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Petya V. Krasteva, Krista Michelle Giglio, Holger Sondermann. Sensing the messenger: The diverse ways that bacteria signal through c-di-GMP. *Protein Science*, 2012, 21, pp.929 - 948. <10.1002/pro.2093>. <hal-03117469>

HAL Id: hal-03117469

<https://cnrs.hal.science/hal-03117469v1>

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Sensing the messenger: The diverse ways that bacteria signal through c-di-GMP

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Received 26 April 2012; Accepted 4 May 2012

DOI: 10.1002/pro.2093

Published online 16 May 2012 proteinscience.org

Abstract: An intracellular second messenger unique to bacteria, c-di-GMP, has gained appreciation as a key player in adaptation and virulence strategies, such as biofilm formation, persistence, and cytotoxicity. Diguanylate cyclases containing GGDEF domains and phosphodiesterases containing either EAL or HD-GYP domains have been identified as the enzymes controlling intracellular c-di-GMP levels, yet little is known regarding signal transmission and the sensory targets for this signaling molecule. Although limited in number, identified c-di-GMP receptors in bacteria are characterized by prominent diversity and multilevel impact. In addition, c-di-GMP has been shown to have immunomodulatory effects in mammals and several eukaryotic c-di-GMP sensors have been proposed. The structural biology of c-di-GMP receptors is a rapidly developing field of research, which holds promise for the development of novel therapeutics against bacterial infections. In this review, we highlight recent advances in identifying bacterial and eukaryotic c-di-GMP signaling mechanisms and emphasize the need for mechanistic structure–function studies on confirmed signaling targets.

Keywords: signaling; c-di-GMP; biofilm formation; protein structure; phosphodiesterase; diguanylate cyclase; c-di-GMP receptors

Introduction

In 1987, Benziman and colleagues published a landmark article describing the identification of a novel, nucleotide-based allosteric activator of *Gluconacetobacter xylinus* cellulose synthase activity, c-di-GMP. The initial discovery was accompanied by the detection of an enzymatic activity responsible for c-di-GMP production, a c-di-GMP binding protein module

regulating cellulose synthesis, and an operon controlling c-di-GMP turnover in the bacterium.^{1,2}

Fifteen years later, c-di-GMP has gained appreciation as a second messenger unique to the bacterial world, which functions as a global regulatory molecule to trigger a plethora of physiological responses. Examples include but are not limited to cell differentiation, changes in motility and surface adhesiveness, secretion of extracellular polysaccharides and proteinaceous fimbriae, host cell cytotoxicity, and virulence gene expression (Fig. 1).^{3–6} Signaling cascades employing this small molecule show evidence for multilayer impact, which includes control at the transcriptional, translational, and post-translational levels. Proteins involved in c-di-GMP mediated signal transduction are often characterized by multidomain architecture with such modularity

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Grant sponsor: NIH; Grant number: R01-GM081373 (to H.S.); PEW scholar award in Biomedical Sciences (to H.S.).

