirm

that Trp275 is protected from solvent exposure in the isolated cSH3 domain^{30,31}. The Trp fluorescence emission maximum is blue-shifted compared with free indole (Table 2). We also compared the Stern-Volmer constant (K_{sv}) of the isolated cSH3-WT domain to free indole (Table 2). The fluorescence results show that Trp275 has a single conformation in which the side chain is buried into the hydrophobic core. These results imply that domain-domain interactions in full-length Crk-II perturb the hydrophobic core of the cSH3 domain by switching Trp275 to a more exposed conformation.

The solvent-protection of the side chain of Trp275 is consistent with solution structure of the isolated cSH3 domain (PDB code; 2GGR). The indole ring of Trp275 is packed against several hydrophobic residues, resulting in the burial of 1117 Å² of hydrophobic surface area³². In contrast, Trp275 populates a different χ_1 and χ_2 rotamer in full-length Crk-II and the indole ring is flipped toward the binding interface with the SH2 domain (Fig. 4a).

Rotameric mutation of Trp275 using non-coded amino acid

We sought to design a mutant of cSH3 that would mimic the conformation of Trp275 observed in full-length Crk-II. The energetic consequences of local conformational changes of a single Trp residue are difficult to evaluate using traditional mutagenesis, because substitutions conserving size and molecular structure cannot be made using natural amino acids. We selectively replaced Trp275 in the isolated cSH3 domain by the non-natural amino acid 7-aza Trp (7AW), in which the carbon atom in the 7-position of indole ring is replaced by nitrogen. This mutation does not perturb the structure of the side chain, except for the increased polarity of the ring. The 7-position of the Trp275 side chain is buried in the hydrophobic core of the isolated cSH3 and we reasoned that the side chain of 7AW would reorient to expose the 7-position to solvent, mimicking the effects of changing the χ_1 rotamer of Trp275 in Crk-II (Fig. 4a). Thus the 7AW substitution can be used as a “rotameric mutation”. The effect of 7AW on the stability of the cSH3 domain was measured using urea-induced denaturation. The stability of the isolated cSH3-7AW domain was decreased by almost ~1 kcal mol⁻¹ relative to wild type isolated cSH3 (Table 2 and Supplementary Fig. 6). The *m*-value also decreased, indicating that the hydrophobic core is slightly perturbed in cSH3_{7AW}, compared to wild-type cSH3.

We performed fluorescence quenching experiments to confirm that the 7AW substitution perturbed the rotameric conformational state of Trp275^{30,31}. We compared the K_{sv} of the isolated cSH3-7AW domain to free 7-azaindole, because of the difference in intrinsic quenching efficiencies of W and 7AW (Table 2 and Fig. 4b and 4c). The result for the cSH3-7AW domain shows that the protein populates major (~89%) and minor (~11%) conformations corresponding to the buried and exposed conformation of the side chain of 7AW, respectively. In addition, the ¹H-¹⁵N HSQC spectrum of the cSH3-7AW showed that several resonances from residues near 7ATrp275 are split into two peaks, further confirming the presence of two different conformations (**data not shown**). In contrast, the spectrum of the isolated wild type cSH3 domain showed single resonance peaks for each residue. These results demonstrate that the incorporation of 7AW induces the rotation of the side chain toward the surface of the cSH3 domain (Fig. 4a). We used these data in combination with the unfolding free energy of the cSH3-7AW ensemble to estimate an energetic penalty of

2.04 ± 0.38 kcal mol⁻¹ for rotation of Trp275 in the autoinhibited form of Crk-II (for details see Supplementary Methods). The substantial energetic penalty incurred when the side chain of Trp275 flips out to enable favorable interdomain interactions provides an explanation for the absence of stabilization of the cSH3 domain in the context of full-length Crk-II.

Discussion

Quantifying the energetics of the structural changes upon domain-domain interactions is central to understanding signaling processes at the molecular level. We have developed a self-consistent thermodynamic and kinetic model for the control of the autoinhibition of Crk-II in which favorable domain-domain interactions are balanced by structural changes, including perturbations of Trp275, that reduce the stability of the cSH3 domain, in order to avoid over-stabilization of the autoinhibited state (Fig. 4d). The moderate conformational change of the side chain of Trp275 destabilizes the cSH3 domain by a surprisingly large ~ 2 kcal mol⁻¹. However, packing interactions in the hydrophobic core of SH3 domains are highly conserved and domain stability is very sensitive to perturbations of the hydrophobic core³³.

