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A Novel Dimerization Interface of Cyclic Nucleotide Binding Domain, which is Disrupted in Presence of cAMP: Implications for CNG Channels Gating

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ABSTRACT Cyclic nucleotide binding domain (CNBD) is a ubiquitous domain of effector proteins involved in signalling cascades of prokaryota and eukaryota. CNBD activation by cyclic nucleotide monophosphate (cNMP) is studied well in case of several proteins. However, this knowledge is hardly applicable to cNMP-modulated cation channels. Despite the availability of CNBD crystal structures of bacterial cyclic nucleotide-gated (CNG) and mammalian hyperpolarization-activated cyclic nucleotide-modulated (HCN) channels in presence and absence of the cNMP, the full understanding of CNBD conformational changes during activation is lacking. Here, we describe a novel CNBD dimerization interface found in crystal structures of bacterial CNG channel MlotiK1 and mammalian cAMP-activated guanine nucleotide-exchange factor Epac2. Molecular dynamics simulations show that the found interface is stable on the studied timescale of 100 ns, in contrast to the dimerization interface, reported previously. Comparisons with cN-bound structures of CNBD show that the dimerization is incompatible with cAMP binding. Thus, the cAMP-dependent monomerization of CNBD may be an alternative mechanism of the cAMP sensing. Based on these findings, we propose a model of the bacterial CNG channel modulation by cAMP.

Keywords: cyclic-nucleotide binding domain, CNG channel, Epac2, channel gating

Abbreviations: CNBD, Cyclic nucleotide binding domain; cNMP, cyclic nucleotide monophosphate; cAMP, cyclic adenosine monophosphate; CNG channel, cyclic nucleotide-gated channel; HCN channel, hyperpolarization-activated cyclic nucleotide-modulated channel; MD, molecular dynamics; RMSD, root mean square deviation; PCA, principal components analysis.

Introduction

The cyclic nucleotide binding domain (CNBD) is a part of many cAMP-regulated proteins [1]. Its function has been well studied for catabolite activator protein (CAP), protein kinase A (PKA) and guanine nucleotide-exchange factor (GEF) Epac2. CAP is organized as a dimer, with its interface being on the CNBD helix C [2]. Binding of cAMP induces a coil-to-helix transition in CNBD-distal end of CNBD helix C, and consequent rearrangement of DNA-binding domain, which results in the increase in DNA affinity [3]. In PKA, binding of cAMP to the regulatory subunit results in dissociation of subunits and activation of the catalytic subunit [4]. As for Epac2, it functions as a monomer, but there are two CNBD domains in it (CNBD-A and CNBD-B). In the inactive form, CNBD-A and CNBD-B are bound to each other and occlude the binding site of the small G protein Rap. Upon introduction of cAMP, CNBD-B moves away from the Rap binding site and Epac2 becomes active [5].

Besides aforementioned proteins, there exist a number of tetrameric cation channels whose conductivity is modulated by cyclic nucleotides. These channels lie at heart of many cellular processes and were extensively studied by various biophysical techniques [6]. Important insights into the channels' function were gained from structural studies [6]. Crystallographic structures revealed not only the tertiary structure of channel CNBDs, but also their multimeric states. CNBD from bacterial cyclic nucleotide gated (CNG) channel MlotiK1 crystallizes as a dimer with non-crystallographic symmetry axis [7, 8] and the structure of the monomers resembles that of CNBDs from other proteins. Because the CNBD was co-purified with a bound cAMP, which was difficult to remove by dialysis, the crystallographic structure of the cAMP-free CNBD was determined for mutants with hampered cyclic nucleotide binding [7, 8]. Though some cAMP-dependent structural rearrangements were revealed, the findings did not result in a definitive hypothesis on how the channel activation could proceed [1, 7]. NMR studies provided complementary information on the CNBD structure in solution, both in presence and absence of cAMP, and found no evidence of dimerization [9, 10]. Finally, the 16 Å resolution electron microscopy structure of the whole-length cAMP-bound MlotiK1 channel revealed that in that state the CNBDs are completely dissociated [11]. As for more complex eukaryotic cyclic nucleotide-