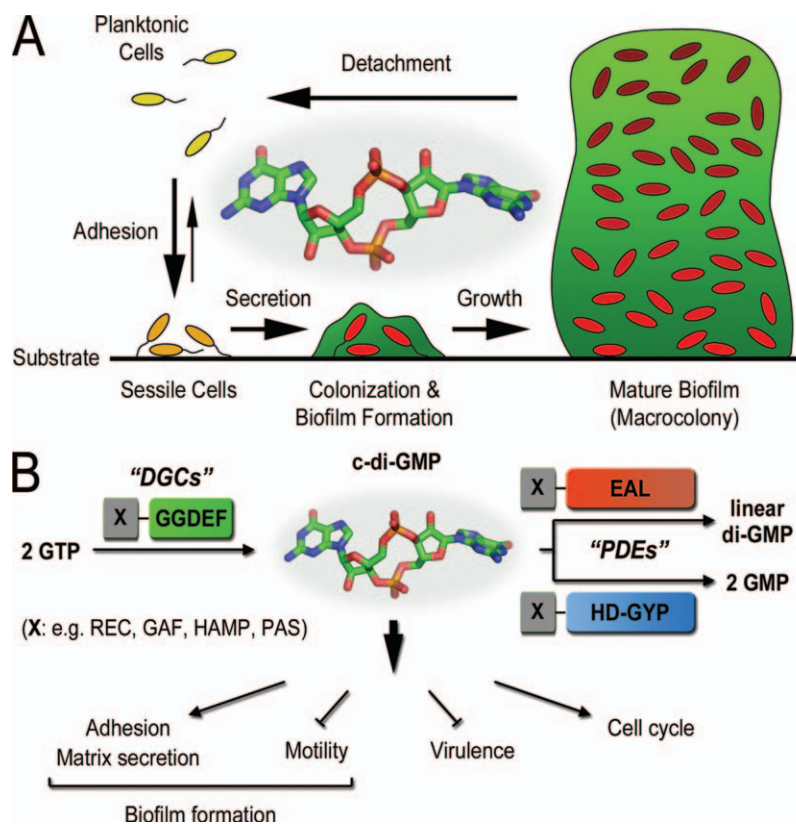


Figure 1. Overview of c-di-GMP signaling. (A) Biofilm formation. High levels of c-di-GMP in bacterial cells are often associated with cell adhesion, matrix secretion, and biofilm formation. (B) C-di-GMP signaling. Opposing activities of diguanylate cyclases (DGCs) with GGDEF domains and phosphodiesterases (PDEs) with EAL or HD-GYP domains control cellular c-di-GMP levels. The domains are often linked to regulatory domains, which control overall enzyme activity and cellular localization. In addition, several cellular programs have been shown to be under c-di-GMP control.

to allow for a variety of regulatory inputs and/or signal ramifications.^{7–9} This complexity is in stark contrast with canonical two-component transduction systems, where upon signal generation a sensor histidine kinase phosphorylates its cognate response regulator to alter the expression of a typically limited number of genes.¹⁰

The identification of a diguanylate cyclase in *Caulobacter crescentus*, responsible for c-di-GMP production during cell cycle-dependent polar differentiation, paired with bioinformatics analyses making use of the increasing number of sequenced genomes have paved the way for many of the discoveries on c-di-GMP signaling in the past decade.^{8,11–13} The metabolic function of a number of c-di-GMP signaling modules has thus been successfully identified based on conserved sequence motifs, predicted secondary structure or domain organization, inter- and intraoperon genetic environment, and phylogenetic patterns of cross-species evolution.⁸

It is now well established that c-di-GMP is generated from two GTP molecules by GGDEF domain-containing diguanylate cyclases, whereas phosphodiesterases containing either EAL or HD-GYP protein domains provide selective signal degradation

[Fig. 1(B)].^{14–18} Comparative cross-genome sequence alignments have characterized GGDEF, EAL, and HD-GYP domains as some of the most abundant protein modules encoded by bacterial genomes, with their nomenclature being a direct result of such bioinformatic analyses: GGDEF, EAL, and HD-GYP correspond to characteristic conserved amino acid motifs found in the corresponding protein domains.^{8,11–13} Interestingly, the number of these signaling modules per species roughly correlates with the organism's adaptational capacities.^{8,19} As an important example, opportunistic pathogens, which are often required to adapt to different ecological niches, typically show the highest number of c-di-GMP metabolizing enzymes per genome.

In general, increased intracellular c-di-GMP levels resulting from higher diguanylate cyclase activity lead to enhanced biofilm formation and inhibit flagellar and pilus-mediated motility.^{4,20} Conversely, low levels of the nucleotide associated with active phosphodiesterase catalysis suppress the maintenance of extracellular adhesins and promote biofilm dispersion and bacterial virulence.^{21,22} Although overexpression studies suggest a redundancy in overall effect, different diguanylate cyclases or

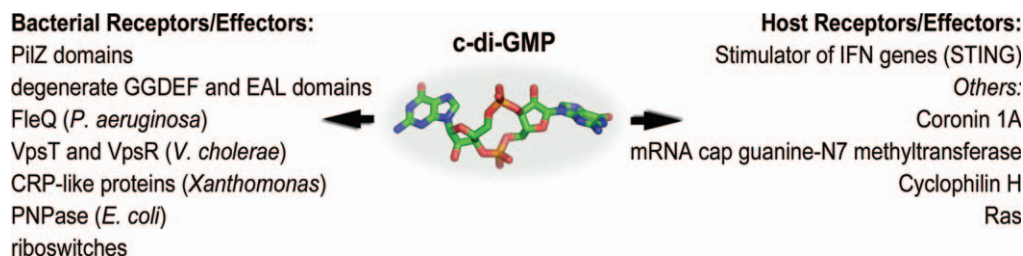


Figure 2. Effectors or receptors for c-di-GMP. Modules and proteins for c-di-GMP binding are diverse. While several bacterial effectors have been studied extensively, only one well-validated target on host cells has been discovered. Others await further characterization.

