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Kinetic Analysis Reveals Differences in the Binding Mechanism of Calmodulin and Calmodulin-like Protein to the IQ Motifs of Myosin-10[¶]

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Abstract

Myo10 is an unconventional myosin with important functions in filopodial motility, cell migration, and cell adhesion. The neck region of Myo10 contains three IO motifs that bind calmodulin (CaM) or the tissue-restricted calmodulin-like protein (CLP) as light chains. However, little is known about the mechanism of light chain binding to the IQ motifs in Myo10. Binding of CaM and CLP to each IQ motif was assessed by non-denaturing gel electrophoresis and by stopped-flow experiments using fluorescence labeled CaM and CLP. Although the binding kinetics are different in each case, there are similarities in the mechanism of binding of CaM and CLP to IQ1 and IQ2: for both IQ motifs Ca²⁺ increased the binding affinity, mainly by increasing the rate of the forward steps. The general kinetic mechanism comprises a 2-step process, which in some cases may involve the binding of a second IQ motif with lower affinity. For IQ3, however, the kinetics of CaM binding is very different from that of CLP. In both cases, binding in the absence of Ca²⁺ is poor, and addition of Ca²⁺ decreases the Kd to below 10 nM. However, while the CaM binding kinetics are complex and best fitted by a multi-step model, binding of CLP is fitted by a relatively simple 2-step model. The results show that, in keeping with growing structural evidence, complexes between CaM or CaM-like myosin light chains and IO motifs are highly diverse and depend on the specific sequence of the particular IQ motif as well as the light chain.

Myosin-10 (Myo10)¹ is an unconventional myosin associated with dynamic actin remodeling and involved in filopodial extension, cell adhesion and mitotic spindle orientation, among its multiple functions (1–6). The basic structure of Myo10 is common to unconventional myosins: the N-terminal head, which binds actin and displays actin-activated ATPase activity is followed by a neck comprised of three light chain-binding IQ motifs and a tail, which in Myo10 includes a putative coiled-coil region, PEST motifs, three PH (pleckstrin homology) domains, a MyTH4 (myosin tail homology 4) domain that binds microtubules, and a FERM (band 4.1/ezrin/radixin/moesin) domain. The presence of MyTH4 and FERM domains relates this myosin with myosins of classes VII and XV involved in hereditary deafness and blindness syndromes (7, 8).

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¹Abbreviations: CaM, calmodulin; CLP, calmodulin-like protein; EGTA, glycol-bis(2-aminoethylether)-*N*, *N*, *N*, *N*-tetraacetic acid; FERM, band 4.1/ezrin/radixin/moesin; HPLC, high pressure liquid chromatography; Myo10, myosin-10; MyTH4, myosin tail homology 4; PH, pleckstrin homology; TA, 2-chloro-(amino-Lys75)-[6-[4-(*N*, *N*-diethylamino)phenyl]-1,3,5-triazin-4-yl]; TEA, triethanolamine.

The regulation of Myo10 by its light chains is poorly understood. As in most other unconventional myosins, calmodulin (CaM) is thought to be the principal light chain of Myo10 (9–11). In addition, however, the epithelial-specific calmodulin-like protein (CLP) also binds to Myo10 as a specific light chain (12), increasing Myo10 expression and stimulating the Myo10-dependent growth of filopodia (13). This raises questions as to how CaM and CLP bind to and compete for the individual IQ sites on Myo10. Because CLP differs from CaM in its Ca^{2+} binding characteristics (14), the effect of Ca^{2+} on the binding and occupancy of each IQ domain by CaM and CLP also remains unexplored.

There is virtually no information on the kinetic mechanism(s) by which CaM and CLP bind to the IQ motifs in the neck of Myo10. Indeed, little is known about the kinetics of binding of CaM (or CaM-like light chain) to any unconventional myosin. Moreover, how Ca²⁺ affects the mechanism of binding of either light chain to the IQ motifs in the neck of Myo10 is unknown. The purpose of this work was to start filling this void. We performed equilibrium and fast-kinetic experiments to elucidate the mechanism of binding of both CaM and CLP to each of the IQ motifs in Myo10. Our results show that while CaM and CLP bind with moderate affinity to the isolated IQ2 domain in the absence of Ca²⁺, both light chains display dramatically increased affinity for each of the three IQ domains in the presence of Ca²⁺. The studies further indicate different binding mechanisms for CLP and CaM to IQ3, suggesting structural differences between the CaM-IQ3 and CLP-IQ3 complexes.

EXPERIMENTAL PROCEDURES

Origin and Preparation of Reagents

CaM was purchased from Calbiochem (LaJolla, CA). CLP was expressed in *E. coli* and purified as described (15, 16). 2-chloro-(amino-Lys75)-[6-[4-(N, N-diethylamino)phenyl]-1,3,5-triazin-4-yl]-CaM (TA-CaM) was synthesized and purified as described (17). Purity was confirmed as reported earlier (18). TA-CLP was synthesized the same way except that bacterially expressed and purified CLP was used as protein. TA-CaM and TA-CLP were kind gifts from Dr. Katalin Török (St. Geoge Hospital Medical School, London, UK) and Dr. Richard Thorogate (London Centre for Nanotechnology, University College London, UK).

Synthetic Peptides

Peptides corresponding to IQ motifs 1–3 were synthesized by the Mayo Peptide Core Facility and purified to homogeneity by HPLC. A second batch of IQ3 peptide was synthesized and purified by Anaspec, Inc. (San Jose, CA). The sequences of the peptides are shown in Figure 1.

Protein-peptide Complex Formation and Non-denaturing Electrophoresis

Aliquots of purified CaM or CLP (10 μ g; 0.6 nmoles) were mixed with equimolar amounts of IQ1, IQ2, or IQ3 peptide in a final volume of 20 μ l sample buffer (25 mM Tris-HCl, pH 6.8, 10% glycerol, 0.1 mM CaCl₂) and incubated at room temperature for 15 min. For competition experiments, equimolar amounts of CLP and CaM were added to the samples containing the pre-formed CaM/IQ and CLP/IQ complexes, respectively, and incubated for an additional 15 min. Non-denaturing electrophoresis was performed in 10% polyacrylamide gels in 25 mM Tris–glycine (pH 8.3), 100 μ M CaCl₂. After running, the gels were stained with Coommassie Brilliant Blue.