gated (CNG) channels and hyperpolarization-activated cyclic nucleotide-modulated (HCN) channels, the crystallographic structures of the cytoplasmic parts revealed similar CNBD structure and tetrameric overall assembly, which proceeds via the interaction between the C-linkers [12–14]. However, no large-scale structural rearrangements, which could result in channel modulation, were detected between the cNMP-free and cNMP-bound structures. Thus, despite the extensive studies, the full understanding of the cyclic nucleotide modulation of both prokaryotic and eukaryotic cation channels has not been reached.

Here, we describe a novel CNBD dimerization interface, present in two different crystallographic structures, the first one is of MlotiK1 CNBD domain R307W mutant with hampered cAMP binding, PDB ID 3CO2 [8], and the other one is of Epac2, PDB IDs 1O7F [15] and 2BYV [16]. This dimerization is incompatible with cAMP binding, and thus the CNBD monomerization upon binding of cAMP may be a mechanism of the cyclic nucleotide sensing. We have studied the stability of the MlotiK1 CNBD homodimer and Epac2 CNBD heterodimer by means of molecular dynamics. No structural rearrangements were observed during the simulations. The dimers were stable on the studied timescale. As a reference, we have also simulated the other MlotiK1 CNBD dimerization interface, described in the previous studies, which is observed in both the cAMP-bound and the cAMP-free structures [7, 8]. This interface fluctuates excessively and partially dissociates during the simulations. On the basis of our findings, we discuss the possibility of CNBD dimerization in the physiological context and propose the mechanism of the bacterial CNG channel gating, based on cAMP-disrupted dimerization of its CNBDs.

Materials and Methods

For molecular dynamics simulations the models were immersed in a water box with 8 Å padding. Sodium and chloride ions were added at a total concentration of 0.2 M in such amounts that the total charge of a system would be equal to zero. Before the production run, the systems were equilibrated in three steps. First, the energy of the system was minimized using standard algorithms [17, 18]. After that, the solvent was equilibrated for 1 ps, with the protein atoms being restrained. Finally, the whole system was released and equilibrated for additional 1 ps.

Atomic coordinates for MlotiK1 CNBD dimers were taken from the structure of R307W mutant (PDB ID 3CO2 [8]) and the model was completed to contain residues 221-350. The residue W307 was mutated back to arginine in order to represent the wild-type protein using the psfgen utility [17], and the structure was equilibrated by the standard procedure. After the equilibration, R307 occupied the conformation similar to that observed in the experimental structures. Atomic coordinates for Epac2 CNBD dimer were taken from the PDB ID 1O7F [15]. Residues 13-167 were taken for CNBD-A, and 305-444 for CNBD-D. All the residues were assumed to be in their standard protonation states, based on by-residue pKa values determined with PROPKA server [19]. In total, three molecular systems were prepared: the Epac2 CNBD heterodimer, the MlotiK1 CNBD homodimer with the novel interface, and the MlotiK1 CNBD homodimer with a previously described interface. Each system was simulated for 100 ns at 310 K with CHARMM27 parameters [20] with integration timestep of 2 fs. The Epac2 and MlotiK1 dimers with a novel interface were simulated using NAMD2 version 2.7 [17], and the MlotiK1 dimer with a previously reported interface was simulated using GROMACS version 4.5.3 [18]. Bonds between the hydrogen atoms and the heavy atoms were kept rigid using the SHAKE algorithm. In namd2 simulations, the temperature was maintained with the Langevin thermostat with the damping coefficient of 5 ps^{-1} , and the pressure of 1 bar was maintained with the Langevin piston barostat with the following parameters: period of 100 fs, decay of 50 fs. In GROMACS simulations, the temperature was maintained with the Berendsen thermostat modified to reproduce the correct sampling of the temperature [21], and the pressure was maintained using the Parinello-Rahman scheme [22].