phosphodiesterases in a given genome often have enzyme specific physiological effects.^{15,23} Oftentimes proteins with similar domain architecture or enzymatic activity trigger distinct physiological responses.^{14,24} This is particularly surprising if one assumes that, as a small hydrophilic molecule, c-di-GMP is freely diffusible in the cell. A number of studies argue instead that once generated, c-di-GMP is a sequestered, rather than general, diffusive signal.^{4,25} Measurements of the intracellular levels of c-di-GMP in several bacterial species indicate concentrations in the micromolar range or lower, without taking into account probable local fluctuations. Based on the fact that most identified c-di-GMP receptors and phosphodiesterases have lower affinity constants for the nucleotide, it has been hypothesized that cellular c-di-GMP exists primarily in a protein-bound form.^{15,25} It is also a reasonable hypothesis that enzymes that produce or degrade c-di-GMP function in close proximity to the effector systems, ensuring signaling specificity and local regulation.

The complexity of c-di-GMP signaling becomes even more apparent if the nucleotide's known signaling targets are taken into account (Fig. 2). Whether functioning as intermediaries in nucleotide signal relay or as final effectors in signaling cascades, the few c-di-GMP receptors identified to date are characterized by an obvious diversity and multilevel impact [Fig. 3(A–D)].

Initial *in silico* prediction identified PilZ domains as protein modules involved in various c-di-GMP signaling pathways and predicted that they might function as direct sensors relaying the second messenger's input.²⁶ Further experimental evidence has corroborated c-di-GMP binding to several PilZ domain-containing proteins, as well as the direct involvement of these sensors in biofilm formation or expression of virulence determinants.^{27–31} These domains are expressed either alone or as fusions with other modules, including but not restricted to EAL, GGDEF, HD-GYP, PAS, and helix-turn-helix motifs.²⁶ A PilZ domain was also identified as the c-di-GMP binding module in *G. xylinus* cellulose synthase, where c-di-GMP regulatory function was first

reported. Phylogenetic and structural analyses showed that PilZ domains have low sequence conservation with the exception of a few interspersed residues responsible for nucleotide docking.²⁸ In addition, recent structural studies have shown that even highly similar PilZ domain-containing proteins can employ a markedly different molecular readout upon ligand binding.³²

Interestingly, many bacteria that utilize c-di-GMP mediated signaling for adaptation lack PilZ domains encoded in their genomes.²⁶ In addition, in some organisms PilZ signaling modules seem to control some, but not all the c-di-GMP dependent processes involved in biofilm formation and pathogenicity. For example, while PilZ domain-containing proteins in *Vibrio cholerae* are not essential for rugosity and exopolysaccharide production, some of them are required for efficient intestinal colonization during environment-to-host transition.^{29,33}

It is now established that the gamut of c-di-GMP receptors spreads far beyond the family of PilZ protein domains. Identified c-di-GMP sensors include but are not limited to bacterial riboswitches, transcription factors of various domain architectures, divergent c-di-GMP turnover domains, and allosteric sites on active or degenerate diguanylate cyclases (Fig. 2).^{22,33–41} Interestingly, high-resolution structural data have shown that this structural diversity of c-di-GMP signal recognition is not limited to the nucleotide-sensing modules, but c-di-GMP itself can adopt a variety of stable conformations [Fig. 3(E)].⁴²

Some of the identified c-di-GMP targets belong to protein families generally involved in sensing or metabolizing different nucleotide-based small molecules. For example, FleQ of *Pseudomonas aeruginosa* and VpsR of *V. cholerae* belong to the AAA+ superfamily of ATPases, but seem to function as c-di-GMP signal effectors independent of ATP binding or hydrolysis.³⁸ As another example, the Clp proteins of *Xanthomonas* sp. are classified as catabolite activator-like proteins, whose homolog in *Escherichia coli* regulates gene expression in a cAMP-dependent manner.^{40,43} Similarly, c-di-GMP turnover domains are themselves homologous to

