Cyclic di-AMP traps proton-coupled K^+ transporters of the KUP family in an inward-occluded conformation

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Abstract

Cyclic di-AMP is the only known essential second messenger in bacteria and archaea, regulating different proteins indispensable for numerous physiological processes. In particular, it controls various potassium and osmolyte transporters involved in osmoregulation. In *Bacillus subtilis*, the K^+/H^+ symporter KimA of the KUP family is inactivated by c-di-AMP. KimA sustains survival at potassium limitation at low external pH by mediating K^+ ions uptake. However, at elevated intracellular K^+ concentrations, further K^+ accumulation would be toxic. In this study, we reveal the molecular basis of how c-di-AMP binding inhibits KimA. We report cryo-EM structures of KimA with bound c-di-AMP in detergent solution and reconstituted in amphipols. By combining structural data with functional assays and molecular dynamics simulations we reveal how c-di-AMP modulates transport. We show that an intracellular loop in the transmembrane domain interacts with c-di-AMP bound to the adjacent cytosolic domain. This reduces the mobility of transmembrane helices at the cytosolic side of the K^+ binding site and therefore traps KimA in an inward-occluded conformation.

33 **Introduction**

34 In bacteria and archaea, potassium ions are essential, playing roles in
35 osmoregulation[1,2], pH homeostasis[3–5], regulation of protein synthesis[6], enzyme
36 activation[7,8] such as the ribosome[9,10], membrane potential adjustment[11] and electrical
37 signaling[12–14]. K^+ homeostasis must be strictly maintained as deviations and
38 fluctuations of potassium levels have lethal consequences for the cell[15,16]. In Gram-
39 positive bacteria like *Bacillus subtilis* the second messenger cyclic di-AMP (c-di-AMP)
40 is essential for the regulation of channels and transporters that maintain potassium
41 homeostasis[17]. At elevated intracellular K^+ concentrations, c-di-AMP is present at
42 increased concentrations and regulates potassium-transporting proteins in two ways:
43 Firstly, it directly binds its target proteins, leading to the inhibition of potassium
44 uptake systems like K^+ channel KtrAB[18,19] and K^+/H^+ symporter KimA[17,20] or the
45 activation of potassium exporters like CpaA[21] and KhtTU[22]; secondly, c-di-AMP
46 suppresses gene expression by binding to the *ydaO* riboswitch[23] that among others
47 controls the transcription of the *kimA* and *ktrAB* genes. The ability of c-di-AMP to
48 modulate both the expression and the activity of the same proteins makes it a key
49 player in potassium homeostasis[24]. However, while the control of gene expression is
50 well understood, it remains unclear how c-di-AMP binding controls the activity of
51 potassium transporters and channels.

52 The c-di-AMP-sensitive, high-affinity K^+/H^+ symporter KimA from *B. subtilis* is
53 particularly required for the uptake of potassium at low external potassium
54 concentrations in acidic environments. KimA exploits the inward-directed proton
55 gradient to accumulate K^+ against its concentration gradient. As a member of the
56 amino acid-polyamine-organocation (APC) superfamily it has the classical LeuT-fold
57 with the first ten transmembrane helices (TMHs) adopting a 5+5 inverted repeat.
58 TMHs 11 and 12 connect the transmembrane domain (TMD) to the cytosolic domain
59 (CD) of KimA, which consists of four alpha helices and a five-stranded beta sheet[25].
60 KimA is a homodimer, stabilised by the swapping of the cytosolic domains with respect
61 to the transmembrane domain. A long loop connects the swapped cytosolic domain
62 to the last helix of the TMD. The dimeric cytosolic domains adopt a fold similar to a
63 phosphopantetheine adenylyltransferase (PPAT) domain and have been suggested to
64 bind c-di-AMP[25,26]. However, no structural information is available for the c-di-AMP

binding site and, consequently, the inhibition mechanism of potassium uptake by c-di-AMP remains elusive. In this study, we report new structural and functional insights into the binding of c-di-AMP to KimA and how it inhibits the transporter.

Results

C-di-AMP binds to KimA in a cooperative manner and inhibits potassium uptake

The uptake of potassium ions through KimA is known to be impaired by c-di-AMP[16,19]. Here, we confirmed the hypothesis and showed that binding of c-di-AMP leads to a significant decrease of potassium uptake velocity under *in vivo* conditions. We co-produced KimA with the inactive adenylate cyclase variant $CdaA_{D171N}$[27] in *Escherichia coli* LB2003 cells, a strain that lacks all endogenous potassium uptake systems[28], and determined potassium uptake into the potassium-depleted cells, obtaining a V_{max} of 269.3 ± 24.0 nmol*mg dw^{-1}*min^{-1} and a K_m of 0.49 ± 0.09 mM. Upon the co-expression of KimA with active, c-di-AMP-synthesising CdaA, the V_{max} was reduced by ~64% to 96.7 ± 10.1 nmol*mg dw^{-1}*min^{-1}, while the K_m did not change (0.66 ± 0.31 mM) (Figure 1a, Table 1).

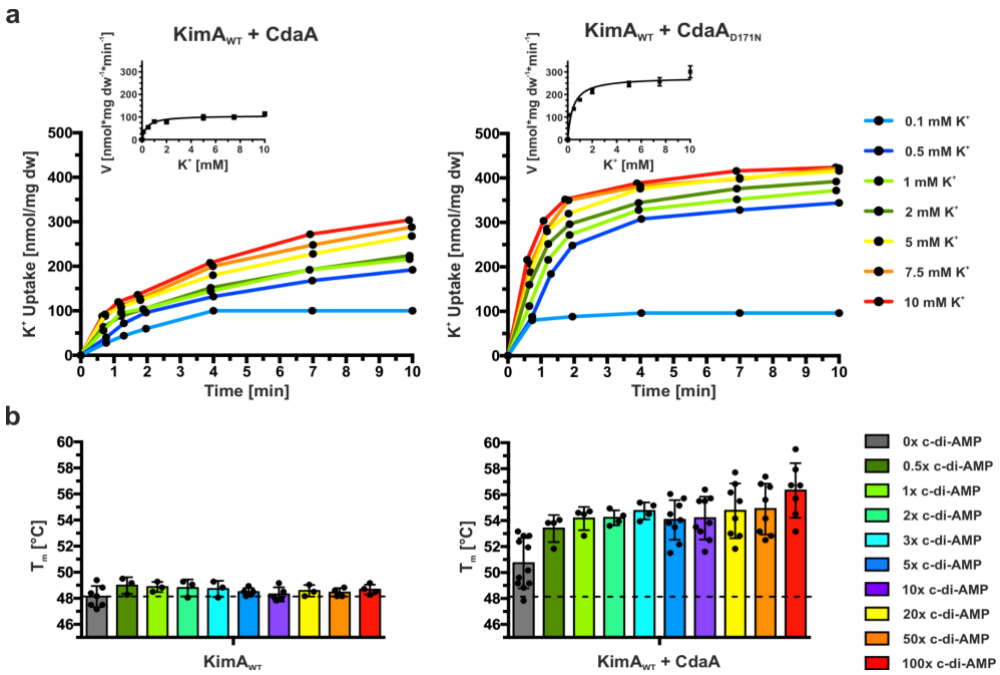


Figure 1: The effect of c-di-AMP binding to KimA *in vivo* and *in vitro* a) Whole-cell K^+ uptake assay in *E. coli* LB2003 cells producing KimA with active (CdaA) or inactive ($CdaA_{D171N}$) diadenylate cyclase. Michaelis-Menten plot shown in graph in graph. Representative experiments shown (n= 3). **b)** Melting temperatures of KimA purified from cells with CdaA present (KimA WT + CdaA) or absent (KimA WT) and incubated with an increasing c-di-AMP concentration given in x-fold molar excess over KimA. Determined with Differential Scanning Fluorometry (DSF). Dashed line indicates T_m of KimA WT w/o c-di-AMP addition. Data points represent the average and error bars the standard deviation of measurements from at least biological triplicate.