The structures and the trajectories were analyzed using VMD [23]. For determination of the root mean square deviations of the atomic coordinates, all trajectory frames were aligned using the backbone atoms of both protomers. Principal components analysis (PCA) [24] of the mass-weighted covariance matrix of the backbone atoms' coordinates was conducted using the tools `g_covar` and `g_anaeig` of the GROMACS suite [18].

Results and Discussion

Analysis of the novel CNBD dimerization interface

The interface is present in crystallographic structures of two different proteins, bacterial CNG channel MlotiK1 R307W mutant (PDB ID 3CO2 [8]) and human guanine nucleotide exchange factor Epac2 (PDB IDs 1O7F [15] and 2BYV [16]). The protomers are located slightly asymmetrically. There are two main interactions present (Fig. 1). First, there are extensive β -sheet-like interactions of strand 4 with 6' and strand 6 with 4' (prime denotes the other protomer in the dimer). It appears that the β -roll of one protomer continues into the β -roll of the second protomer. Second, there are numerous hydrophobic contacts between the internal surfaces of the β -strands 4 and 5. In case of MlotiK1 CNBD, the dimer is further stabilized by the contact between the N-terminal helix of one protomer and β -strands 4 and 5 of the other (Fig. 1, (a)). In case of Epac2 CNBD, the dimer is additionally stabilized by the contact between the region C-terminal to CNBD-A and β -roll of CNBD-B (Fig. 1, (b)).

The described interfaces possess contact surface areas of 800 Å² and 1100 Å² in MlotiK1 and Epac2 CNBD dimers, correspondingly, as calculated by PISA server [25]. However, in both dimers the protomers are not independent *in vivo*. The only physiological form of the MlotiK1 channel is tetrameric, with the distance between the N-terminal tails of CNBD domains in the range 10-20 Å, as judged from the crystallographic structure of the transmembrane part [26]. In Epac2 CNBD-A and CNBD-B are simply a part of the same polypeptide chain. It means that in both cases the local concentrations of the interacting protomers are extremely high, of the order of 10% v/v and higher. Thus, the usual expectations about the contact surface area of physiological interfaces (as e.g. in [27]) are not applicable here, as the local concentration of interacting partners in case of Epac2 and MlotiK1 is highly elevated, compared to the usual concentrations of the interacting proteins in cytosol, and the proteins would dimerize even when the association constants are lower than usual. The more thorough consideration of the effects arising with domain linking (tethering) may be found in [28].

To sum up, from the general considerations it appears very plausible that the described dimerization takes place in physiological settings.

Incompatibility of the dimerization with cAMP binding

The described dimers possess a very notable feature, which may be directly relevant for their function. Both dimers are observed in the absence of the CNBD's ligand cAMP. Comparison with the cAMP-bound structures (PDB ID 1VP6 for MlotiK1 CNBD [7] and PDB ID 3CF6 for Epac2 CNBD [29]) shows that the presence of cAMP is incompatible with the dimerization. First, the cAMP, bound to the first protomer, occludes the place of binding of the second protomer, and second, the movement of the C-helix, induced by cAMP binding, would result in the steric clash of this helix with the second protomer (Fig. 1). Thus, the dimerization via the analyzed interface is impossible in the presence of cAMP. We propose that the dissociation of CNBD dimers upon rise of the cAMP concentration may underlie the function of CNBD in the MlotiK1 and Epac2.

Molecular dynamics simulations

To analyze stability of the observed dimers on the atomic level, we conducted the molecular dynamics (MD) simulations. We simulated the MlotiK1 and Epac2 CNBD dimers for 100 ns. Both dimers reveal a high degree of stability. The average RMSD values for the backbone atoms are 2.3 Å for MlotiK1 dimer and 2.1 Å for Epac2 dimer, and RMSD values for the backbone atoms at the contact site are 1.8 Å for MlotiK1 dimer and 1.8 Å for Epac2 dimer. Time-dependence of the RMSD for both systems is shown in Fig. 2.