Equilibrium Fluorescence

Equilibrium binding of IQ peptides to TA-CaM and TA-CLP was measured at 25°C in a Varian Cary Eclipse spectrofluorometer. Measurements were made in media containing 120

mM KCl, 30 mM Tes-Triethanolamine (Tes-TEA, pH 7.2), 5 mM MgCl₂, 5 mM NaN₃, and 10 mM EGTA (absence of Ca^{2+}) or 0.2 mM EGTA and enough $CaCl_2$ to obtain 100 μ M free Ca^{2+} (presence of Ca^{2+}). For experiments in the presence of Ca^{2+} , $CaCl_2$ was included in both syringes. The concentration of either TA-CaM or TA-CLP was 34 nM. The excitation wavelength was 370 nm and the emission wavelength was 420 nm.

Stopped Flow Rapid Mixing Experiments

Stop-flow measurements of changes in fluorescence of TA-CaM or TA-CLP were performed in an Applied Photophysics SX.18MV reaction analyzer at 25 °C. The excitation wavelength was 365 nm, and the emitted light was detected using a 390 nm cut-off filter. The final mix contained 17 nM TA-CaM or TA-CLP and the concentration of IQ peptides indicated in the legend of the figures, in buffer containing 30 mM Tes-TEA (pH 7.2), 120 mM KCl, 5 mM MgCl₂, 0.2 mM EGTA, and enough CaCl₂ to obtain 100 μ M free Ca²⁺ (or 10 mM EGTA for measurements in the absence of Ca²⁺).

Analysis of Results and Modeling

Preliminary analysis of the fluorescence traces was performed by fitting mono- or multiexponential functions to the data. This was done to obtain an approximate idea of whether a single-step or multi-step kinetic model might be appropriate. Fitting to specific kinetic models was then performed by global non-linear regression analysis using the Dynafit software (www.biokin.com). Global analysis refers to the fact that all progress curves are fitted at once rather than individually. This allows greater discriminatory power between models and enhanced accuracy in determining the kinetic parameters. The software allows direct fitting of predetermined kinetic models to experimental data. The differential equations derived from the kinetic models are solved numerically rather than analytically. In this way, several models of different complexity can be easily tested. When the choice of the best model was not obvious, kinetic models were chosen on the basis of Akaike information criteria (19). These are based on information theory and statistical methods, and take into account the number of parameters as well as the differences between the model and data ² (weighted sum of least square differences). Thus, the tighter fitting for models with a larger number of parameters is balanced against the requirement to use the minimal number of parameters for the fit, with a penalty imposed for the total number of parameters. We have used this procedure before with great success to analyze kinetic data of the binding of peptides corresponding to the calmodulin-regulated domain of the calcium pump to calmodulin (18, 20).

RESULTS

CLP and CaM Bind to Each IQ Motif of Myo10, and CLP Effectively Competes with CaM for IQ Binding

Figure 1 shows the sequence of the three IQ motifs in Myo10. Although the main conserved sequence features of an IQ motif are recognizable in each of these motifs, there are substantial variations and substitutions compared to the consensus sequence IQxxxRGxxxR. For example in IQ1 the canonical Q is replaced by R, and the first canonical R is replaced by L. Likewise, in IQ2 the G is replaced by an A, and in IQ3, the I is replaced by a bulkier F. In view of the different characteristics of these motifs, we studied the binding of the two known light chains of Myo10, CaM and CLP, to the isolated IQ motifs in more detail.

To qualitatively verify the binding of the IQ peptides to each of the light chains, we first mixed IQ peptides and CaM or CLP in stoichiometric amounts in a Ca²⁺- containing buffer, and observed the complexes formed by non-denaturing electrophoresis. It should be noted that under these conditions, differences in the mobility of the bands on the gel mainly reflect

differences in charge and shape, rather than in size, of the protein-IQ peptide complexes. The results are shown in Figure 2. In panel A, each sample contained an equal amount of CaM. Lane 1 shows CaM alone, Lane 2 contained CaM and CLP mixed in equal amounts, demonstrating that CaM and CLP are easily distinguishable in these native gels. Lane 3 shows an equimolar mixture of IQ1 peptide and CaM. The double band indicates the presence of both free CaM and the CaM-IQ1 peptide complex (labeled a in Figure 2). Adding an equimolar amount of CLP to the CaM-IQ1 sample results in the formation of the CLP-IO1 complex (broad band labeled b in lane 4) and a decrease in the band corresponding to the IQ1-CaM complex. Similarly, lane 5 shows that mixing CaM and IQ2 leads to the virtually quantitative formation of the IQ2-CaM complex (labeled c). When CLP is added, the band corresponding to the IQ2-CaM complex is no longer visible and a new band of lesser mobility is formed along with the reappearance of the band for free CaM (lane 6). The new band (labeled d) corresponds to the CLP-IQ2 complex. Lastly, when IQ3 is added to CaM (lane 7) the presence of a band of lesser mobility, together with the disappearance of the band corresponding to free CaM, indicates the formation of the CaM-IQ3 complex (labeled e). Addition of CLP (lane 8) promotes the formation of a new band of even lesser mobility, corresponding to the IQ3-CLP complex (labeled f), together with the reappearance of the band corresponding to free CaM. These results show that while CaM forms a complex with each of the three IO peptides, CLP effectively competes with CaM for IO binding, especially for binding to IQ2 and IQ3.

Figure 2B shows the reciprocal experiment where all samples contained an equal amount of CLP. Lane 1 shows the band corresponding to CLP alone and lane 2 shows the bands corresponding to CLP and CaM. In lane 3, the complex (b) between CLP and IQ1 is seen as slowly migrating, diffuse band. This lane also shows a band corresponding to remaining free CLP, indicating that the CLP-IQ1 complex is not very stable under the conditions of the gel electrophoresis. Addition of CaM (lane 4) does not lead to a significant decrease in the amount of IQ1-CLP complex. Thus, the amount of IQ1-CaM complex formed appears to be small in the presence of CLP. Lane 5 shows the complex (d) between IQ2 and CLP. Addition of CaM (lane 6) results in the formation of some CaM-IQ2 complex (c), but there is little free CLP, indicating that CaM does not very effectively compete with CLP bound to IQ2. Similar results were obtained for the CLP-IQ3 complex (f), as shown in lanes 7 and 8 of panel B. Taken together, the results show that, at least in the presence of Ca²⁺, complexes between the IQ peptides and CaM or CLP are formed and are stable enough to be visualized by non-denaturing gel electrophoresis. Furthermore, at least in the case of IQ2 and IQ3, CLP can effectively compete with CaM for the IQ motifs in Myo10 whereas CaM appears to be less effective in competing with CLP bound to these IQ peptides.