90 **Table 1: Kinetic parameters of KimA variants in the presence or absence of c-di-AMP (i.e., active**
91 **(CdaA) or inactive ($CdaA_{D171N}$) diadenylate cyclase).** Determined by fitting the Michaelis-Menten plot
92 with Michaelis-Menten equation. Mean and standard deviation from three independent whole-cell
93 potassium uptake assays are shown.

Variant	V_{max} [nmol/(mg dw*min)]	K_m [mM]
$KimA_{WT}$ + CdaA	96.7 ± 10.1	0.66 ± 0.31
$KimA_{WT}$ + $CdaA_{D171N}$	269.3 ± 24.0	0.49 ± 0.09
$KimA_{R337A}$ + CdaA	111.7 ± 25.5	0.54 ± 0.41
$KimA_{R337A}$ + $CdaA_{D171N}$	198.9 ± 35.8	0.39 ± 0.08
$KimA_{Y118A}$ + CdaA	210.6 ± 29.0	18.85 ± 5.82
$KimA_{Y118A}$ + $CdaA_{D171N}$	202.8 ± 19.4	4.8 ± 0.75
$KimA_{N237A}$ + CdaA	207.9 ± 23.6	7.80 ± 2.23
$KimA_{N237A}$ + $CdaA_{D171N}$	161.9 ± 41.8	7.93 ± 4.21
$KimA_{A481WS582W}$ + CdaA	194.2 ± 49.2	0.35 ± 0.15
$KimA_{A481WS582W}$ + $CdaA_{D171N}$	182.0 ± 44.1	0.33 ± 0.08

94

95 In our previous structural study[25], when c-di-AMP was added to KimA purified in
96 styrene maleic-acid lipid particles (SMALPs) before EM grid preparation, binding of c-
97 di-AMP to KimA was not observed. In agreement with this we show here that the
98 melting temperature of detergent-purified KimA did not significantly change upon the
99 titration of increasing concentrations of c-di-AMP in differential scanning fluorometry
100 (DSF) measurements (Figure 1b), suggesting that c-di-AMP did not bind in these
101 conditions. To overcome this limitation, we co-produced KimA with the diadenylate
102 cyclase CdaA in *E. coli* LB2003 cells, as this combination showed KimA inhibition *in*
103 *vivo*, and then purified KimA from this condition. Interestingly, this sample already had
104 a slightly increased melting temperature from 48.1 ± 0.8 °C to 50.7 ± 1.9 °C, prior to
105 the addition of c-di-AMP. The addition of a three-fold molar excess of c-di-AMP led to
106 a further 6.6 °C increase of the melting temperature to 54.7 ± 0.7 °C (Figure 1b).
107 Hence, during purification, a fraction of c-di-AMP appeared to remain bound to KimA,
108 which enabled the binding of further c-di-AMP under *ex vivo* conditions. This suggests
109 a cooperative behaviour of c-di-AMP binding to KimA at least under *in vitro* conditions.
110

111 **Structural characterisation of the binding of c-di-AMP to KimA**

112 To determine the binding site of c-di-AMP and the effect of c-di-AMP binding on the
113 overall structure of KimA, cryo-EM specimens of purified KimA that had been co-
114 expressed with *cdaA* were prepared in both DDM and amphipols. KimA in DDM was
115 additionally incubated with a ten-fold molar excess of c-di-AMP after purification.

116 332k and 296k particles were processed to reconstruct cryo-EM maps at 3.3 Å and 3.8
117 Å resolution, respectively. The overall structure of KimA obtained from the
118 preparation in DDM resembles the previous structure obtained in SMALPs[25] (RMSDs
119 of TMD (residues 30-462): 1.31 Å, CD (residues 462-606): 1.65 Å, global RMSD: 1.74
120 Å). The two TMDs are tilted towards each other, forming a dimer interface at the
121 extracellular side that would enforce a bending of the membrane by ~130° against its
122 natural curvature. As speculated for the preparation of KimA in SMALPs, the tilting
123 likely was caused by a loss of lipids between the TMDs during purification. In contrast,
124 the map of KimA in amphipols shows the upright dimer architecture of the TMDs
125 (Supplementary Figure 1a). At the extracellular side an elongated non-protein density
126 is visible between both protomers, indicating that amphipol molecules wrapped
127 around both TMDs individually. The structure resembles the upright dimer that was
128 previously obtained following molecular dynamics (MD) simulations of KimA in a lipid
129 bilayer[25], with an RMSD of only 3.4 Å (Supplementary Figure 1b). The pivot point for
130 the tilting of TMDs is located around the C-terminal end of TMH 12, which extends out
131 of the membrane (residues 459-464). Otherwise, the architectures of the TMDs
132 (RMSD 0.55 Å for residues 30-462) and of the cytosolic domains (RMSD 0.66 Å for
133 residues 462-606) of KimA reconstituted in amphipols and in DDM are very similar
134 (global RMSD 1.26 Å). Also, the dimer interfaces that are mainly formed by the long
135 connecting loops (residues 462-474) and by a short beta-sheet between the two
136 cytosolic domains are not affected (Figure 2a). A comparison of the resulting new
137 structures of KimA in DDM and in amphipols with the previously reported structure
138 from SMALPs[25] suggests that all of them represent a similar inward-occluded
139 conformation. In agreement with this, a strong density for the bound substrate
140 potassium ion is observed in the 3.3 Å cryo-EM map (Figure 2b). The potassium ion is
141 coordinated by Tyr43 (2.9 Å), the carbonyl and carboxyl of Asp36 (2.8 Å and 3.3 Å,
142 respectively), hydroxyl and carbonyl of Thr230 (2.9 and 3.2 Å, respectively) and Tyr377
143 (3.2 Å). The map from KimA obtained in SMALPs suggested that Ser125 is also part of
144 the K^+ binding site, but in MD simulations this was not the case[25]. The higher resolution
145 map shows that the distance between Ser125 and the potassium ion (4.9 Å) is too long
146 for coordination (Figure 2b), confirming the MD simulations. In contrast to the
147 previous map, no extra densities are localized below Asp36 and Tyr377. Asp36 and

Tyr377 are suggested to function as an intracellular gate. Potassium ions below them were hypothesized to hinder the opening of the gate by a trans-inhibition mechanism[25]. An explanation for the lack of the inhibitory ions could be the locking of KimA in the inward-occluded conformation by other means, namely the binding of c-di-AMP, which could prohibit K^+ binding to the trans-inhibitory site from the cytosolic side. In fact, both cryo-EM maps of KimA in DDM and in amphipols clearly show two non-protein densities in the CDs, which were modelled as c-di-AMP. Each binding site is formed mostly by the cytosolic domain of one monomer (Figure 2c): One of the phosphodiesters of c-di-AMP is coordinated through two hydrogen bonds and a salt bridge between the non-esterified oxygen atoms and the amide groups of Gln581 and Ser582 and the guanidinium group of Arg337. One adenosine moiety forms hydrogen bonds with the backbone carbonyls of Val506 and Pro479. The hydroxyl group of the ribose of the second adenosine moiety forms hydrogen bonds with His579 and the backbone amide of Asn580. Only the second adenosine moiety is coordinated by residues of the neighbouring protomer; it is stacked between Val457 and Leu461 of TMH 12 that extends out of the membrane and Arg337, located in the loop between TMHs 8 and 9. Arg337 is the only residue of the TMD that engages in a strong interaction with c-di-AMP, making it a prime candidate for the transmission of the inhibitory effect of bound c-di-AMP.