Analysis of the dependence of the RMSD on residue number shows that on average all the residues of the simulated dimers behave similarly (Fig. 3). Outliers are the unstructured regions between the α -helices $\alpha A'$ and αA of CNBD-A and α -helices $\alpha A''$ and $\alpha A'$ of CNBD-B of Epac2. However, these regions are distal to the dimerization interface and their increased mobility should not affect the dimerization. Also, the N- and C-termini of all proteins reveal greater mobility, which is expected for the exposed ends of a polypeptide chain. As for the contact site, its residues in both proteins do not display any increased mobility compared to other residues. Fluctuations of the backbone atoms are presented in Fig. 3.

In MlotiK1 CNBD homodimer, the protomers are positioned slightly asymmetrically. As a consequence, the difference in mobilities of the pin between β -strands 4 and 5 (4-5 pin), and of the β -strands is observed. The 4-5 pin of the protomer B is exposed to the solvent, and reveals a higher mobility as compared

to the buried 4-5 pin of the protomer A (Fig. 3). Also, a small rearrangement of the 4-5 pin of the protomer B is observed for one MlotiK1 protomer. This may be ascribed either to MD artifacts or to the probable errors in the initial crystallographic model due to its relatively low resolution (2.9 Å).

As for the Epac2 CNBD heterodimer, the fluctuations of the contact site residues of both protomers (β -strands 4 and 5) are similar. The 4-5 pin is much longer in CNBD-A, it does not reveal any secondary structure and it does not participate in the dimerization. As a consequence, it fluctuates stronger than the neighboring residues (Fig. 3). β -strands 4 and 5 of CNBD-A are also more mobile than those of CNBD-B, probably as a consequence of the mobile pin.

To sum up, the molecular dynamics show that the CNBD dimers observed in crystallographic structures remain stable on the studied timescale, the residues of the contact sites fluctuate similarly to other residues, and thus the observed interaction is significant.

Molecular dynamics simulations of the previously reported MlotiK1 CNBD dimer

Previously, it was proposed that the MlotiK1 may dimerize via the α -helices A' and A [7, 8] (Fig. 4). However, this notion is challenged by a number of other studies [6, 9–11, 30]. Here, we test this dimerization interface by means of molecular dynamics. As the CNBDs are dissociated in the cAMP-bound state [11], we used the interface from the cAMP-free mutant crystallographic structure as a starting conformation (PDB ID 3CO2, [8]). The dimer reveals excessive fluctuations, which sometimes result in a partial dissociation (Fig. 4 and 5) of the protomers (for example, at the time mark ~61 ns). There, the interaction is reduced to the α -helices A', and the contact area is of the order of 300 Å².

In order to further characterize dynamics of this dimer, we have carried out the principal components analysis [24] of the mass-weighted covariance matrix of the backbone atoms' coordinates. The analysis reveals that the conformational changes are dominated by the first three modes (Fig. 5). There, the protomers move largely independently as they do not change their own conformations, but move relative to each other for as much as 20 Å during the trajectory (Fig. 5). The nature of the contacts between the protomers is changed dramatically (Fig. 5).

The situation is different for the novel type of dimerization. There, only the first mode somewhat corresponds to relative domain motions (Fig. 5). However, its amplitude is lower than the amplitude of any of the first three modes of the previously reported dimer, and the dimerization interface is preserved. This mode is rather a result of the overall flexibility of CNBD.

Thus, we conclude that the previously reported type of dimerization is probably unspecific and simply reflects the hydrophobic nature of the interface. It should also be noted, that in this interface the contact site surface area is lower than in the dimerization interface proposed above, $\sim 600 \text{ \AA}^2$ versus $800 \text{ \AA}^2$ and $1100 \text{ \AA}^2$. The α -helices A', which continue from the transmembrane helices S6, are directed almost opposite to each other (Fig. 4), whereas in the novel interface the α -helices A' cross at much lower angle (90° , Fig. 3). To sum up, it appears that if the CNBD dimerization occurs, it is much more likely to proceed via the newly reported interface, described above, and not via the previously reported one.