Quantitative Analysis of Binding of CLP and CaM to IQ 1, 2, and 3 of Myo10

To obtain quantitative measurements of CaM and CLP binding to the individual IQ motifs in Myo10, we performed binding experiments using fluorescence labeled TA-CaM and TA-CLP. The advantage of labeling CaM or CLP with the TA fluorescent group is that the fluorescence is very sensitive to the environment, creating the opportunity to detect different steps during the binding process. This approach has previously been used with success to study the kinetic mechanism of the binding of CaM to the plasma membrane Ca^{2+} pump (18, 21). Figure 3 shows the results of titrations of TA-CaM and TA-CLP with the IQ peptides both in the presence and absence of Ca^{2+} . In the absence of Ca^{2+} , binding of IQ1 to TA-CaM and TA-CLP increases the fluorescence of the labeled proteins about 6-fold in the case of TA-CaM and 3-fold in the case of TA-CLP (Figure 3A). The affinity of the interaction in the absence of Ca^{2+} is low (see Table 1), both for TA-CaM and TA-CLP. In the presence of 100 μ M free Ca^{2+} the increase in TA-CaM and TA-CLP fluorescence upon IQ1 binding is smaller (about 2-fold both for TA-CaM and TA-CLP; Figure 3B). The shape

of the curve is similar for IQ1 binding to TA-CaM and TA-CLP. It shows a high affinity component and a smaller, low-affinity increase in fluorescence. In Table 1 we report the K_d values of the high affinity component. The data clearly show that Ca^{2+} promotes a large increase in the affinity of binding of IQ1 to both TA-CaM and TA-CLP.

The binding of IQ2 to TA-CaM and TA-CLP in the absence of Ca^{2+} results in an increase in the fluorescence of both proteins, followed by a small decrease in fluorescence at higher IQ2 concentrations (Figure 3C). The low affinity components (at high peptide concentrations) in the titration curves for IQ1 and IQ2, although small, are important when interpreting the results of rapid kinetic experiments (see below). Ca^{2+} produces an increase in the affinity of both CaM and CLP for IQ2. As observed for IQ1, the change in fluorescence upon binding of IQ2 is much smaller in the presence of 100 μ M Ca^{2+} (Figure 3D). Table 1 shows that the affinity of IQ2 for TA-CaM and TA-CLP is comparable, although CaM appears to bind IQ2 more strongly than CLP in the presence of Ca^{2+} .

Figure 3E and 3F shows the effect of IQ3 binding to TA-CaM and TA-CLP in the absence and presence of Ca²⁺, respectively. IQ3 binds with very low affinity to TA-CaM and TA-CLP in the absence of Ca²⁺ (Figure 3E). In the presence of Ca²⁺, however, IQ3 binds to both proteins with very high affinity (Figure 3F, and see Table 1). A comparison of the fluorescence curves in Figure 3F shows that while binding of IQ3 to TA-CaM produces an increase in fluorescence, binding to TA-CLP produces a small but significant *decrease* in fluorescence, suggesting a difference in the mechanism of binding of IQ3 to these proteins. This is different from the results obtained with IQ1 and IQ2, where binding of these peptides to both TA-CaM and TA-CLP produces similar changes in fluorescence.

Kinetic Analysis of TA-CaM and TA-CLP Binding to IQ1, IQ2 and IQ3

To study the mechanism of binding of CaM and CLP to the IQ motifs in Myo10, we performed stopped-flow experiments mixing TA-CaM or TA-CLP with peptides IQ1, IQ2 and IQ3, in the absence and in the presence of Ca²⁺. To interpret these experiments we considered for each reaction several kinetic models. The models were fitted to the experimental data by means of the Dynafit software (www.biokin.com). This approach has the advantage that the fitting is directly done by numerical integration based on the chemical equations involved rather than on exponential approximations. Figures 5 to 9 show the results of association and dissociation experiments of TA-CaM and TA-CLP with the different IQ peptides in the absence and in the presence of Ca²⁺. The kinetic models used to fit the results are presented in Figure 4. Table 2 lists the model chosen and the values of the relevant constants obtained in each case. These will be discussed later to provide a comparative interpretation.

(i) Binding to IQ1—Figure 5A shows an experiment in which 17 nM TA-CaM were mixed with increasing concentrations of IQ1 (from 1 to 8 μ M) in the absence of Ca²⁺, and the fluorescence was recorded over time. This shows that binding of IQ1 to TA-CaM increases the fluorescence, and that the increase in fluorescence is complete in about 0.5 s. Panel B shows the decrease in fluorescence when IQ1 dissociates from TA-CaM. Dissociation proceeds in a biphasic fashion with a fast phase (k₁ = 2.9 s⁻¹) and a slower one (k₂ = 0.37 s⁻¹).

Figure 5C shows the corresponding experiment where 17 nM TA-CLP were mixed with increasing concentrations of IQ1 (from 1 to 8 μ M) in the absence of Ca²⁺. As observed with TA-CaM, the binding of IQ1 to TA-CLP increases the fluorescence, and the increase in fluorescence is complete in about 2 s. Figure 5D shows the decrease in fluorescence when IQ1 dissociates from TA-CLP. Dissociation proceeds in a biphasic fashion with a faster (k₁ = 0.66 s⁻¹) and a slower phase (k₂ = 0.19 s⁻¹). For both TA-CaM and TA-CLP, model 1

(see Figure 4 and Table 2) provides an excellent fit to the data. The model indicates that TA-CaM (or TA-CLP) initially binds a single molecule of IQ1, and then is able to bind a second molecule, with lower affinity. For comparison, we also show in Figure 5A and 5C the fits (stippled red lines) obtained when a simpler one-step binding model was assumed for the IQ1 interaction with TA-CaM (or TA-CLP). This clearly shows that the simple model is inadequate to fit the data, especially at the higher peptide concentrations. Taken together, the results indicate that the kinetic mechanism of IQ1 binding to CaM and CLP in the absence of Ca²⁺ is essentially the same.

Figure 6 shows the kinetic analysis of the interaction of TA-CaM and TA-CLP with IQ1 in the presence of Ca^{2+} . The progress curves in Figure 6A were recorded upon mixing of 17 nM TA-CaM with different concentrations of IQ1 (from 0.1 to 0.8 μ M) in the presence of 100 μ M Ca^{2+} . The reaction was faster than in the absence of Ca^{2+} . In preliminary fittings, 2 exponential functions were needed to fit the experimental data, suggesting a multi-step reaction. The time course of the dissociation of IQ1 from TA-CaM in the presence of 100 μ M Ca^{2+} is shown in Figure 6B. Again the dissociation was biphasic with a faster ($k_1 = 2.5 s^{-1}$) and a slower ($k_2 = 0.8 s^{-1}$) component. The model required to fit the experimental results was more complex than that for fitting the data in the absence of Ca^{2+} and corresponds to model 4 in Figure 4. After binding a first IQ1 peptide, CaM may either undergo a subsequent conformational change (to yield the TA-CaM-IQ1* state), or it may bind a second IQ1 in no preferred order of addition.