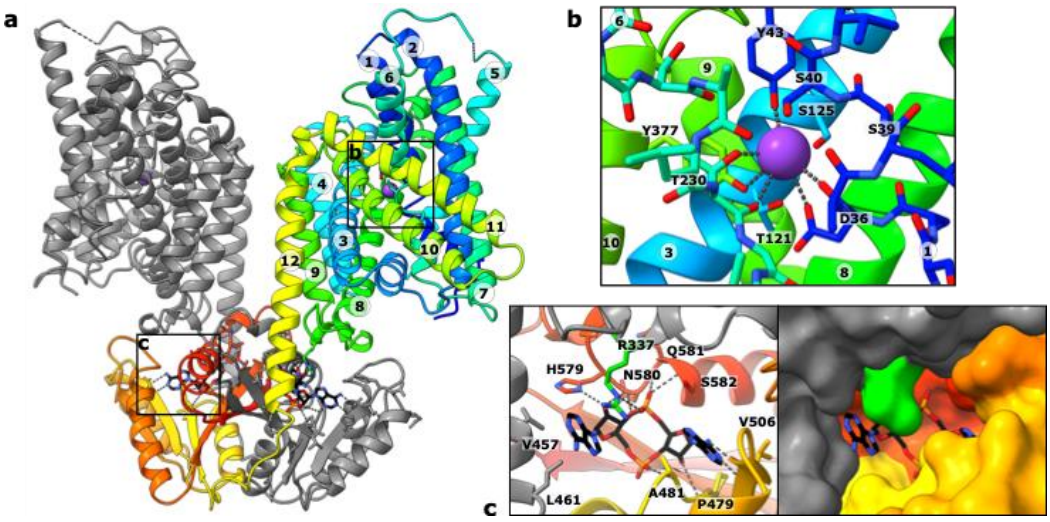


Figure 2: Cryo-EM structure of KimA with c-di-AMP bound. a) KimA homodimer with c-di-AMP bound between the swapped CDs and TMDs of the grey and rainbow-coloured monomers. **b)** K^+ binding site of one TMD. The substrate potassium ion is coordinated by Tyr43, Asp36, Thr230 and Tyr377 with distances ranging between 2.8 and 3.3 Å. **c)** The c-di-AMP binding pocket is mostly formed by residues of the cytosolic domain. Only Arg337 of the TMD strongly interacts with c-di-AMP. KimA was purified after co-production with CdaA and solubilized with DDM. A ten-fold molar excess of c-di-AMP was added after purification.

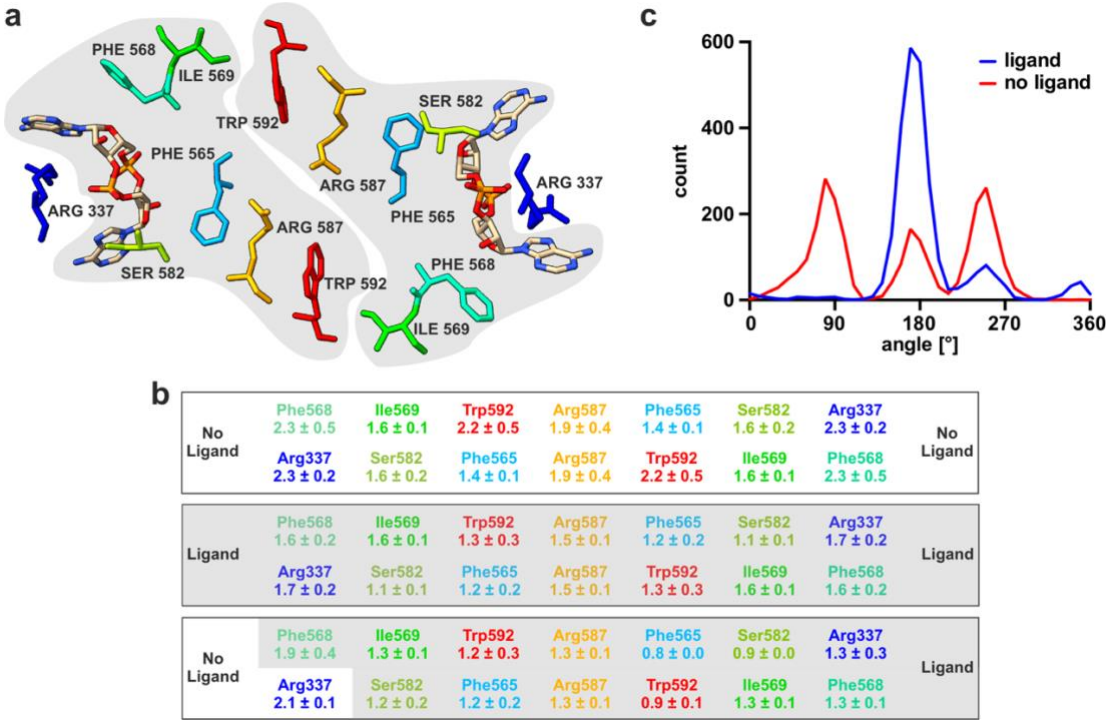


Figure 3: Cooperativity network between cytosolic domains of KimA dimer. a) Top view of dimeric CDs highlighting bound c-di-AMP and residues linking both binding sites. Protomer affiliation highlighted in grey. **b)** RMSF (Å) of residues connecting binding pockets in the absence, presence or with asymmetric binding of c-di-AMP. Priming of unoccupied binding site by ligand binding to the opposing protomer is highlighted in grey. Data in top row are from 3 x 2.2 µs simulations, as for the middle row, and data from the bottom row are from 3 x 500 ns simulations. **c)** The presence of c-di-AMP has a considerable effect on the range of χ_2 angles sampled by the Trp592 sidechain.

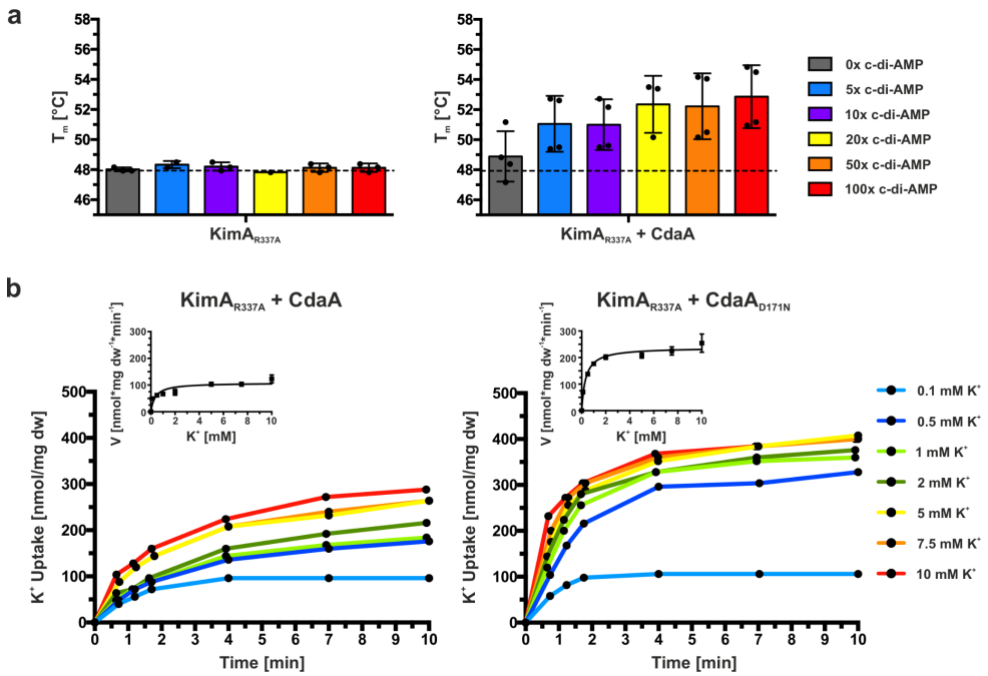
C-di-AMP binding to one binding site primes the other