The hypothetical binding energy between the cSH3 and SH2 domains of Crk-II that would be expected if exposure of the Trp275 side chain were not energetically unfavorable is important in assessing whether additional conformational changes contribute to the free energy difference between the closed and open states. The intramolecular binding energy between the two domains can be estimated using the relationship $K_{a,intra} = K_{a,inter} * C_{effective}$, in which $C_{effective}$ is the effective local concentration of the binding domain and $K_{a,inter}$ is the intermolecular binding affinity³⁴. The intermolecular binding constant between the cSH3 and SH2 domains of Crk-II is not known; however, the Abl SH3 domain binds to a proline-rich peptide sequence derived from the DE-loop of the Crk-II SH2 domain with $K_{a,inter} \sim 360$ μ M³⁵. A theoretical investigation of the intramolecular interaction of SH3 domains showed that the effective local concentration is ~ 80 mM when two domains are separated by the optimum linker length³⁴. Thus, the hypothetical intramolecular binding energy, in which the unfavorable interaction of Trp275 is not considered, would be ~ -3.2 kcal mol⁻¹. This result is consistent with our estimation of free energy difference between the closed state and open states in which the Trp275 side chain is buried (Fig. 4d). Thus, the offset of the favorable interdomain interaction (~ -3.2 kcal mol⁻¹) by unfavorable conformational change from buried to exposed conformations of Trp275 (~ 2 kcal mol⁻¹) reduces the free energy difference between the closed and open states to ~ 1.2 kcal mol⁻¹. The self-consistency between this latter quantity and the difference measured independently (Table 1) indicates that the conformational change associated with Trp275 is the dominant contribution to tuning the free energy difference between autoinhibited and activated-like states of Crk-II. The intramolecular binding free energy of the cSH3 domain to other regions of Crk-II is relatively small, compared to other SH3-ligand interactions, perhaps because the intramolecular binding interface of Crk-II is not optimized for ideal binding³⁶.

Two other models have been proposed for control of autoinhibition of Crk-II. In the first model, autoinhibition of chicken Crk-II is controlled by interactions between the nSH3 and cSH3 domain mediated *via* isomerization of Pro237³⁷. However, that work made use of a

truncated Crk construct (nSH3-cSH3) that lacked the SH2 domain. The nSH3 and cSH3 domains do not directly interact in the structure of the full-length human protein¹⁹. In addition, the amino acid sequence adjacent to Pro237 in the chicken Crk-II sequence differs from the corresponding human and mouse sequences (Supplementary Fig. 4). These differences suggest that the regulatory mechanism of chicken Crk-II may be different from that of human and mouse Crk-II. In the second model, the autoinhibited state of human Crk-II is stabilized by rigidification of the linker region (residues 224–236) tethering the nSH3 and cSH3 domains¹⁹. However, heteronuclear NOE data (Fig. 3d) indicate that the linker region experiences extensive conformational fluctuations on picosecond-nanosecond time scales. In addition, the H/D exchange data indicate that the linker region undergoes slower interconversion between associated (exchange non-competent) and dissociated (exchange competent) states with minor free energy differences.

In conclusion, intramolecular domain-domain interactions induce differential effects on the thermodynamic stabilities of individual domains to finely adjust the overall functional response of Crk-II. The interplay of these competing effects acts as the driving force to induce the basal level of the open (activated-like) state required for activation of Crk-II. The thermodynamically unfavorable conformational change of the cSH3 domain upon autoinhibition contributes to timely activation in response to cellular stimuli. The mode of regulation revealed by the present study is likely to be found in other multi-domain proteins. In the limiting case, order-to-disorder transitions may maximally offset favorable domain-domain interactions³⁸.

Methods

Protein Expression and Purification

Detailed procedures for preparation of protein samples are described in Supplementary Methods. Intactness of segmentally labeled Crk-II was confirmed by comparing the NMR chemical shifts and R_2 relaxation rate constants with those of uniformly ¹⁵N labeled Crk-II which was prepared in conventional protein preparation method (Supplementary Fig. 7). Isolated cSH3 domain was confirmed to be monomeric by measuring the concentration dependent NMR transverse relaxation rate constants (Supplementary Fig. 7).

Ligand Binding Assays

A fluorescence-based titration assay was used to measure the affinity constants of both C3G-ligand¹⁵ and a phosphopeptide-ligand³⁹. All ligand binding measurements were performed at 25°C in a stirred 1 cm pathlength cuvette. The excitation wavelength was 295 nm. C3G-ligand, HPPPPLPPKRRK-ε-fluorescein-G-NH₂, was prepared by solid-phase peptide synthesis. Dissociation constants were calculated by assuming the formation of a 1:1 complex and by fitting corrected fluorescence intensities to the equation;

$$F = [\Delta F] \left[P + K_D + L - \left\{ (P + K_D + L)^2 - 4PL \right\}^{1/2} \right] [2P]^{-1} \quad (1)$$

where F and F_{max} are the change and maximum change in protein fluorescence, respectively. P is the total protein concentration, L is the total ligand concentration, and K_D is the equilibrium dissociation constant. Experiments were performed in triplicate.