Implications for CNG Channels Gating

Bacterial CNG channels consist of three domains: four-helical trans membrane (TM) voltage sensing-like domain, two-helical TM ion channel pore domain, and CNBD. Mammalian CNGs and homologous to them HCNs are more complex and have elongated N- and C-termini and, most important, a C-linker, consisting of several α -helices, between the pore domain and CNBD. Presence of this linker undoubtedly results in qualitative differences. There, the CNBD is no longer connected directly to the S6 helix of the ion pore, and the distance between the CNBD and the membrane is much larger. Moreover, the crystallographic structures of the cytoplasmic parts of HCN2 and HCN4 reveal tetrameric ring-like assemblies, where the interaction is mediated by the C-linker [12, 14, 31]. Finally, the gating process itself is different in mammalian and bacterial CNG channels at least in one aspect: while in the former it is strictly cooperative (Hill coefficients in the range 3-4), in the latter it is not (Hill coefficients of ~ 1.5) [7, 8, 6]. Thus, the differences between the mammalian and bacterial channels are significant, and we will limit ourselves to the discussion of only the bacterial CNG's gating.

The MlotiK1 CNBD has been extensively studied experimentally [7–11, 30, 32]. It is a general conclusion that this domain behaves as a monomer in solution [9,

10, 30]. However, as we have already pointed out, the concentration of the protein used in these experiments is of the order of 0.1% v/v, meanwhile in the physiological context in the functional tetrameric channel there are four CNBDs in a very close proximity, with the effective local concentration of the order of 10-20% v/v. Thus, the proposed dimerization could easily go undetected in the experiments. Finally, we would like to make a remark about the study of Cukkemane et al. [30]. There, three different methods were used to determine the dissociation constant K_D of cAMP (isothermal titration calorimetry, tryptophan fluorescence and 8-NBD-cAMP fluorescence). In these three techniques the authors used different protein concentrations (of the order of $50\text{ }\mu\text{M} \approx 0.1\% \text{ v/v}$, $5\text{ }\mu\text{M}$ and $0.5\text{ }\mu\text{M}$), perhaps unintentionally. The resulting K_D were 107 ± 11 , 80 and $67.8 \pm 8.7\text{ nM}$, correspondingly [30]. The discrepancies between the determined K_D values may result from the methodological differences between the techniques, but the other explanation is also possible: cAMP competes with the CNBD dimerization, and as a consequence, it has a higher apparent K_D at higher CNBD concentrations. Thus, the experimental results do not contradict the proposition of MlotiK1 CNBD dimerization, and probably support it.

The possibility of the cAMP-disrupted dimerization of CNBD allows us to formulate the hypothesis on the gating of MlotiK1. In the absence of cAMP, the CNBDs are in the dimeric form and the channel is closed (Fig. 6). Binding of cAMP prevents the CNBD dimerization. Thus, in presence of cAMP, the CNBD domains are completely separated, in accord with low resolution electron microscopy structure [11], and the channel is open (Fig. 6).

Conclusions

In this study, we have described the novel dimerization interface of cyclic nucleotide-binding domains, which is disrupted in presence of cAMP. Two 100 ns molecular dynamics simulations of the dimers from different proteins demonstrate that the CNBD dimer is stable on this timescale, and that the contact residues preserve their positions. The previously reported interface, on the opposite, is found to be unstable during the 100 ns molecular dynamics simulation. Possibility of cAMP-disrupted dimerization of CNBD allows us to formulate the hypothesis of bacterial CNG channel gating. In the absence of cAMP, the CNBDs are in the dimeric form and the channel is closed. Binding of cAMP prevents the CNBD dimerization and thus promotes the channel opening. This hypothesis fits and explains the available experimental data on channel function. Finally, the novel interface may be involved in some yet unknown interactions in cases where the CNBD function is not determined, as for CNBD-A domain of Epac2.