The structural insights were used to further investigate the binding of c-di-AMP to KimA. In particular, we strove to understand how pre-binding of c-di-AMP facilitates further c-di-AMP binding, as suggested by DSF measurements. To this end, MD simulations were run of the KimA dimer either with or without c-di-AMP, with c-di-AMP remaining stably bound to KimA throughout the simulations where present, with an RMSD relative to the protein of 0.25 ± 0.07 nm over 3 repeats of ca. 2.2 µs. Analysis of the data led to the identification of a cooperativity pathway between the two binding sites (Figure 3a): Ser582, which is in direct contact with c-di-AMP, connects through Phe565 and Arg587 with Trp592. Trp592 links both CDs through a contact with Ile569 across the dimer interface. Ile569 is a direct neighbour of Phe568 which, like Phe565, delimits the c-di-AMP binding pocket. The root mean square fluctuation (RMSF) of all these residues was significantly reduced compared to the apo-protein when c-di-AMP was present in both binding sites, suggesting a stabilising effect of c-di-AMP binding on the CD (Figure 3b, Supplementary Videos 1 and 2). Surprisingly, if

199 only one binding site was occupied by c-di-AMP, a similar stabilising effect was
200 observed and all connecting residues of the apo-protomer also showed lower RMSF
201 as if c-di-AMP was bound (Figure 3b, grey highlighted, Supplementary Video 3). It is
202 possible that the more stable sidechain of Trp592 (Figure 3c) restricts the motion of
203 its counterpart Ile569 in the other protomer, which via the described network
204 facilitates the priming of the second, empty c-di-AMP binding site.
205

206 **Arg337 increases the affinity for c-di-AMP but is not essential for the inhibition**

207 As Arg337 was the only residue of the TMD identified to strongly interact with c-di-
208 AMP in the binding pocket, it was suspected to transmit the inhibitory effect of c-di-
209 AMP binding from the cytosolic domain to the TMD. DSF measurements showed that
210 the mutated variant $KimA_{R337A}$, which should be unable to link c-di-AMP binding to the
211 TMD, was still able to bind c-di-AMP when co-expressed with *cdaA;* however, a 20-
212 fold excess was required for reaching the same thermal stabilisation as with a three-
213 fold excess for the wildtype (Figure 4a), suggesting a decreased affinity for c-di-AMP.



214
215 **Figure 4: Arg337 is essential for high-affinity binding of c-di-AMP. a)** Melting temperatures of KimA
216 R337A purified from cells with CdaA absent ($KimA_{R337A}$) or present ($KimA_{R337A}$ + CdaA), and incubated
217 with an increasing c-di-AMP concentration given in x-fold molar excess over KimA. Determined with
218 Differential Scanning Fluorometry (DSF). Dashed line indicates T_m of KimA WT without c-di-AMP
219 addition. Data points represent the average and error bars the standard deviation of measurements
220 from at least biological triplicate. **b)** Whole-cell K^+ uptake assay in *E. coli* LB2003 cells producing
221 $KimA_{R337A}$ with active (CdaA) or inactive ($CdaA_{D171N}$) diadenylate cyclase. Michaelis-Menten plot shown
222 in graph. Representative experiment shown (n= 3).

223 To determine how the mutation affects the inhibition by c-di-AMP, *in vivo* potassium
224 uptake studies were performed. In the presence of active CdaA, the potassium uptake
225 velocity remained rather high (Figure 4b), but the inhibition by c-di-AMP was not
226 completely abolished. The V_{max} value for K^+ uptake by $KimA_{R337A}$ in the presence of
227 active CdaA was reduced by ~44% to 111.7 ± 25.5 nmol*mg dw^{-1}*min^{-1} when
228 compared to the co-expression of $KimA_{R337A}$ with inactive CdaA, where a V_{max} value
229 for K^+ of 198.9 ± 35.8 nmol*mg dw^{-1}*min^{-1} was determined (Table 1). To address
230 whether the remaining inhibition was related to c-di-AMP binding to KimA or caused
231 by the production of c-di-AMP in general, variant $KimA_{A481WS582W}$ that should be unable
232 to bind c-di-AMP because of a sterically blocked binding pocket, was analysed. The
233 loss of c-di-AMP binding was confirmed by DSF measurements (Supplementary Figure
234 2a). This loss of binding was accompanied by the insensitivity of the *in vivo* potassium
235 uptake through $KimA_{A481WS582W}$ to inhibition by c-di-AMP (Supplementary Figure 2b).
236 The uptake velocities were comparable in the presence of an active and inactive
237 cyclase, respectively (Table 1). In conclusion, Arg337 appears to be important for high-
238 affinity c-di-AMP binding but is not necessarily required for communicating c-di-AMP
239 binding to the TMD for inhibition.
240
241 **C-di-AMP-induced inhibition of KimA**
242 C-di-AMP does not directly interact with the TMD, and long-range communication is
243 required because the c-di-AMP binding site is ~37 Å away from the substrate K^+
244 binding site. The question remains how c-di-AMP binding to the CDs of KimA controls
245 K^+ transport at a distance. To address this question, we further evaluated the MD
246 simulations performed in the presence and absence of c-di-AMP. They showed a
247 significant stabilisation of the whole intracellular loop between TMH8 and TMH9 upon
248 c-di-AMP binding, with the largest decrease of the RMSF for Arg337 from 2.3 ± 0.2 Å
249 to 1.7 ± 0.2 Å. However, KimA with an Arg337 to alanine mutation showed a similar
250 reduction of flexibility of the loop in the presence of c-di-AMP (Figure 5a). This
251 suggests that inhibition by c-di-AMP is transmitted to the TMD via this loop.
252 The MD simulations further revealed that one consequence of c-di-AMP being
253 removed from the system was a movement of TMH6 away from TMH3, leading to an
254 opening of the TMD at the cytosolic end. This effect can be seen by measuring the