The free energy difference between the open and closed states of Crk-II was calculated using eq (2). In our three state binding model (Fig. 1c), the equilibrium constant (K_O) for the opening and closing process between the closed and open states of Crk-II was calculated from the apparent binding affinity (K_D) of the C3G ligand and the binding affinity (K_{int}) of C3G ligand:

$$\Delta G^\circ = -RT \ln K_O \quad (2)$$

where, $K_O = K_{int} / (K_D^{apparent} - K_{int})$, $K_{int} = [O][L] / [OL]$ and

$$K_D^{apparent} = ([O] + [C])[L] / [OL] \quad (3)$$

Determination of the bimolecular rate constant, k_{on}

The bi-molecular rate constant, k_{on} , was determined from the exchange rate constant, k_{ex} , estimated from a NMR-titration experiment. The k_{ex} for the ligand binding to the Crk-II is defined by the equation:

$$k_{ex} = K_O k_{on} [L] + k_{off} = k_{on} (K_O [L] + K_{int}) \quad (4)$$

where k_{on} and k_{off} represent bimolecular on-rate and unimolecular off-rate constants, respectively, between the open and ligand-bound states (Fig. 1c), $[L]$ represents the free ligand concentration, and K_{int} represents the dissociation constant of C3G-ligand to Crk-I. The reported k_{on} was calculated with K_O of 0.074 ($p_{open} = 7.4\%$ in Crk-II) and $[L]_{free} = 0.077 \mu\text{M}$ (protein:ligand = 1:0.22). The NMR titration experiment (Fig. 3c) showed that the exchange rate is $165 \text{ s}^{-1} < k_{ex} < 790 \text{ s}^{-1}$ at this protein:ligand ratio resulting in $5.8 \times 10^9 \text{ M}^{-1} \text{ s}^{-1} < k_{on} < 2.8 \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}$. The observed K_D for fluorescein-labeled C3G-ligand is 85-fold lower than fluorescein-free C3G-ligand²⁴. Despite this effect of fluorescein, the label is necessary to quantify the concentration of ligand in solution because the ligand sequence lacks spectroscopic probes and measurements of peptide concentrations using amino acid analysis have large errors. Thus, the observed k_{on} rate was corrected by the ratio of K_D for label-free and fluorescein-labeled ligand. The correction yields estimates of $6.5 \times 10^7 \text{ M}^{-1} \text{ s}^{-1} < k_{on} < 3.1 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$.

NMR spectroscopy

A Bruker DRX600 NMR spectrometer equipped with a CryoProbe was used. Chemical shifts for different protein states and complexes were measured in a buffer containing 10 mM sodium phosphate (pH 7.0), 100 mM NaCl, 2 mM DTT at 298 K. Chemical shift changes were calculated as:

$$\Delta\delta = \sqrt{(\Delta\delta_{HN})^2 + (0.1\Delta\delta_N)^2} \quad (5)$$

The ^1H - ^{15}N NOE spectra were collected for segmentally labeled Crk-II ($\text{SH2}^{14}\text{N}$ - $\text{nSH3}^{14}\text{N}$ - $\text{cSH3}^{15}\text{N}$) and for the isolated cSH3 domain with and without ^1H saturation for 4.0 s at 600 MHz equipped with a CryoProbe. Recycle delays for NOE experiment and control experiments were 3.0 s and 9.0 s, respectively. The delay between NOE and control experiments was 18.0 s. ^1H - ^{15}N HSQC, 3D-HNCO, HNCACB, CBCA(CO)NH, and HN(CA)CO experiments were used for backbone assignment.

Equilibrium Unfolding Measurements

Urea-denaturation experiments were performed at 298 K by monitoring the change in fluorescence signal intensity recorded at 340 nm and 398 nm for wild type cSH3 and cSH3-7AW, respectively. Details of experimental conditions are described in Supplementary Methods.

H/D Exchange Rate Measurements

H/D exchange experiments were performed at 290 K on a Bruker DRX 600 MHz NMR spectrometer equipped with a CryoProbe. Samples for H/D exchange experiments were exchanged into a buffer containing 10 mM sodium phosphate (pD 6.6), 200 mM NaCl, 2 mM DTT (d11) in 100% D_2O , using Micro Bio-spin columns (BIO-RAD). H/D-exchange rates were determined from peak intensity changes in a series of ^1H - ^{15}N HSQC experiments. Reported exchange rates are the average of two or three repeated experiments. Exchange rates for residues whose exchange was dominated by local conformational fluctuations could not be determined, because the intrinsic rate of exchange is too high at the experimental pH (pD = 6.6). The folding rate for the cSH3 domain has previously been measured to be $82.6 \pm 2.9 \text{ s}^{-1,32}$; thus the system is within the EX2 regime.