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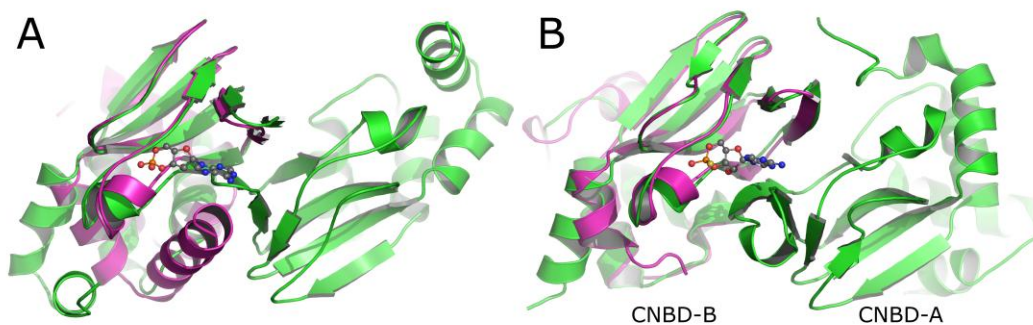


Fig. 1 Dimers of cyclic nucleotide binding domains observed in crystallographic structures. (a) Superposition of the cAMP-bound CNBD monomer (*magenta*) and the CNBD dimer in the absence of cAMP (*green*) for CNG ion channel MlotiK1 (PDB IDs 1VP6 and 3CO2). (b) Superposition of the cAMP-bound CNBD monomer (*magenta*) and the CNBD dimer in the absence of cAMP (*green*) for guanine nucleotide-exchange factor Epac2 (PDB IDs 3CF6 and 1O7F) correspondingly. cAMP is shown in grey, in balls and sticks representation. Note that the presence of cAMP is incompatible with dimer formation.

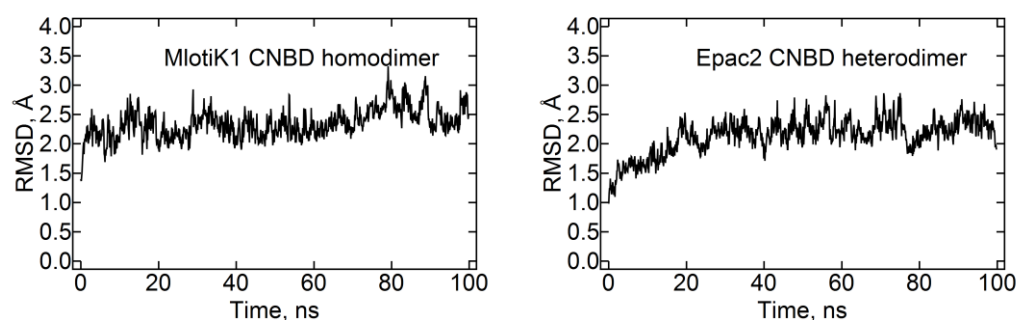


Fig. 2 RMSD of atomic coordinates of the backbone atoms during the trajectory of molecular dynamics simulations of MlotiK1 CNBD homodimer and Epac2 CNBD heterodimer. RMSD is measured relative to the initial (crystallographic) structures.