The association and dissociation data for the interaction of TA-CLP with IQ1 in the presence of 100 μ M Ca²⁺ are shown in Figure 6, panels C and D. Both the association and the dissociation were slower than in the case of TA-CaM. However, the model used to fit the data is similar to that for binding of IQ1 to TA-CaM in the presence of Ca²⁺ (model 4), except that the binding of the second molecule of IQ1 does not occur until the first one is bound. The model is illustrated in Figure 4 as model 3.

(ii) Binding to IQ2—The results of the progress curves when 17 nM TA-CaM or TA-CLP were mixed with different concentrations of IQ2 in the absence of Ca^{2+} are shown in Figure 7A and 7C, respectively. For both proteins, the association appears to have 2 phases, with a faster increase in fluorescence followed by a slower phase. This is more apparent at higher concentrations of IQ2. Panels B and D of Figure 7 shows the kinetics of dissociation of IQ2 from TA-CaM and TA-CLP, respectively, in the same (Ca^{2+} -free) conditions. Again, the dissociation is biphasic with a fast component ($L_1 = 0.34 \, \text{s}^{-1}$ for TA-CaM and 0.13 s⁻¹ for TA-CLP) and a slower one ($L_2 = 0.036 \, \text{s}^{-1}$ and $0.023 \, \text{s}^{-1}$, respectively, for TA-CAM and TA-CLP). The model that gave the best fit of these data corresponds to model 4 in Figure 4. According to this model TA-CaM and TA-CLP bind one molecule of IQ2, and then can either undergo a conformational stabilization step, or bind a second molecule of IQ2. Although the same model fitted the data for TA-CLP, the overall reaction for both association and dissociation was slower for TA-CLP than for TA-CaM (compare Figure 7, panels A and B with panels C and D).

Figure 8A and 8C shows the corresponding association data when 17 nM TA-CaM or TA-CLP were mixed with different concentrations of IQ2 in the presence of $100~\mu M$ Ca²⁺. The results resemble those obtained in the absence of Ca²⁺, except that the reaction is faster and we used lower concentrations of IQ2 peptide because of the higher affinity of TA-CaM and TA-CLP for IQ2 in Ca²⁺ conditions (see Table 1 and Figure 3). The dissociation reaction (Figure 8B and 8D) is also biphasic, although significantly slower than the same reaction in the absence of Ca²⁺ (compare Figure 8B with 7B, and Figure 8D with 7D). The kinetic model used for the binding of IQ2 to TA-CaM and TA-CLP in the absence of Ca²⁺ was equally able to fit the data in the presence of Ca²⁺ (model 4, Figure 4).

(iii) Binding to IQ3—We did not analyze the kinetics of binding of TA-CaM and TA-CLP to IQ3 in the absence of Ca²⁺ because the interaction is so weak that the data were not reliable (see also Figure 3 and Table 1). Therefore, the experiments on the kinetics of binding of IQ3 to TA-CaM and TA-CLP were performed only in the presence of 100 µM Ca²⁺. In Figure 9A we show the changes in fluorescence when 17 nM TA-CaM was mixed with increasing concentrations of IQ3 from 0.05 to 2 µM. The fluorescence changes produced by IQ3 binding are very complex. At lower IQ3 concentrations, binding produces a small increase in fluorescence that follows approximately mono-exponential kinetics. However, at higher IQ3 concentrations the progress curves show an increase, a decrease, and, at longer times, a new increase. At the highest IQ3 concentrations, the initial increase is so fast that it happens mostly during the dead time of the instrument (~ 1.3 ms). The dissociation curve is complex as well (Figure 9B). It shows an extremely rapid increase in fluorescence, a pronounced decrease, and a slow increase. These patterns indicate a multistep reaction pathway for both the association and the dissociation of IQ3 to and from TA-CaM. The model that best fit the experimental data corresponds to model 5 in Figure 4. It includes the possibility of the addition of a second IQ3 molecule, as well as two stabilization steps. Several models of lesser or similar complexity were tried but were unable to fit the data.

In contrast to the complex behavior of TA-CaM, binding of 17 nM TA-CLP to IQ3 leads to a small decrease in fluorescence (Figure 9C) along a curve that could be fitted by the sum of 2 exponential functions. We assayed several concentrations of IO3 from 0.1 to 1.6 µM, but for the sake of clarity only the extreme low and high concentrations are shown. Figure 9D shows the changes in fluorescence upon dissociation of IQ3 from TA-CLP. This shows a relatively slow ($k = 0.015 \text{ s}^{-1}$) increase in fluorescence. The small total changes in fluorescence, and consequently the relatively larger noise in the data, prevented in this case a sophisticated model analysis, but model 2 in Figure 4 fit the results well. In this model, IQ3 binds to TA-CLP, and initial binding is followed by a second step resulting in an altered conformational state of the complex. Another model, in which IQ3 can bind to two alternative conformations of TA-CLP, was also able to fit the experimental data, but the Akaike information criteria were slightly better for model 2. Regardless of the actual model that is used to fit the data, it is clear that TA-CLP binds to IQ3 in a different manner than TA-CaM. This is a unique characteristic of the IO3 motif, since in the case of IO1 and IO2 the models that fitted the binding progress curves were identical, or very similar, for TA-CaM and TA-CLP.

DISCUSSION

The purpose of this work was to establish kinetic parameters for the binding of the two known light chains of Myo10, CaM and CLP, to each of the IQ motifs in this myosin. To independently assess binding, we performed non-denaturing gels that allowed us to identify the complexes of IQ peptides with either CaM or CLP. The complexes were only detected in the presence of Ca²⁺, suggesting that both CaM and CLP bind the individual IQ motifs with higher affinity in the Ca²⁺-bound holo-form. These qualitative assays also showed that, in an equimolar ratio, CLP was able to partially displace CaM from its complex with the IQ motifs, and appeared to be most effective at competing with CaM for IQ2 and IQ3. Thus, it is likely that CLP, which is expressed only in specific tissues, can displace CaM from Myo10 in vivo, especially because CaM levels are thought to be limiting in cells (22).