Fluorescence Quenching Experiments

Steady state quenching experiments were performed at 298K. Samples were prepared in a buffer containing 20mM Na-phosphate (pH 7.2), 100mM NaCl, 3mM TCEP. The concentration of each sample was adjusted to ensure that the OD was 0.1 or less at the excitation wavelength to prevent inner filter effects. Acrylamide was used as a quencher. Observed signal intensity was corrected for both dilution effects and for the inner filter effect upon adding acrylamide using $\text{OD}_{295\text{nm}} = 0.23 \text{ M}^{-1} \text{ cm}^{-1}$. The excitation wavelengths for cSH3-WT (and free indole) and cSH3-7AW (and 7-azaindole) were 295 nm and 310 nm, respectively. Quenching data for homogeneously emitting samples (free indole, 7-azaindole and cSH3-WT) were analyzed using the Stern-Volmer equation³⁰:

$$F_o/F = (1 + K_{sv}[Q])\exp(V[Q]) \quad (6)$$

For cSH3-7AW, the Stern-Volmer equation was re-written to take into account heterogeneous fluorescent components³¹:

$$F/F_o = f_1/(1 + K_{sv,1}[Q]) + (1 - f_1)/(1 + K_{sv,2}[Q]) \quad (7)$$

where F_o and F represent fluorescence intensity in the absence and presence of quencher, Q , respectively. $K_{sv,1}$ and f_1 is the Stern-Volmer constant and population fraction of the

exposed conformation and $K_{sv,2}$ is the Stern-Volmer constant of the buried conformation. V represents the sphere of action. Application of equation 7 to the quenching data of the cSH3-WT domain yielded unrealistic fitting, indicating that equation 6 is adequate for fitting the data.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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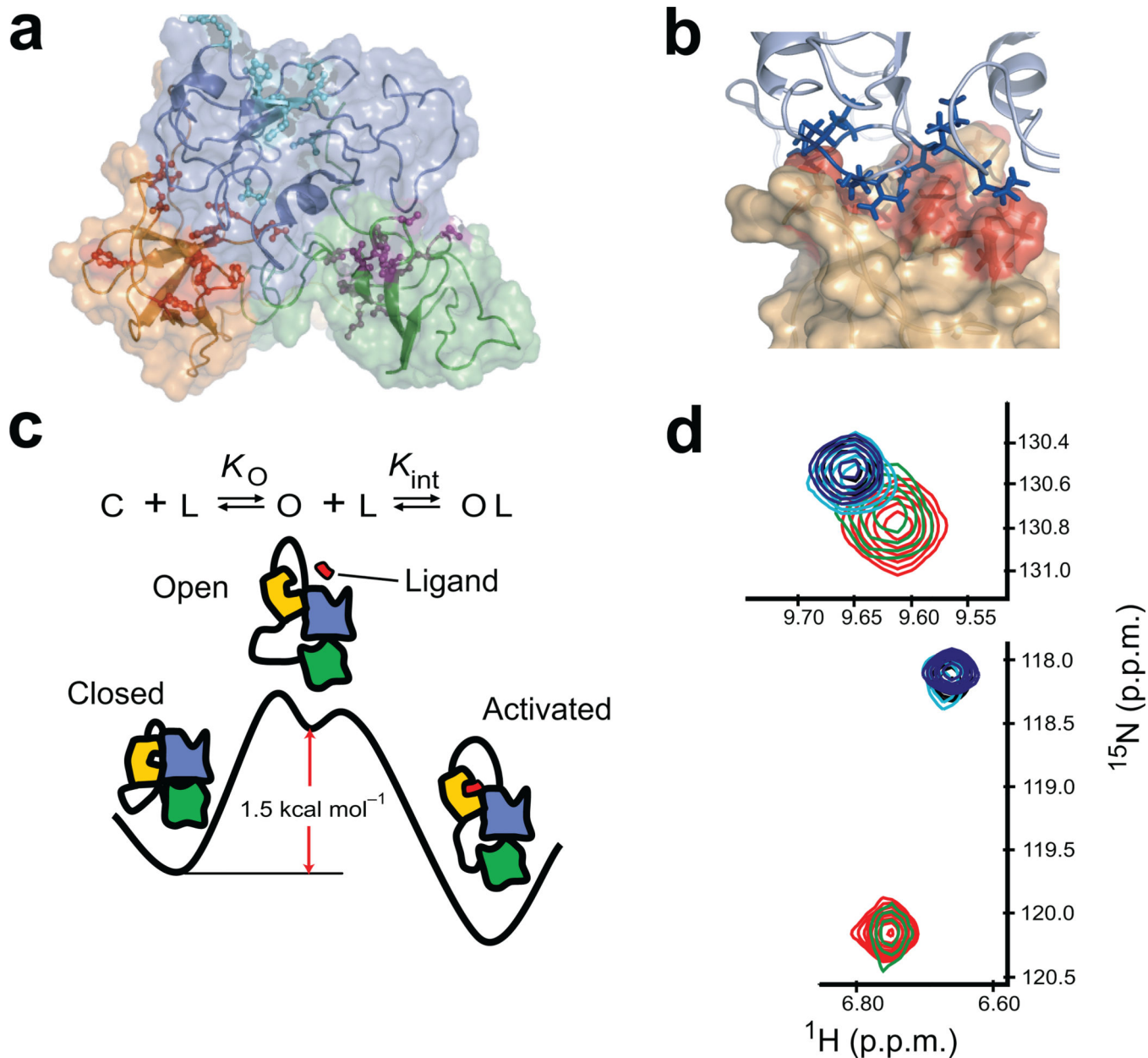