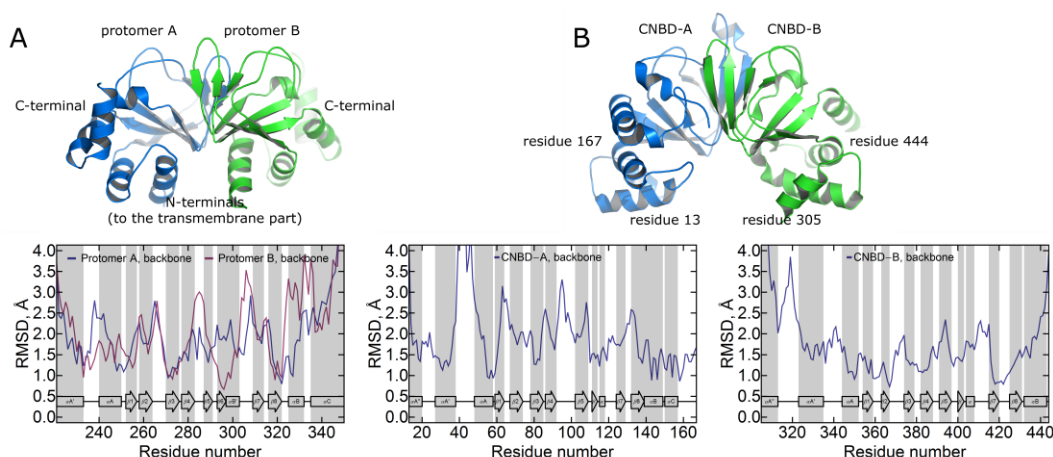


Fig. 3 Backbone fluctuations as a function of the residue number. (a) MlotiK1 CNBD homodimer. (b) Epac2 CNBD heterodimer. Regions with defined secondary structures are highlighted. The secondary structure is labeled as in [7].

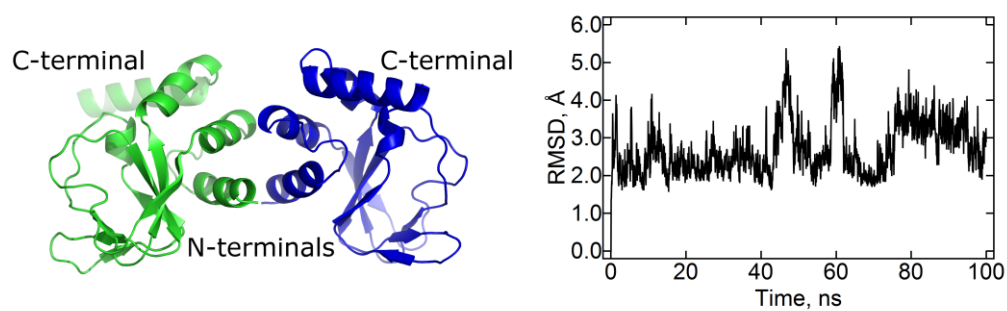


Fig. 4 Structure of the alternative dimerization interface, proposed in the previous studies [7, 8] and its fluctuations during the molecular dynamics simulations. Fluctuations of the atomic coordinates of all the backbone atoms are shown. Two partial dissociation events are observed at ~46 ns and ~61 ns, where the whole-dimer RMSD exceeds 5 Å and the contact site area is reduced to ~320 Å².