To obtain quantitative information on the individual binding affinities of CaM and CLP for each IQ motif of Myo10, equilibrium titrations were performed with TA-CaM and TA-CLP. One observation that was initially unexpected (but is in agreement with the native gel electrophoresis data on complex formation) is that in general, Ca²⁺ increased the affinity of

both TA-CaM and TA-CLP for each of the IQ motifs. This was more evident for IQ1 and IQ3 than for IQ2 (see Table 1). In the presence of 100 μM Ca $^{2+}$, the affinity of TA-CaM and TA-CLP for IQ1 was between 30 and 100 times higher than in the absence of Ca $^{2+}$, yielding Kd values of $\sim\!50-150$ nM. A high concentration of Ca $^{2+}$ (100 μM) was chosen to saturate all 4 Ca $^{2+}$ binding sites in both CaM and CLP; thus the (+ Ca $^{2+}$) values reported in Table 1 reflect the affinity of the fully Ca $^{2+}$ -loaded light chain for the specific IQ motif. The difference in the affinity in the absence and presence of Ca $^{2+}$ is even more pronounced for the binding of TA-CaM and TA-CLP to IQ3. In the presence of Ca $^{2+}$, the affinity is very high (Kd values between 5 and 10 nM), whereas binding in the absence of Ca $^{2+}$ was virtually undetectable. The effect of Ca $^{2+}$ on the affinity of TA-CaM and TA-CLP for IQ2 is more modest, but still significant.

IQ motifs were initially thought to be preferred binding sites for (Ca²⁺-free) apo-CaM (9, 10). However, more recently it has been shown that this is not always the case, and depends strongly on the sequence context of the IQ motifs and whether they are arranged in tandem (23, 24). Martin and Bayley (25) showed that Ca²⁺ increases the affinity for CaM of a tandem made of IQ motifs 3 and 4 of MyoV. The data were interpreted to indicate that in the presence of Ca²⁺, CaM forms a bridge between contiguous IQ motifs with high affinity (26). A recent study on the IQ domain of Myo1c (a myosin-I family member) also found that CaM bound tightly to the tandem IQ motifs both in the presence and absence of Ca²⁺ (27). In addition, it has been pointed out that if IQ motifs have hydrophobic residues at positions 1-5-8-14 (as is typically found in Ca²⁺-dependent CaM binding motifs) they may show Ca²⁺ dependency as well (23). All three IQ motifs of Myo10 do have hydrophobic residues in these positions (Figure 1), except for a Gln residue in position 8 of IQ3. It may be of interest in future studies to determine if and how changes in these hydrophobic residues affect the Ca²⁺ sensitivity of CaM and CLP binding to the Myo10 IQ motifs.

An interesting observation obtained from the stopped-flow experiments is that Ca^{2+} has very little effect on the rate of dissociation of either TA-CaM or TA-CLP from IQ1 and IQ2. By contrast, the rate constants derived from the kinetic analysis indicate that the forward reactions are faster in the presence of Ca^{2+} . In particular, Ca^{2+} increases the rate constant k_1 for the association of CaM with IQ1 and of CLP with IQ2, as well as the rate (k_5) of binding a second IQ peptide by CaM and CLP (see Table 2). These results suggest that CaM and CLP could be prone to forming "bridges" between different (adjacent) tandem IQs in Myo10 in the presence of Ca^{2+} . The results are in keeping with the findings of Martin and Bayley (26) who reported the formation of very high affinity 1:1 complexes of CaM with tandem IQ motifs from MyoV in the presence of Ca^{2+} . It should be noted, however, that our experiments showed that binding of a second IQ peptide to CaM or CLP required the peptide to be in large excess, suggesting the low affinity of such a possible second binding event.

IQ3 shows very different kinetic characteristics when compared to IQ1 and IQ2: (i) for both TA-CaM and TA-CLP, there is little or no binding in the absence of Ca²⁺; (ii) binding of TA-CaM to IQ3 shows an overall increase in fluorescence, while binding of TA-CLP shows a decrease in fluorescence. By contrast, both TA-CaM and TA-CLP show a similar pattern of overall fluorescence change upon binding to IQ1 or IQ2; (iii) the progress curve of TA-CaM binding to IQ3 shows a very complex multi-step pattern, whereas the mechanism of TA-CLP binding to IQ3 seems much simpler (compare model 5 with model 2 in Figure 4). We emphasize that the binding schemes in Table 2 must be interpreted with caution, as these are merely models yielding optimal fits of the experimental data. The exact chemical nature of the intermediate steps is undetermined, thus definite conclusions as to the structure and conformation of different TA-CaM or TA-CLP complexes with a single or two IQ peptides must await high-resolution structural determination. However, regardless of the uncertainties

associated with the precise structures of the complexes, the binding of TA-CLP to IQ3 follows a very different kinetic path compared to TA-CaM.

Multiple effects contribute to the fluorescence changes induced by binding of the IQ peptides to TA-CaM or TA-CLP. These include conformational changes of TA-CaM/TA-CLP upon interaction with the peptides, as well as possible changes in the fluorescence yield of the complexes caused by induced secondary structure stabilization of the IQ peptides upon binding. Therefore, the structural identity of the kinetic intermediates postulated by the models (such as TA-C-IQ* and TA-C-IQ^) remains undetermined. However, the distinct kinetic steps likely correspond to major structural transitions in TA-CaM/TA-CLP and/or the peptide(s), which affect the environment of the fluorescent TA moiety.

When discussing the kinetic differences in light chain binding to these IQ motifs, it is important to keep their sequence differences in mind. None of these IQ motifs comply with all the characteristics of the canonical IQ motif IQxxxRGxxxR. IQ1 contains an R residue instead of a Q in the initial IQ, and the RG is changed to LG. Although an R at position 2 in this motif can participate in a network of H-bonds like Q does, it is a charged and more bulky residue, potentially rationalizing why IQ1 has lower affinity for the light chain than IQ2 and IQ3. IQ2 is close to the canonical sequence, the only modification being RA instead of RG. The effect of substitution of the canonical G by A in this position has recently been studied in a reference IQ motif, and was found to decrease the affinity for CaM in the absence of Ca²⁺ but increase the affinity for Ca²⁺-bound CaM (28). However, of the three Myo10 IQs, IQ2 still has the highest affinity for both Ca²⁺-free CLP and CaM (Table 1), suggesting that other residues within the IQ motif are equally important for binding to CaM and CLP in the absence of Ca²⁺. Finally in IQ3, the aliphatic I in position 1 is replaced by F (a bulky aromatic residue). Whether this replacement is sufficient to confer the observed different properties to IQ3 in regard to the effect of Ca²⁺ on light chain binding remains to be addressed in future work. However, it is interesting to note that an F>A mutation in IQ3 drastically reduced CLP, but not CaM, binding (29).