Figure 1. Scheme for the autoinhibition and activation of Crk-II
(a) Solution structure of Crk-II (SH2; blue, nSH3; yellow, cSH3; green) (pdb id; 2EYZ¹⁹). Active site residues of each domain are shown in stick-and-ball format. **(b)** Expanded view of the interface between the active site of the nSH3 and SH2 domain in Crk-II. Active site residues are shown in red on the surface representation of the nSH3 domain. Interface residues of SH2 domain are shown in stick format. **(c)** Schematic representation of a three-state equilibrium of ligand binding to the nSH3 domain in Crk-II (SH2; blue, nSH3; yellow, cSH3; green, ligand; red). K_O is the equilibrium constant between the closed and open state, and K_{int} is the intrinsic binding affinity of the ligand to the nSH3 domain in the open state. C and O represent the closed and open state of Crk-II, respectively. L represents the ligand that binds to the nSH3 domain. **(d)** Change in the chemical shift of W169 (upper panel) and

A172 (lower panel) as a function of C3G-ligand concentration. The ratios of protein and ligand are 1:0 (red), 1:0.22 (green), 1:0.63 (cyan), 1:0.79 (black) and 1:1.19 (blue).

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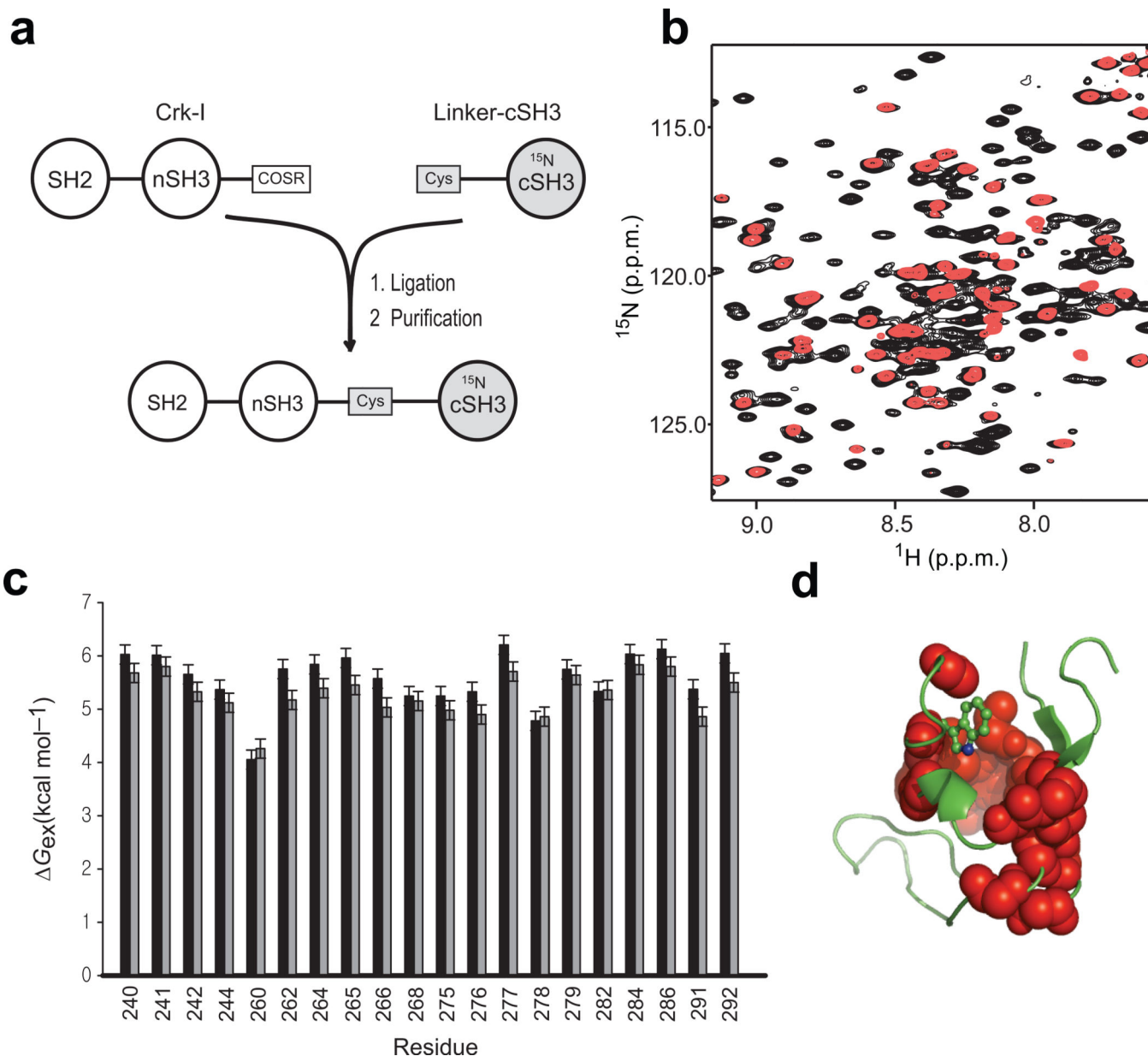