255 distance between two residues sitting underneath the intracellular gate, Tyr118
256 (TMH3) and Asn237 (TMH6). These typically interacted closely (<4 Å) when c-di-AMP
257 is bound (Figure 5b, Supplementary Figure 3a), which likely contributes to locking the
258 TMD in an inward-occluded conformation. When c-di-AMP was removed this
259 interaction was broken (Figure 5b), and the distance frequently shifted from about 4
260 Å to 8 Å (Supplementary Figure 3b). Distancing of Tyr118 and Asn237 can also be seen
261 using principal component analysis (PCA) of the MD data (see Methods), which
262 revealed a rearrangement of the central TMHs in the non-liganded state, with TMH6
263 moving ca. 4 Å away from TMH3 and TMH8 (Figure 5c, Supplementary Figure 4b).
264 Rearrangement of TMH6 was not observed in the liganded state (Supplementary
265 Figure 4a + c). The MD data reveal that, whilst the initial bound K^+ remain tightly bound
266 in the c-di-AMP-bound state, they are free to rapidly exchange with the bulk solvent
267 in the c-di-AMP-free state, suggesting a switch to an inward-open conformation. In
268 agreement with the MD simulations, a mutation of Tyr118 or Asn237 to alanine led to
269 a loss of inhibition by c-di-AMP in potassium uptake assays, while binding of c-di-AMP
270 was still possible (Figure 5d, Table 1, Supplementary Figure 5). The mutations seem to
271 have abolished the observed interaction from the MD simulations, and therefore
272 reduce the ability of c-di-AMP to lock KimA in an inward-occluded state. The
273 significantly increased K_m of 19 mM and 8 mM for $KimA_{Y118A}$ and $KimA_{N237A}$,
274 respectively, is in agreement with the assumption that these residues line the exit
275 pathway to the cytosol and suggests their involvement in ion release.
276 In conclusion, the MD simulations together with the functional data suggest that the
277 binding of c-di-AMP (Figure 5e-1) communicates to the TMD via a stabilisation of the
278 TMH8-TMH9 loop (Figure 5e-2, blue), which in turn affects the positioning of TMH8
279 and TMH3 relative to TMH6 (Figure 5e-3 pink, green, yellow). These conformations
280 would appear to regulate the switching between inward-occluded and inward-open,
281 thereby presenting a possible mechanism of c-di-AMP inhibition of KimA.

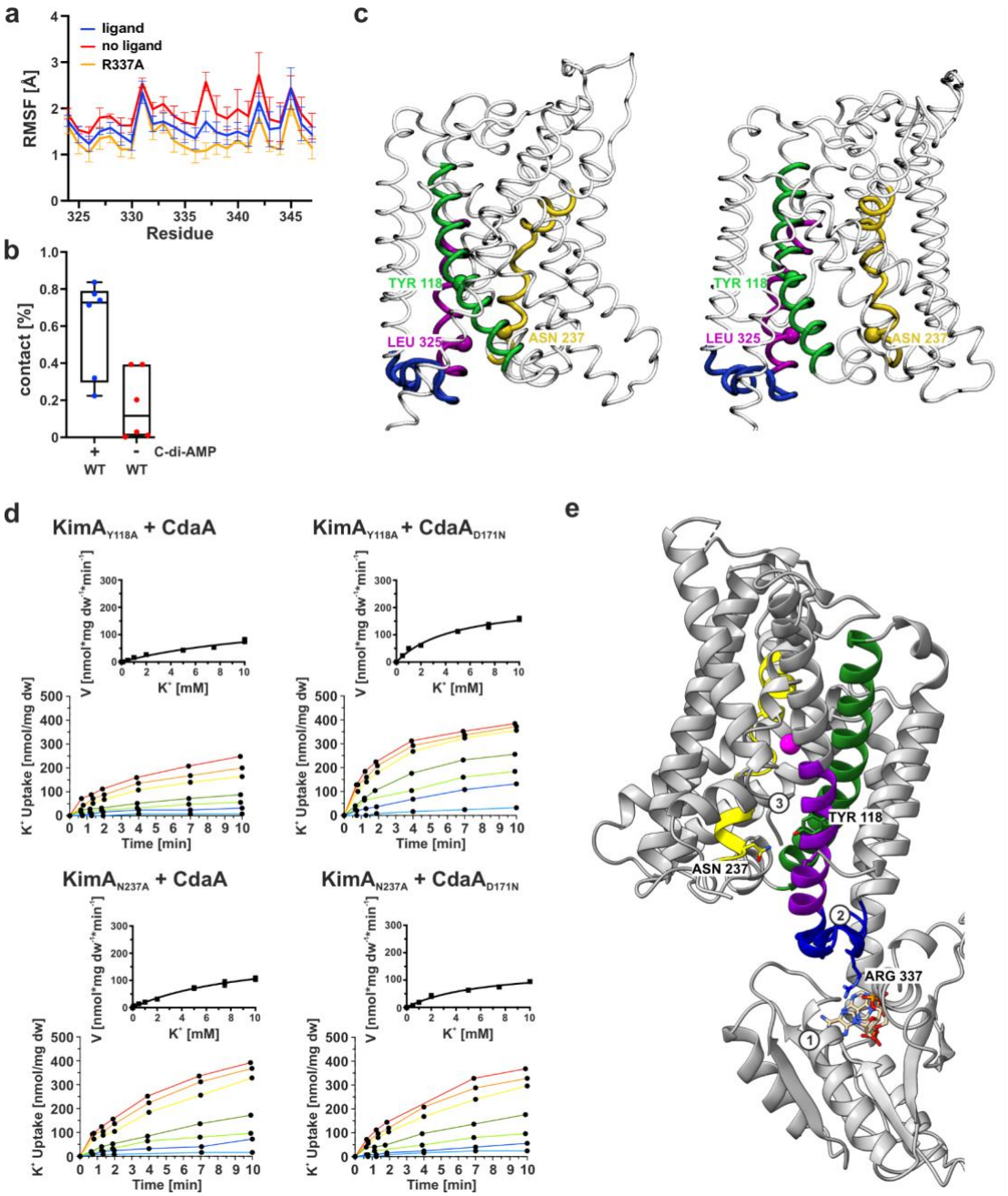


Figure 5: Transduction of inhibition from CD to TMD upon c-di-AMP binding. a) RMSF of binding loop between TMH 8 and 9 determined by MD simulations. **b)** MD simulations reveal an increased likelihood of interaction (distance ≤ 4 Å) between Tyr118 and Asn237 upon c-di-AMP binding compared to when no ligand is bound or when Arg337 is changed to alanine. Interactions were analyzed over 3 x 2.2 µs simulations for the system with and without c-di-AMP. **c)** Views of the KimA TMD in the eigenvector 1 extreme states from simulation data with no ligand bound, showing a move of TMH6 (yellow) away from TMH3 (green) and TMH8 (pink). The loop connecting TMH8 and 9 is shown in blue. The same data for the ligand-bound state can be found in Supplementary Figure 4 along with structures for eigenvector 2 and the relative view from the input model. **d)** Reduced inhibition of potassium uptake in whole-cell K^+ uptake assays when Tyr118 or Asn237 were mutated to alanine. K^+ uptake for $KimA_{Y118A}$ or $KimA_{N237A}$ variant in the presence of active CdaA or inactive $CdaA_{D171N}$. Added K^+: light blue: 0.1 mM, blue: 0.5 mM, light green: 1 mM, green: 2 mM, yellow: 5 mM, orange: 7.5 mM, red: 10 mM. Michaelis-Menten diagram shown in graph. Representative experiment shown (n= 3). **e)** Model for signal transduction: Upon c-di-AMP binding to CD (1), the binding loop (blue) rigidifies (2), TMH8 and 3 become restricted in their mobility, leading to more constant interaction between the cytosolic ends of TMH3 and 6, e.g. Tyr118 and Asn237 (3). The opening of the intracellular gate is abolished.