A pending question concerns the changes that will occur in terms of light chain binding to the IQ1-3 neck region of Myo10 upon a rise in Ca²⁺. Given the complexity of the issue, one can only speculate at this point. However, in light of our results some general points may be legitimate: (a) Binding of both CaM and CLP are expected to increase with a rise of Ca²⁺; (b) no significant change in the rate of dissociation of either CaM or CLP may occur, but rebinding could be impaired by one light chain forming a tight bridge between neighboring IQ motifs, especially between IQ1 and IQ2; (c) regarding which light chain would prevail, in the "in vitro" conditions CaM and CLP show comparable affinities. However, "in vivo" the conditions might differ, considering that CaM has many more targets than CLP. Therefore, its free concentration in the cell is greatly reduced (22). Taking this into account, it is probable that in the tissues where it is expressed CLP is the preferred light chain on most, if not all IQ motifs of Myo10.

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hMyo10-IQ1: FEVSHAAMVIRAHVLGFLARKQYR

763 786
hMyo10-IQ2: RKVLYCVVIIQKNYRAFLLRRRFL

786 810
hMyo10-IQ3: LHLKKAAIVFQKQLRGQIARRVYRQ

Consensus IQ: IQxxxRGxxxR

FIGURE 1.

Sequence of the three IQ motifs in Myo10. The underlined residues are in the positions of the consensus IQ motif IQxxxRGxxxR, and the residues indicated in bold are the hydrophobic residues in positions 1-5-8-14. The first and last residues of each IQ peptide are numbered according to their position in full-length human Myo10.

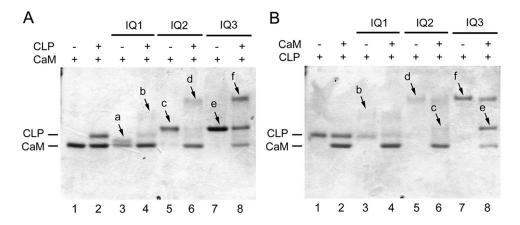


FIGURE 2.

Identification of CaM-IQ and CLP-IQ complexes by non-denaturing gel electrophoresis. Complexes were formed and the gels were run as described in the Methods. In panel A, all lanes contained 10 μg CaM. Lanes 2, 4, 6 and 8 also contained 10 μg of CLP. Lanes 3 and 4 contained an amount of IQ1 equimolar with the amount of CaM, lanes 5 and 6 contained equimolar amounts of IQ2, and lanes 7 and 8 contained equimolar amounts of IQ3. In panel B, all lanes contained 10 μg of CLP, lanes 2, 4, 6 and 8 also contained 10 μg CaM, lanes 3 and 4 contained IQ1, lanes 5 and 6 contained IQ2 and lanes 7 and 8 contained IQ3, all in equimolar amounts.

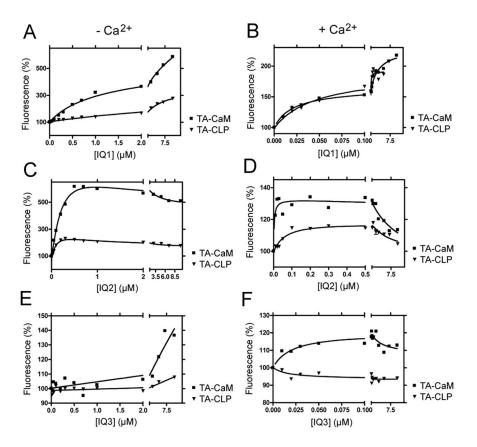


FIGURE 3. Titration of TA-CaM and TA-CLP with the IQ motifs of Myo10, in the absence (A, C and E) and in the presence (B, D and F) of Ca^{2+} . Titrations were done by monitoring the fluorescence of TA-CaM or TA-CLP at 25°C in the media described in the Experimental Procedures. When the measurements were done in the absence of Ca^{2+} the buffer contained 10 mM EGTA. When the measurements were done in the presence of Ca^{2+} , the buffer contained 0.2 mM EGTA and enough $CaCl_2$ to obtain 100 μ M free Ca^{2+} . In each case, fluorescence is represented as a percentage of the fluorescence of the sample in the absence of IQ peptide.

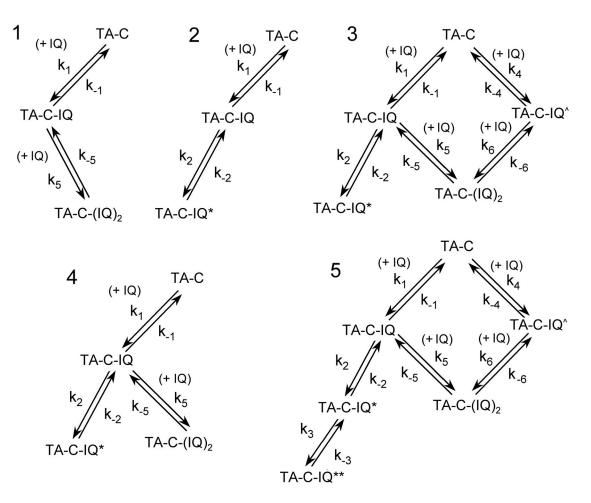


FIGURE 4.

Schematic representation of the kinetic models used for fitting the data obtained in the stopped flow experiments described in Figures 5–9. TA-C stands for either TA-CaM or TA-CLP. TA-C-IQ*, TA-C-IQ**, and TA-C-IQ^ denote different conformations of the complex of TA-CaM or TA-CLP with IQ peptide.

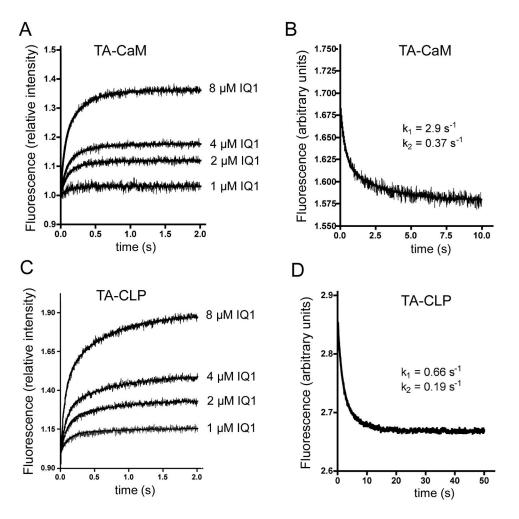


FIGURE 5. Kinetics of binding of TA-CaM and TA-CLP to IQ1 in the absence of Ca^{2+} . (A) and (C): Time course of fluorescence changes upon mixing 17 nM TA-CaM (A) or 17 nM TA-CLP (C) with the indicated concentrations of IQ1. The experimental data were best fitted (continuous blue line) with the model and kinetic constants shown in Table 2. For comparison, the fit using the simpler one-step model is also shown as stippled red line, clearly indicating a poorer fit especially at higher IQ1 concentrations. (B) and (D): Progress curves of the dissociation of TA-CaM (B) or TA-CLP (D) from IQ1. 34 nM TA-CaM or 34 nM TA-CLP were pre-mixed with 2 μ M IQ1, and dissociation was initiated by adding 10 μ M CaM (B) or 10 μ M CLP (D). The progress curves were fitted with a sum of 2 exponentials, yielding the constants shown in the figure. All media contained 10 mM EGTA. In all cases, the indicated concentrations are after final mixing.