Figure 2. NMR analysis of the cSH3 domain within full-length Crk-II
(a) Schematic diagram outlining preparation of segmentally labeled Crk-II (residues from 208 to 304 are ¹⁵N-labeled). **(b)** Overlay of the ¹H-¹⁵N HSQC spectrum of uniformly ¹⁵N-labeled Crk-II (black) and segmentally labeled Crk-II (red). **(c)** NMR-detected H/D-exchange free energy of unfolding of the cSH3 domain in isolated (black), and Crk-II (grey) The error bars represent standard deviation of three repeated experiments. **(d)** Residues detected in the H/D-exchange experiment are shown as spheres on the structure of the cSH3 domain.

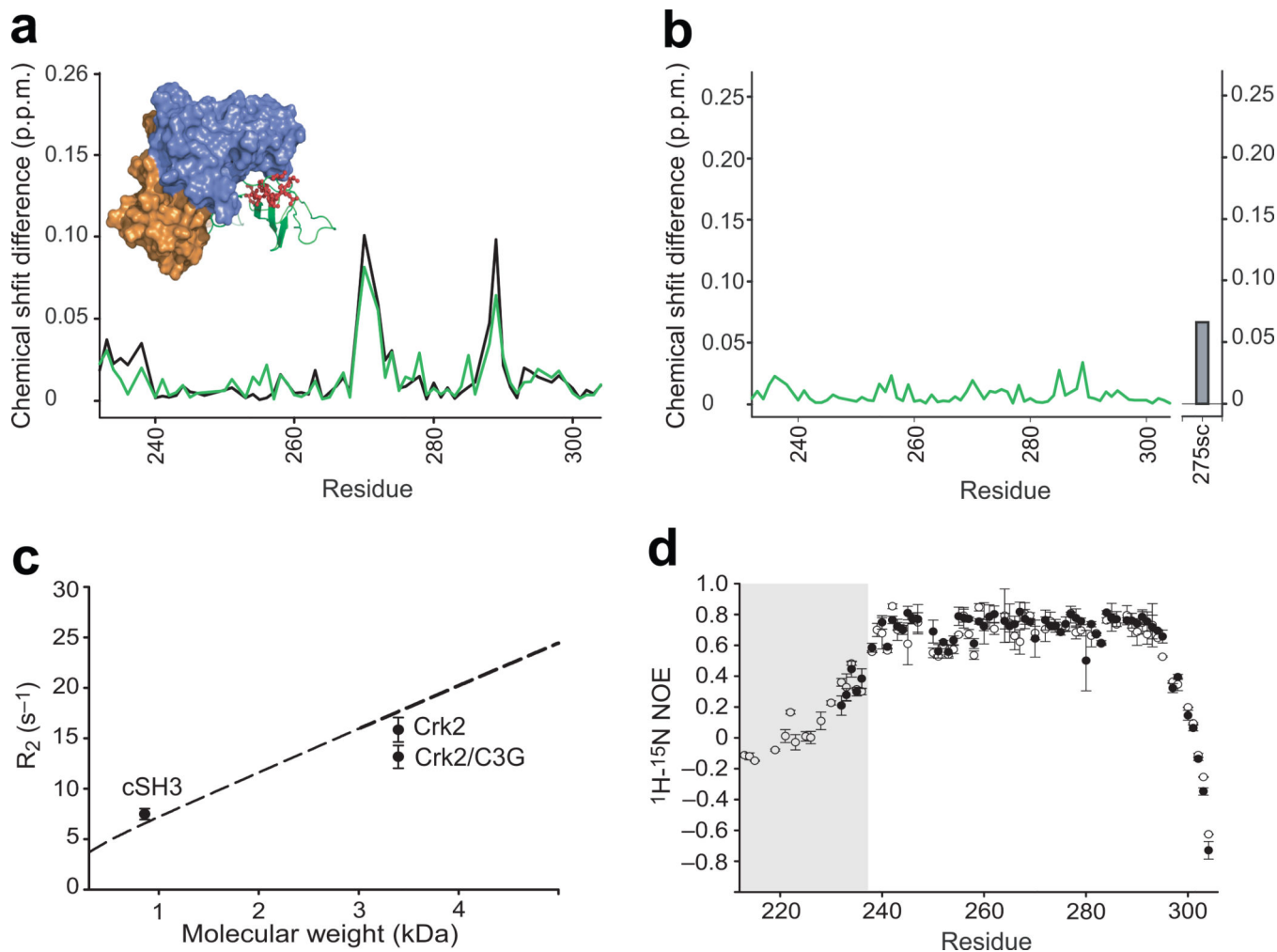


Figure 3. Structural and dynamic analysis of the cSH3 domain within Crk-II

(a) Chemical shift differences (δ) of the cSH3 domain in Crk-II (black), and C3G-ligand bound Crk-II (green), relative to the isolated cSH3 domain. The inset shows the locations of the residues whose resonances changed markedly between isolated cSH3 and Crk-II (top 25% in the deviation plot). (b) Chemical shift differences of the cSH3 domain in C3G-ligand bound Crk-II, compared to those within Crk-II. The difference for the $^{15}\text{N}^{\epsilon 1}$ resonance of Trp275 is shown as a bar. (c) Comparison of average R_2 rates of the cSH3 domain in isolation (Mw=8,569 Da), in Crk-II, and in Crk-II/C3G (~33,830 Da). The dashed line represents the molecular weight dependent R_2 rates calculated using isotropic rotational correlation times calculated from Stokes' law. Error bars represent the standard deviation of average R_2 . (d) $\{^1\text{H}\}$ - ^{15}N heteronuclear NOE measurements for the cSH3 domain in isolation, residues 232–304, (closed circles) and within Crk-II, residues 208–304, (open circles). The residues corresponding to the linker region, residues 208–236, are shaded in the plot. Error bars represent the propagated uncertainties of two repeated experiments. The uncertainty was estimated using background noise of the spectrum.