Discussion

Our results provide first structural and functional insights into how c-di-AMP inhibits proteins of the KUP transporter family. Interestingly, the binding mode differs from other c-di-AMP-regulated transport proteins such as potassium channel KtrAB[19], osmolyte transporters OpuC[29] and OpuA[30], and potassium exporter KhtTU[22], where c-di-AMP binds at dimer interfaces in a symmetric binding pocket. In KimA, two c-di-AMP molecules bind to the dimer in an asymmetric binding pocket. Different to the binding sites in the RCK_C domains of KtrA and KhtT and in the CBS domains of OpuA and OpuC, in which two arginine residues, one per protomer, coordinate both phosphate esters, only one of the two phosphate esters of c-di-AMP is coordinated by an arginine (Arg337) in KimA. In fact, not even Arg337 is essential for the binding of c-di-AMP or its inhibitory function, although it appears to increase the affinity. Instead, the major linker to the TMD is the entire binding loop between TMH8 and 9, which becomes more restricted in its movement once c-di-AMP is bound. This stabilisation may limit the freedom of movement of TMH8, which then restrains TMH3 and TMH6, for instance through interactions between Leu325, Tyr118 and Asn237. MD simulations indicate that through these interactions, opening of the inner gate is hindered, arresting KimA in an inward-occluded conformation. Consequently, the rocker switch movement can no longer take place (Figure 6). When c-di-AMP is removed, KimA is once again able to open up to exchange K^+ with the bulk solvent, as seen in our simulations.

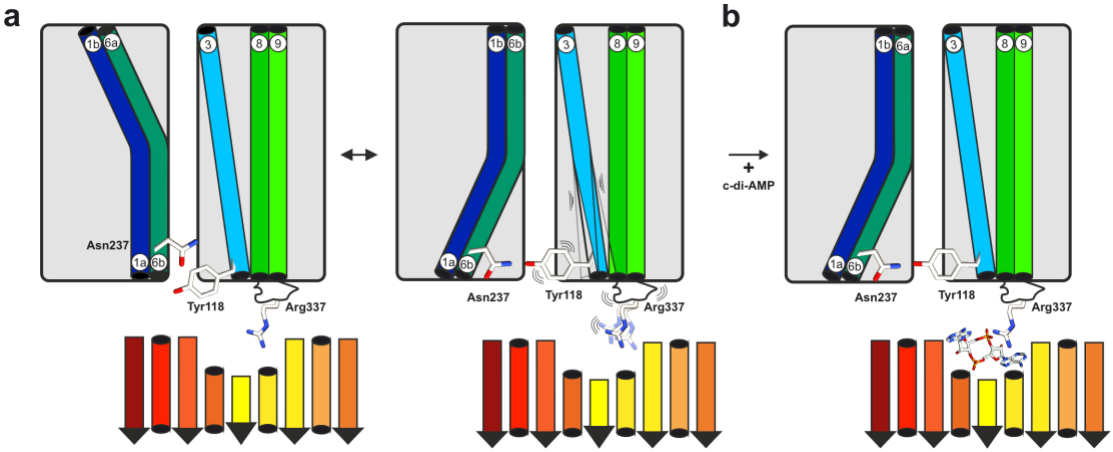


Figure 6: Schematic model of c-di-AMP-induced inhibition of KimA. a) KimA undergoes a rocker switch movement to fulfil proton-coupled potassium uptake. The binding loop is unrestricted in its movement. Tyr118 and Asn237 are only in loose contact. **b)** C-di-AMP binding rigidifies the binding loop. Helix 8 leans on helix 3, stabilizing the interaction between helix 3 and 6 bringing Tyr 118 and Asn237 closer. KimA is then locked in an inward-occluded conformation, preventing the opening of the inner gate.

326 To elucidate how conserved the inhibition mechanism is among other KUPs we
327 performed a structure-based sequence alignment with AlphaFold predictions of four
328 other KUPs of gram-positive bacteria as well as Kup from the gram-negative *E. coli*,
329 which have been functionally but not structurally characterized (Supplementary
330 Figure 6). While Kup from *E. coli* is not regulated by c-di-AMP because the second
331 messenger is lacking in *E. coli*, KimA from *Listeria monocytogenes* ($KimA^{Lmo}$)[26], and
332 KupA and KupB from *Lactococcus lactis* IL1403[31] showed growth inhibition by c-di-
333 AMP. Further, c-di-AMP binding was shown for KupA and with lower efficiency for
334 KupB. The regulation of KimA from *Staphylococcus aureus* ($KimA^{Sau}$) by c-di-AMP
335 remains unclear[26]. In agreement with the lack of c-di-AMP in *E. coli*, Kup from *E. coli*
336 does not share any of the characteristic residues Arg337, Tyr118 or Asn237 with KimA
337 from *B. subtilis*. In contrast, Arg337 is conserved in $KimA^{Lmo}$, $KimA^{Sau}$ and KupA. KupB
338 lacks an equivalent amino acid, which could explain the weaker c-di-AMP binding.
339 KupB is still inhibited by c-di-AMP, supporting our assumption that Arg337 is not
340 required for transmission of the inhibition to the TMD. Residues Tyr118 and Asn237
341 are only conserved in $KimA^{Sau}$, while KupA, KupB and $KimA^{Lmo}$ share a glycine and a
342 serine, respectively, at those positions. In the binding pocket, $KimA^{Sau}$ shows
343 conservation for Pro479, His579, Asn580, Gln581 and similarity for Val457 and Ser582
344 (Ile/Thr in $KimA^{Sau}$, respectively). The KUPs show conservation only for Gln581 and like
345 $KimA^{Sau}$ have a Thr at the position of Ser582, with the exception of Kup from *E. coli*,
346 which has a lysine and leucine. Therefore, inhibition of $KimA^{Sau}$ by c-di-AMP is likely to
347 be similar to KimA from *B. subtilis*, while for the other homologs some variations are
348 to be expected.
349
350 A surprising observation was that c-di-AMP only bound to KimA *in vitro* when at least
351 a proportion of c-di-AMP was co-purified along with KimA. Structurally, there is no
352 obvious reason for this observation. The MD simulations showed that c-di-AMP stably
353 binds to both the upright- and the tilted-dimer arrangement. The superposition of the
354 previously solved structure in the absence of c-di-AMP with the c-di-AMP-bound
355 structures does not show any significant conformational changes in the binding
356 pocket, apart from a general stabilisation when c-di-AMP is bound. However, binding
357 of the first c-di-AMP molecule by KimA appears to be unfavourable. A direct hand over

358 of c-di-AMP from the cyclase could lower the energetic barrier for ligand binding.
359 Binding of a second molecule then appears to be strongly cooperative within dimeric
360 KimA. In agreement with this hypothesis, the only membrane-bound diadenylate
361 cyclase CdaA in *B. subtilis* is described as the major cyclase maintaining the internal c-
362 di-AMP concentration required for cell growth[32]. In other bacteria like the pathogenic
363 *L. monocytogenes*[33,34] and *S. aureus*[35] as well as *L. lactis*[36] CdaA is the only c-di-AMP
364 cyclase. Since the majority of proteins regulated by c-di-AMP are membrane bound,
365 the co-localisation of the cyclase appears advantageous if direct interaction is
366 necessary.
367
368 In summary, we revealed key elements for the inhibition of potassium uptake through
369 KimA by c-di-AMP. We show the first structure of a KUP transporter with bound
370 secondary messenger c-di-AMP and confirmed its binding *in vivo* and *in vitro*. Based
371 on our findings we propose a network for the inhibition of KimA by c-di-AMP at a
372 distance. Structures of outward-open and inward-open KimA are necessary to confirm
373 the proposed transport cycle and validate the mode of inhibition by c-di-AMP.
374
375 **Material and Methods**
376 *Cloning of KimA variants using site-directed mutagenesis*
377 Point mutations were introduced into pB24KimA using site-directed mutagenesis
378 (SDM). Primer pairs of 20-40 bp length including a 1-3 bp mismatch were used to
379 introduce point mutations (Supplementary Table 1). PCR product was digested with
380 DpnI to remove template DNA and subsequently used to transform *E. coli* DH5α.
381 Colonies grown on LB agar plates with 100 µg/ml ampicillin were picked and cultured
382 in 5 ml LB with 100µg/ml ampicillin over day. Plasmids were isolated from cells using
383 NucleoSpin Plasmid, Mini kit for plasmid DNA (Macherey-Nagel). Correct mutation of
384 the plasmids was verified by sequencing performed by MicroSynth GmbH Göttingen.
385
386 *Expression and protein purification of KimA*
387 A colony of *E. coli* LB2003 cells transformed with the expression plasmids for KimA or
388 variants thereof (pB24KimA) and, if indicated, CdaA variants (pB33CdaA) where grown
389 at 37°C o/d with 180 rpm shaking in 5 ml KML (1% KCl, 1% tryptone, 0.5% yeast extract