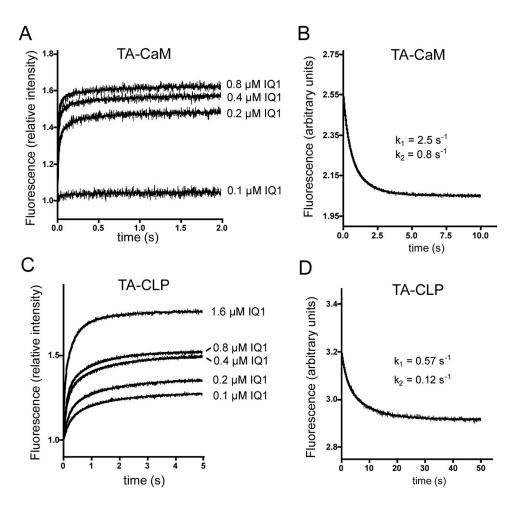


FIGURE 6. Kinetics of binding of TA-CaM and TA-CLP to IQ1 in the presence of 100 μM free Ca^{2+} . (A) and (C): Time course of fluorescence changes upon mixing 17 nM TA-CaM (A) or 17 nM TA-CLP (C) with the indicated concentrations of IQ1. The experimental data were fitted with the models and kinetic constants shown in Table 2. (B) and (D): Progress curves of the dissociation of TA-CaM (B) or TA-CLP (D) from IQ1. 34 nM TA-CaM or 34 nM TA-CLP were pre-mixed with 0.2 μM IQ1, and dissociation was started by adding 1 μM CaM (B) or 1 μM CLP (D). The progress curves were fitted with a sum of 2 exponentials, with the constants shown in the figure. All media contained 0.2 mM EGTA and enough $CaCl_2$ to obtain 100 μM Ca^{2+} . In all cases, the indicated concentrations are after final mixing.

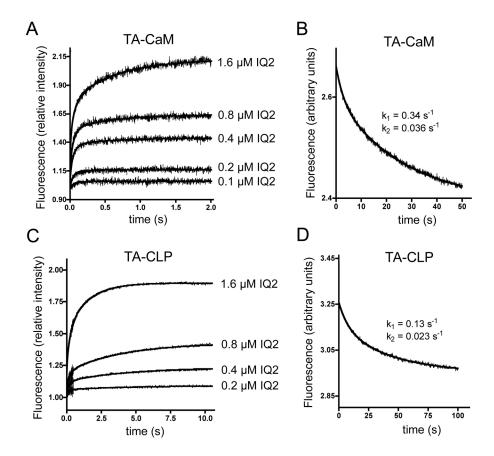


FIGURE 7. Kinetics of binding of TA-CaM and TA-CLP to IQ2 in the absence of Ca^{2+} . (A) and (C): Time course of fluorescence changes upon mixing 17 nM TA-CaM (A) or 17 nM TA-CLP (C) with the indicated concentrations of IQ2. The experimental data were fitted (smooth black lines) with the models and kinetic constants shown in Table 2. (B) and (D): Progress curves of the dissociation of TA-CaM (B) or TA-CLP (D) from IQ2. 34 nM TA-CaM or 34 nM TA-CLP were pre-mixed with 0.2 μ M IQ2, and dissociation was initiated by adding 1 μ M CaM (B) or 1 μ M CLP (D). The progress curves were fitted with a sum of 2 exponentials, with the rate constants shown in the figure. All media contained 10 mM EGTA. In all cases, the indicated concentrations are after final mixing.

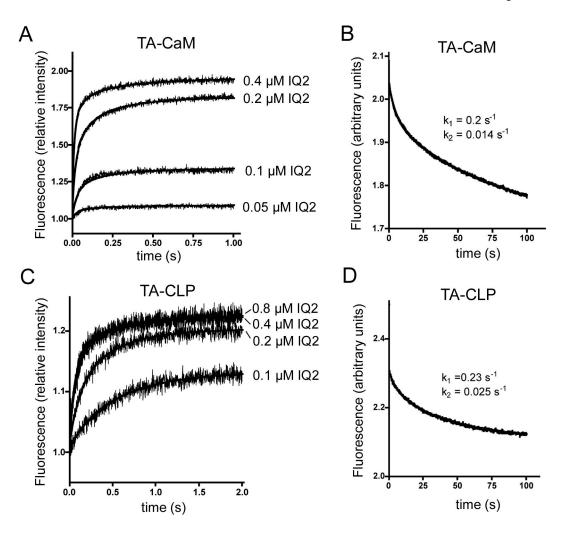


FIGURE 8. Kinetics of binding of TA-CaM and TA-CLP to IQ2 in the presence of 100 μ M free Ca²⁺. (A) and (C): Time course of fluorescence changes upon mixing 17 nM TA-CaM (A) or 17 nM TA-CLP (C) with the indicated concentrations of IQ2. The experimental data were fitted (smooth black lines) with the models and kinetic constants shown in Table 2. (B) and (D): Progress curves of the dissociation of TA-CaM (B) or TA-CLP (D) from IQ2. 34 nM TA-CaM or 34 nM TA-CLP were pre-mixed with 0.2 μ M IQ2, and dissociation was started by adding 1 μ M CaM (B) or 1 μ M CLP (D). The progress curves were fitted with a sum of 2 exponentials, with the constants shown in the figure. All media contained 0.2 mM EGTA and enough CaCl2 to obtain 100 μ M Ca²⁺. In all cases, the indicated concentrations are after final mixing.