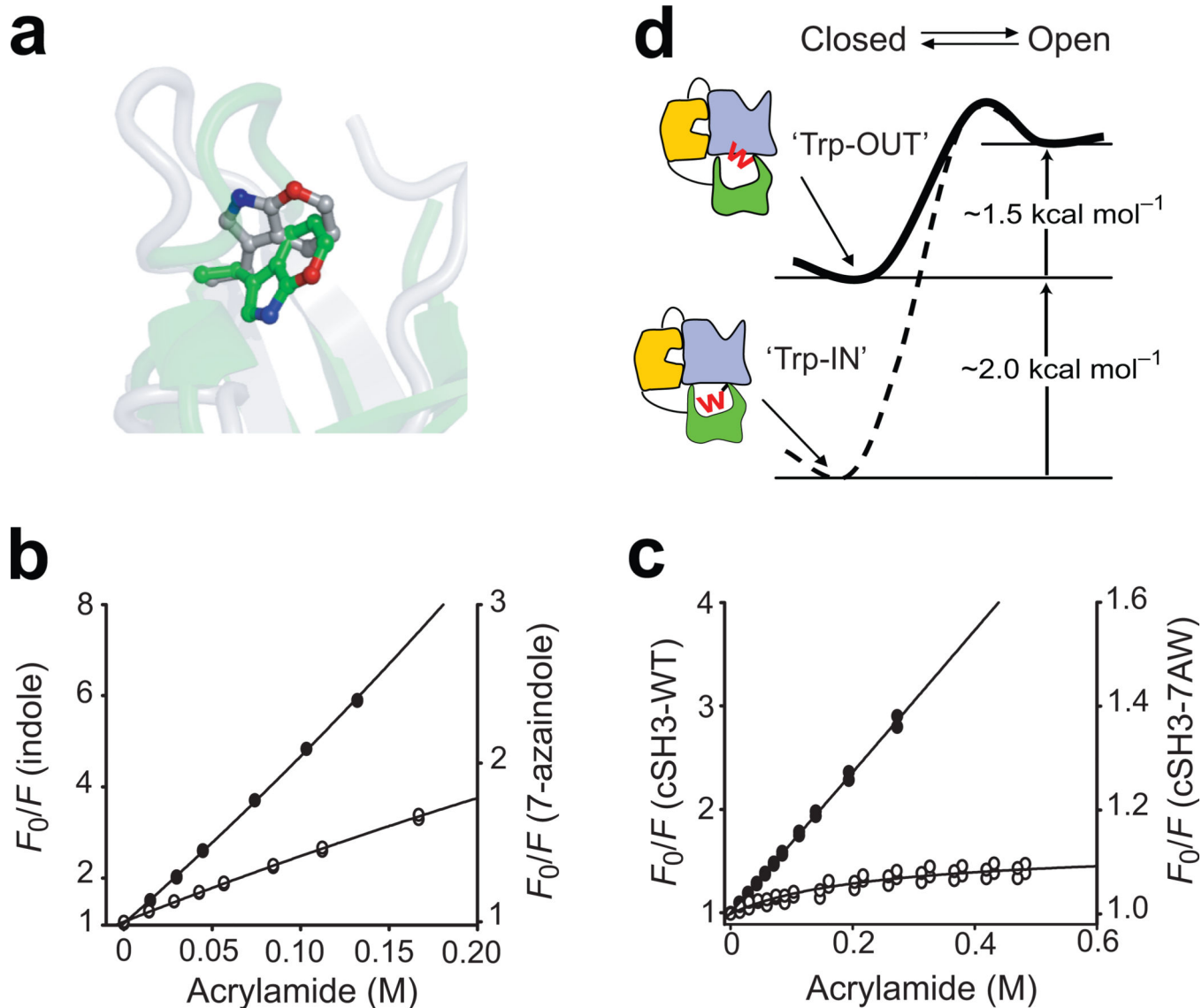


Figure 4. Conformation of Trp275 modulates the stability of the cSH3 domain

(a) Comparison of the position of the indole ring of Trp275 in the structures of isolated cSH3 domain (green, pdb id: 2GGR³²) and full-length Crk-II (silver, pdb id: 2EYZ¹⁹). The 7-position of the Trp residue is highlighted in red. Steady-state fluorescence quenching experiments for (b) free indole (closed circles) and 7-azaindole (open circles) and (c) isolated cSH3-WT (closed circles) and isolated cSH3-7AW (open circles). The solid lines represent the best fit model using the Stern-Volmer relationship (Methods). (d) Schematic representation of the effects of the conformational change of Trp275 on the equilibrium between the open and closed states of Crk-II (W represents Trp275). Reduced stability of the cSH3 upon interdomain interactions reduces the activation barrier between the open and closed states of Crk-II.

Table 1

Binding constants of the C3G-ligand to the nSH3 domain of various states of Crk

Proteins	K_D (nM)	1 Stability (referenced to Crk-I) (kcal mol ⁻¹)
² Crk-I	22.5 ± 1.3	-
Crk-II	285.5 ± 19.0	1.5 ± 0.1

¹Stabilities were calculated from K_O (Methods)²ref. 8

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Table 2

Thermodynamic and fluorescence quenching parameters of the wild type cSH3 domain and the cSH3 domain which contains 7AW at position 275.

Proteins	G° (kcal mol ⁻¹)	m (kcal mol ⁻¹ M ⁻¹)	K_{sv} (M ⁻¹)	λ_{max} (nm)
cSH3-WT	3.3 ± 0.3	0.7 ± 0.1	6.7 ± 0.2	325
Indole	-	-	34.2 ± 1.0	340
cSH3-7AW	2.5 ± 0.4	0.5 ± 0.1	4.7 ± 1.3 ¹ 0.001 ± 0.005 ²	373
7-azaindole	-	-	5.0 ± 0.1	382

¹ Stern -Volmer constant of exposed species deduced from the fit of the data.

² Stern -Volmer constant of buried species deduced from the fit of the data.