390 (w/v)) with 100 µg/ml ampicillin and 30 µg/ml chloramphenicol if needed. An
391 overnight culture of 200 ml KML with respective antibiotics was inoculated from the
392 o/d culture and incubated at 37°C with 180 rpm. 6 l KML were inoculated with the o/n
393 culture to an OD_{600} of 0.1 and incubated at 37°C with 180 rpm. Gene expression was
394 induced with 0.002% arabinose at an OD_{600} of 1. Cells were grown for 1.5 h and then
395 harvested by centrifugation. Cells were resuspended in buffer containing 420 mM
396 NaCl, 180 mM KCl, 50 mM Tris-HCl pH 8 supplemented with 1 mM EDTA, 0.1 mM
397 PMSF, 0.3 mM benzamidine and DNase I. Cells were disrupted by passing through a
398 cell homogeniser (Stansted Pressure Cell Homogeniser FPG 12800) at 1 kbar. Cell
399 debris was removed by centrifuging at 15,000 x *g* for 15 min. Membranes were
400 harvested o/n by centrifuging at 100,000 x *g*. Membranes were resuspended in
401 aforementioned buffer (100 mg/ml) and homogenised. Membranes were solubilised
402 for 1 h at 4°C with 1% of DDM (Glycon). Unsolubilised proteins were removed by
403 centrifuging 30 min at 135,000 x *g*. Supernatant was incubated with 2 ml Ni^{2+}-NTA
404 resin for 1 h at 4°C. Ni-NTA was washed with 50 column volumes of buffer containing
405 140 mM NaCl, 60 mM KCl, 20 mM Tris-HCl pH 8 and 0.04% DDM supplemented with
406 50 mM imidazole. Protein was eluted by using 500 mM imidazole in aforementioned
407 buffer. Protein was further purified via size exclusion chromatography using a
408 Superose6 Increase column (GE Healthcare/Cytiva) preequilibrated with
409 aforementioned buffer without imidazole addition. Protein was concentrated and
410 incubated with c-di-AMP if needed for DSF, HPLC-MS and cryo-EM experiments.
411
412 *Preparation of KimA in amphipols*
413 Protein expression and purification was performed as described above until binding
414 to Ni^{2+}-NTA. The beads were washed with 50 column volumes of buffer (140 mM NaCl,
415 60 mM KCl, 20 mM Tris-HCl, pH 8) containing 60 mM imidazole and a reduced DDM
416 concentration of 0.025%. KimA was eluted by making use of the C-terminal HRV-3C-
417 Protease cleavage site. Therefore, the beads were incubated with 3C-Protease for
418 1.5 h at 4°C. The beads were washed twice with buffer containing 0.025% DDM. The
419 elution fraction containing KimA was concentrated and filtered. For the
420 detergent/amphipol exchange, KimA at a concentration of 12 mg/ml and amphipols
421 PMAL C8 (10% in water; Anatrace) were incubated at w/w ratio of 1:10 for 1 h at 4°C.

422 To remove excess detergent, the sample was incubated with biobeads at a detergent-
423 to-biobeads weight ratio of 1:100 overnight. The biobeads were removed and the
424 sample was loaded onto a Superose6 Increase 200 10/300 GL column (GE Healthcare)
425 equilibrated to cryo-EM buffer (50 mM NaCl, 50 mM KCl, 20 mM Tris-HCl, pH 8). KimA-
426 containing fractions were pooled and concentrated to 2.25 mg/ml for cryo-EM.
427
428 In vivo *whole cell potassium uptake*
429 Protocol was adapted from[25,37]. A colony of *E. coli* LB2003 cells transformed with the
430 expression plasmids for KimA variants (pB24KimA) and CdaA variants (pB33CdaA) was
431 grown at 37°C o/d with 180 rpm shaking in 5 ml KML with 100 µg/ml ampicillin and
432 30 µg/ml chloramphenicol. 75 ml of K_{30} minimal media supplemented with the
433 respective antibiotics were inoculated with 3 ml of o/d KML culture and incubated o/n
434 at 37°C. 500 ml of K_{30} minimal media with antibiotics were inoculated to an OD_{600} of
435 0.15 and incubated at 37°C with 180 rpm. Gene expression was induced at an OD_{600}
436 of 0.4-0.6 with 0.002% arabinose. Cells were grown 1.5 h before being harvested with
437 6000 x *g* at 20°C for 10 min. The cell pellet was washed twice in 10 ml 120 mM Tris-
438 HCl pH 8. Cells were adjusted to an OD_{600} of 30 and incubated for 5 min at 37°C in a
439 water bath (130 rpm shaking). To permeabilize cells for internal potassium, 1 mM
440 EDTA was added and cells were incubated at 37°C for 7 min shaking at 130 rpm in a
441 water bath. Cells were centrifuged with 4500 x *g* for 7 min at 20°C and washed twice
442 in 200 mM HEPES TEA pH 7.5 to remove EDTA and internal potassium. Cells were
443 adjusted to an OD_{600} of 30. For uptake, cells were diluted to an OD_{600} of 3 with 200 mM
444 HEPES TEA pH 7.5 and energized with 0.2% glycerol and 0.002% arabinose. Different
445 potassium concentrations (0.1, 0.5, 1, 2, 5, 7.5, 10, 15 mM) were added and 1 ml
446 samples were taken at 0, 1, 2, 4, 7, 10 min after addition. Potassium uptake was
447 stopped as cells were centrifuged through 200 µl of silicone oil (ρ = 1.04) at 17000 x *g*
448 for 2 min. Media and oil was removed and the 1.5 ml reaction tube tip containing the
449 cell pellet was cut off. Cell pellet was resuspended in 1 ml of 5% TCA. Cells were lysed
450 by freezing at -20°C and subsequent boiling at 95°C for 10 min. Supernatant was
451 diluted with 3 ml of 6.7 mM CsCl and 4 ml 5 mM CsCl. Cell debris was pelleted by
452 centrifuging 20 min at 4000 x *g*. Internal potassium concentration was determined
453 using flame atomic absorption spectroscopy (F-AAS).