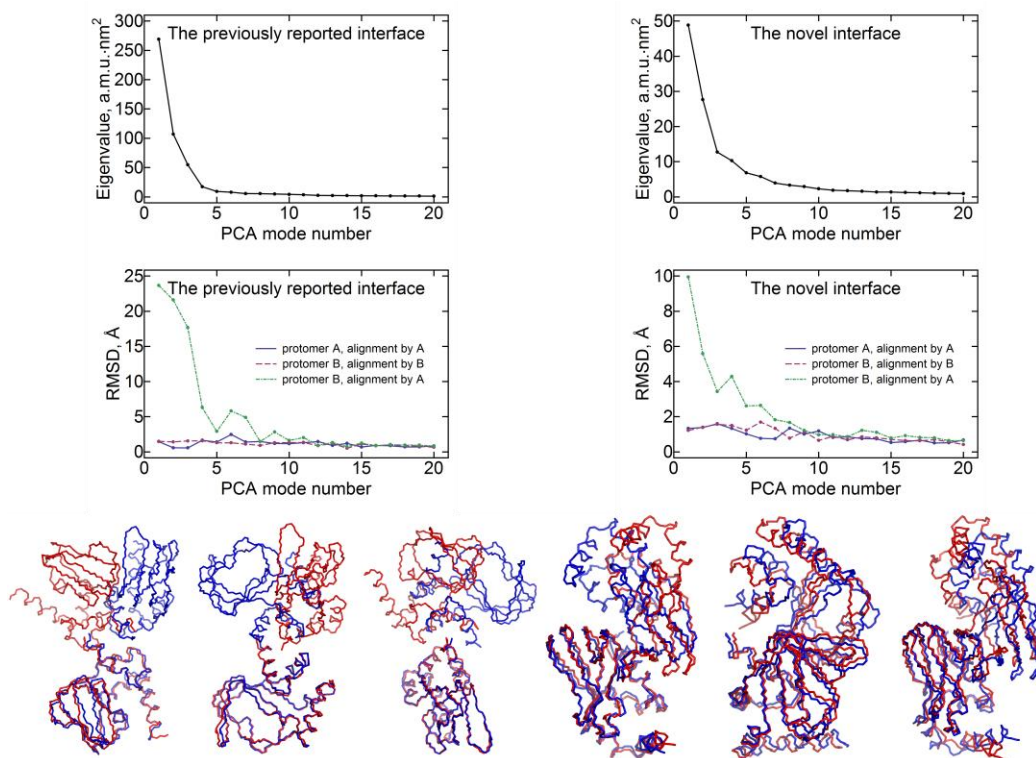


Fig. 5 Principal components analysis of the simulated trajectories of MlotiK1 CNBD dimers that interact via the previously proposed interface or via the novel one. In the top row, the first 20 eigenvalues of the mass-weighted covariation matrices are shown. Note that the scales of the graphs are different. In the middle and the bottom rows, extreme projections of the trajectory on the corresponding eigenvectors are analyzed. In the middle row, there are the root mean square deviations of the backbone atoms of the protomers A and B, with the whole structure being aligned either by the protomer A or by the protomer B. It can be seen that the first three modes of the previously reported dimer trajectory correspond largely to the relative motions of the protomers. In the bottom row, backbone traces of the extreme conformations along the first three eigenvectors are shown for the both simulations. The structures are aligned by one of the protomers. The view is chosen so that the corresponding conformational changes are seen most clearly.

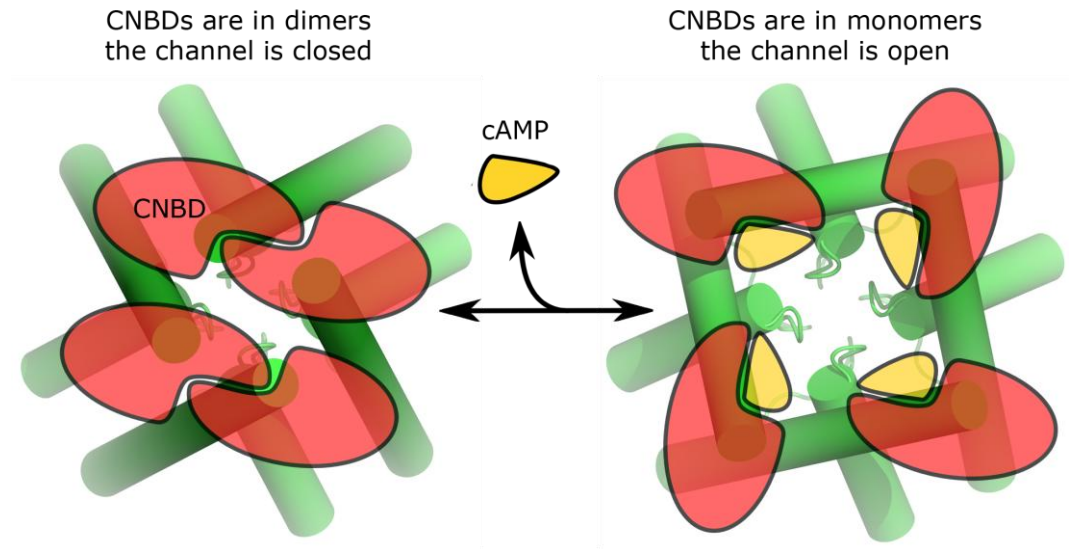


Fig. 6 Model of the bacterial CNG channel activation by cAMP. The view is from the cytoplasm perpendicular to the membrane plane. Only the pore helices S5 and S6 are shown. In absence of cAMP, the CNBDs are in dimers, the channel's S6 helices are close to each other and the pore is occluded. In presence of cAMP, the CNBDs dissociate, and the ion pore opens. PDB IDs 2AHY and 3E86 were used to represent the closed and the open state of the pore correspondingly [33, 34]. Helices of the voltage sensor-like domain are not shown.