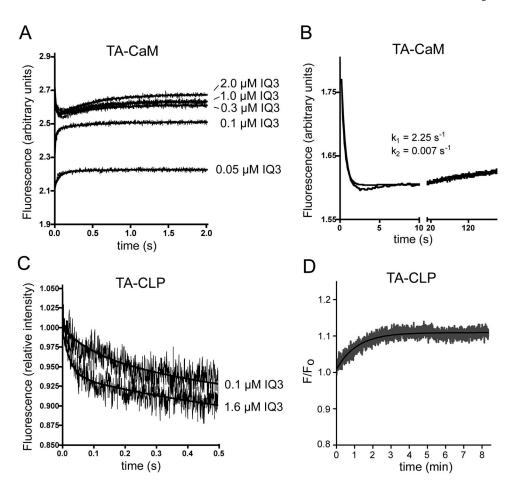


FIGURE 9. Kinetics of binding of TA-CaM and TA-CLP to IQ3 in the presence of 100 μM free Ca²⁺. (A) and (C): Time course of fluorescence changes upon mixing 17 nM TA-CaM (A) or 17 nM TA-CLP (C) with the indicated concentrations of IQ3. In panel C, the data obtained with 0.2, 0.4 and 0.8 μM IQ3 were omitted from the figure for the sake of clarity, but the accuracy of the fit was comparable to that shown in the figure. The experimental data were fitted (smooth black line) with the models and kinetic constants shown in Table 2. (B) and (D): Time course of the dissociation of TA-CaM (B) or TA-CLP (D) from IQ3. 34 nM TA-CaM or 34 nM TA-CLP were pre-mixed with 0.2 μM IQ3, and dissociation was started by adding 1 μM CaM (B) or 1 μM CLP (D). In panel B, the progress curve could be fitted with a sum of 2 exponentials, with the constants indicated in the figure. In contrast, dissociation of IQ3 from TA-CLP is slower ($k = 0.015 \text{ s}^{-1}$) and results in an increase of fluorescence

(panel D). All media contained 0.2 mM EGTA and enough CaCl2 to obtain 100 µM Ca²⁺. In

all cases, the indicated concentrations are after final mixing.

 $\label{eq:Table 1} \textbf{Table 1}$ Parameters of TA-CaM and TA-CLP binding to Myo10 IQ peptides a

Protein	Condition ^b	K _d IQ1 (nM)	K _d IQ2 (nM)	K _d IQ3 (nM)
CaM	- Ca ²⁺	5078 ± 1888	708 ± 48	> 50000
Calvi	+ Ca ²⁺	152 ± 78	6.9 ± 6.1	9.4 ± 3.1
CLP	- Ca ²⁺	4938 ± 338	134 ± 27	(NM) ^C
CLI	+ Ca ²⁺	57 ± 35	46 ± 7	7.2 ± 4.4

 $^{^{}a}$ Equilibrium titrations were performed as described in the Experimental Procedures. Affinities for the high-affinity binding sites were determined from the binding isotherms and are reported as K_{d} values.

 $^{^{}b}$ The buffer for the Ca $^{2+}$ -free (- Ca $^{2+}$) condition contained 10 mM EGTA, whereas 100 μ M Ca $^{2+}$ was present in the Ca $^{2+}$ containing (+ Ca $^{2+}$) condition

 $^{^{}c}$ Not measurable.

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Table 2

Kinetic parameters for binding of TA-CaM and TA-CLP to IQ motifs in ${
m Myo}10^a$

			TA-CaM					TA-CLP		
Peptide)I	IQ1)ı	1Q2	1Q3	I	IQ1	1Q2	2	1Q3
Condition	- Ca ²⁺	+ Ca ²⁺	– Ca ²⁺	+ Ca ²⁺	+ Ca ²⁺	– Ca ²⁺	+ Ca ²⁺	- Ca ²⁺	+ Ca ²⁺	+ Ca ²⁺
Model	1	4	4	4	2	1	3	4	4	2
$k_1 (\mu M^{-1} s^{-1})$	0.662 ± 0.044	173.4±1.1	78.74±33.35	11.81±1.536	450.2 ± 65.18	1.03 ± 0.048	4.24±1.65	4.9719±0.2134	52.14±17.51	103.43±5.857
$k_{-1} (s^{-1})$	8.423±0.346	7.128±0.125	12.503±0.7889	27.187±0.8348	118.04±7.249	8.03±0.2	7.50±1.31	5.6748±0.2905	29.03±18.07	2.54±0.6693
$k_2 (s^{-1})$		0.3153±0.1174	7.5534±0.6942	0.4685 ± 0.3594	2.072±2.847		4.861±3.026	2.9254±0.2536	5.137±3.794	2.166±0.8612
$k_{-2} (s^{-1})$		3.8825±0.2155	1.471±0.1936	2.9365±0.2101	4.215±4.782		0.808 ± 0.2353	0.35875±0.02875	1.157 ± 0.05252	0.015^{b}
$k_3 (s^{-1})$					1.066±1.868					
$k_{-3} (s^{-1})$					3.283±3.466					
$k_4 (\mu M^{-1} s^{-1})$					406.98±34.3		12.915±1.221			
$k_{-4} (s^{-1})$					8.5281±0.3125		2.0773±0.1243			
$k_5 (\mu M^{-1} s^{-1})$	0.0941 ± 0.0148	90.21 ± 9.54	3.196±1.696	31.651±0.8853	<0.0001	0.231±0.017	0.4284 ± 0.6298	0.2129 ± 0.05354	27.93±29.4	
$k_{-5} (s^{-1})$	3.036 ± 0.079	68.251±5.44	1.175 ± 0.1493	14.027±0.6736	165.9±78.83	1.091 ± 0.022	3.526±2.9	0.30602±0.02266	5.989±0.7487	
$k_6 (\mu M^{-1} s^{-1})$					6.755±1.608		0.3229±0.2473			
$k_{-6} (s^{-1})$					<0.0005		<0.0001			
rta-c	1	1	1	1	1	1	1	1	1	1
ľTA-C-IQ	1.41	2.33	1.466	4.336	1.538	2.14	1.939	2.19	1.11	0.935
ľTA-C-1Q2	1.88	1.61	3.89	1.449	0.273	2.40	4.694	0.98	1.19	
rta-c-10^					1.223		1.290			
ľta-c-iq*		4.69	1.041	6.849	1.341		1.960	9.04	1.31	0.878
ľTA-C-IQ**					2.768					

a kx refers to the forward rate constants for each of the steps in the models considered, k-x refers to the reverse rate constants, rTA-x refers to the relative fluorescence of the species considered, taking the fluorescence of TA-C as 1.

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 $b_{\rm Estimated}$ value as data noise prevented detailed model analysis.