454

455 *Differential Scanning Fluorometry/Thermal Shift Assay (DSF/TSA)*

456 When ligands bind their target, they stabilize or destabilize the protein leading to a
457 shifted protein melting temperature. The melting temperature is detected *via* the
458 increasing fluorescence of the 7-diethylamino-3-(4-maleimidophenyl)-4-
459 methylcoumarin (CPM) dye which binds cysteine residues that become accessible
460 during the unfolding process[38,39]. 0.5 mg/ml purified KimA protein, coexpressed with
461 and without *cdaA*, was incubated with 0-100x molar excess of c-di-AMP over night at
462 4°C. The next day 16.67 µg/ml CPM dye was added with subsequent incubation for
463 15 min on ice in the dark. 10 min centrifugation at 17,000 x *g* at 4°C removed
464 aggregates. Melting curves were recorded in a Rotor-Gene Q 5Plex HRM system
465 (Qiagen) using a temperature range of 25-85°C with 1°C steps each 30 s and 90 s
466 prewarm. CPM was excited at 365 nm and emission was detected at 460 nm. Samples
467 were measured in technical triplicates. Melting temperatures were determined at the
468 inflection point of the resulting fluorescence curve.

469

470 *Cryo-EM specimen preparation and data acquisition*

471 UltrAuFoil R1.2/1.3 400 mesh gold grids were glow-discharged twice before the
472 application of 3 µl of either a 4.0 mg/ml solution of KimA in DDM or 2.25 mg/ml KimA
473 reconstituted in amphipols. The sample was then plunge-frozen using a FEI Vitrobot
474 Mark IV. The chamber and Whatman 595 blotting paper were equilibrated at 4°C and
475 100% relative humidity.

476 A Titan Krios (Thermo Scientific) equipped with a Gatan K3 camera in counting mode
477 and energy filter was used for imaging with the software EPU (Thermo Scientific). The
478 fluence over a broken hole was adjusted to 1.1 electrons/$Å^2$ per frame. Micrographs
479 were acquired as 50-frame movie stacks in 2.7 s or 2.3 s exposures, respectively, at a
480 nominal magnification of 105,000x with a resulting pixel size of 0.83 Å. Defocus values
481 were set in the range of -1.1 to -2.5 µm.

482

483 *Image processing and model refinement*

484 Micrographs were processed using Relion-3.1[40] (Supplementary Figure 7). The Relion
485 implementation of MotionCor2[41] was used for drift correction and dose weighting.

486 Gctf[42] was used for the initial CTF estimation. Initially 950k coordinates were picked
487 from 2,349 micrographs by Topaz[43] after training the neural network with 726
488 particles that were manually picked from 30 micrographs of KimA in DDM. After two
489 rounds of 3D classification 332k particles remained that yielded a 3.6 Å map of KimA.
490 Per particle drift correction and dose-weighting during Bayesian polishing[44] and two
491 iterations of CTF refinement improved the resolution to 3.5 Å (C1) and 3.3 Å (C2
492 symmetry applied) (Supplementary Figure 8a).
493 In the case of KimA in amphipols, 5.18 million coordinates were picked from 4,303
494 micrographs by Topaz after training the neural network with 530 particles that were
495 manually selected from 20 micrographs. Bad picks were removed by two consecutive
496 3D classifications yielding 1.99 million and 861k particles, respectively. A lowpass
497 filtered map of KimA solubilized in SMA[25] was used as an initial 3D template.
498 Additional rounds of 3D classification resulted in a set of 296k particles from the 3,417
499 best micrographs yielding a 4.1 Å map of KimA. The resolution was also improved by
500 Bayesian polishing and two iterations of CTF-refinement to 4.0 Å (C1) and 3.8 Å (C2)
501 (Supplementary Figure 8b). Further improvement of the map was achieved through
502 focused classification of symmetry expanded particles. The particles of the last
503 refinement were C2 symmetry expanded. Then density outside a wide mask that
504 contained one TMD and the cytosolic domain of the other protomer was subtracted
505 from the particle image with a Relion Particle Subtraction job. 3D classification of
506 these particles with four classes, local searches only and the same mask used for the
507 particle subtraction as a reference mask yielded one class with 307k particles. 3D-
508 autorefinement of these particles resulted in a 3.7 Å map of one half of the dimer
509 (Supplementary Figure 8c).
510 Individual domains of the model of KimA in SMA (pdb ID 6s3k)[25] were rigid body fitted
511 into the map of the upright dimer with *Coot*[45] and manually rebuilt. Models were
512 optimised using Phenix real space refinement[46] (Supplementary Table 2).
513
514 *Molecular dynamics simulations*
515 Atomistic simulations were built using the coordinates of dimeric KimA bound to c-di-
516 AMP from this study. The systems were described with the CHARMM36m force
517 field[47,48] and built into 6:3:1 POPE, POPG, cardiolipin membranes with TIP3P waters

518 and K^+ and Cl^- to 150 mM, using CHARMM-GUI[49,50]. Three bound K^+ from each subunit
519 were preserved from the input structure. The c-di-AMP molecules were
520 parameterised in CHARMM-GUI, and either included in both subunits, a single subunit,
521 or in neither subunit. Where used, mutations were made in CHARMM-GUI. Each
522 system was minimized and equilibrated according to the standard CHARMM-GUI
523 protocol. Production simulations were run in the NPT ensemble, with temperatures
524 held at 303.5 K using a velocity-rescale thermostat and a coupling constant of 1 ps,
525 and pressure maintained at 1 bar using a semi-isotropic Parrinello-Rahman pressure
526 coupling with a coupling constant of 5 ps[51,52]. Short range van der Waals and
527 electrostatics were cut-off at 1.2 nm. Simulations were run in triplicate, to ca. 2.2 µs
528 for the wild-type systems with c-di-AMP present or apo, or to 500 ns for the Arg337
529 to alanine mutation or asymmetric c-di-AMP occupancy systems. In total, ca. 16 µs of
530 data were gathered.
531 PCA was carried out on the C-alpha atoms of each KimA monomer. The trajectories
532 for each subunit and each of the 3 repeats were concatenated before analysis for a
533 total of 6.6 µs sampling per condition. PCA was performed using the gmx covar and
534 gmx anaeig programs. For both ligand and no ligand conditions, ca. 1700 eigenvectors
535 were found, of which eigenvectors 1 and 2 contributed ca. 50% of the total variance
536 (see Supplementary Figure 9). Projecting these two eigenvectors reveal good overlap
537 between all 6 KimA monomers, with a generally broader distribution along
538 eigenvector 1 for the non-liganded state (Supplementary Figure 10).
539 All simulations were run in Gromacs 2020.1[53]. Data were analysed using Gromacs tools
540 and VMD[54] Plots were made using Prism 9 (GraphPad).
541
542 **Data availability**
543 Data supporting the findings of this manuscript are available from the corresponding
544 authors upon request. A reporting summary for this Article is available as a
545 Supplementary Information file. The source data of the DSF measurements and the
546 whole-cell transport assays are provided as a Source Data file. The cryo-EM maps with
547 amphipols and DDM were deposited in the wwPDB with accession codes EMD-15895
548 and EMD-15894, respectively, and the models with PDB-ID 8B71 and 8B70.
549

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Author contributions

J.V. and I.H. conceived this study. All authors designed the experiments. M.F.F. performed in vivo and DSF experiments. M.F.F., Y.H. and I.T. purified KimA for cryo-EM. J.P.W, J.S.S. and J.V. performed the cryo-EM analysis, and built and validated the atomic models. R.A.C. performed MD simulations. All authors participated in the data analysis. M.F.F., J.P.W. and Y.H. wrote the initial draft, all authors participated in manuscript editing and revision. P.J.S., J.V. and I.H. supervised work and secured the funding for this work.

Competing interests

The authors declare no competing interests.

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