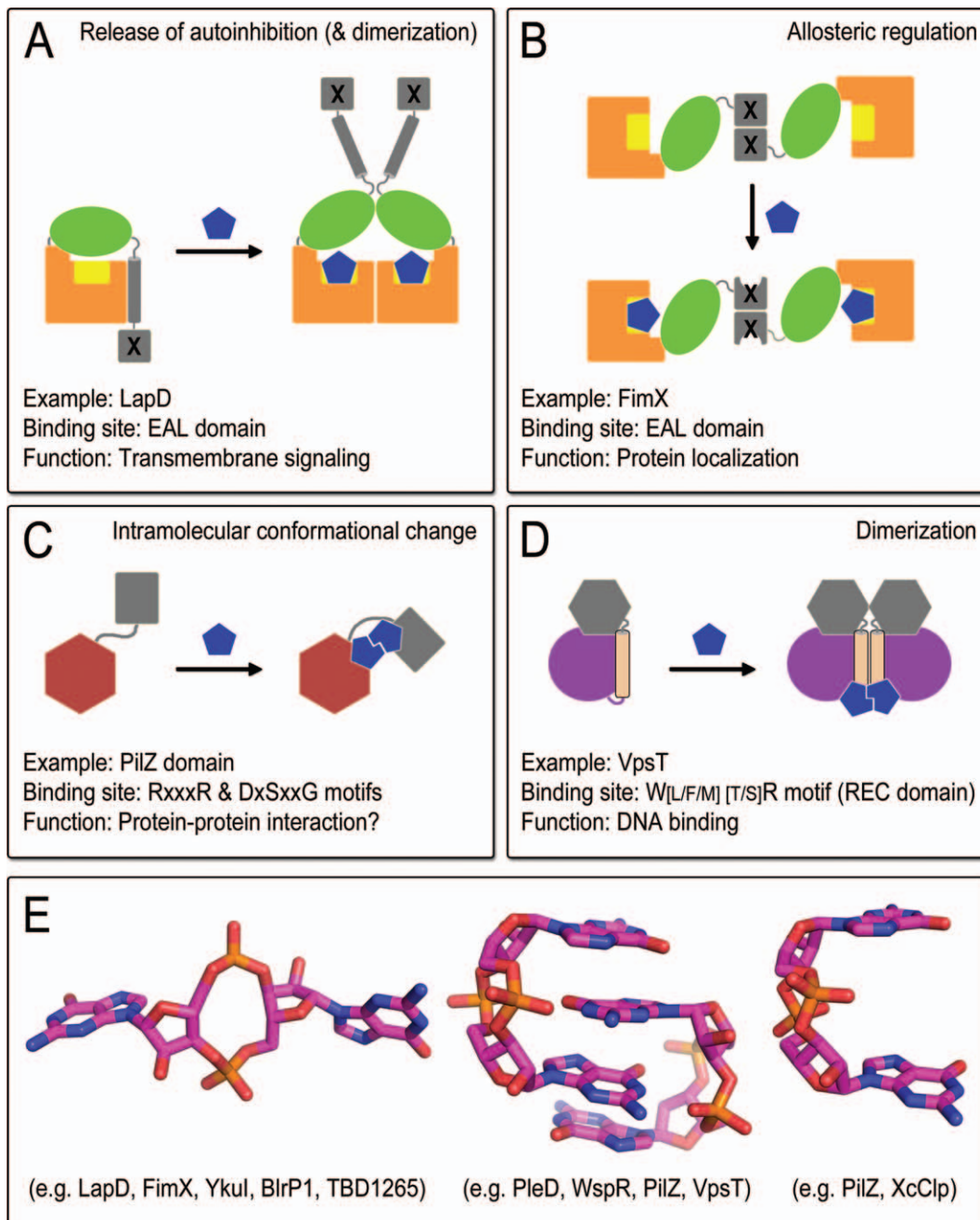


Figure 3. Modes of c-di-GMP effector function. Based on structural studies several modes of c-di-GMP action on effector proteins have been described, including the release of autoinhibitory interactions (A), allosteric regulation (B), major structural rearrangements (C) and/or dimerization (D). In addition, c-di-GMP itself can adopt several distinct conformations when bound to proteins.

protein domains involved in the synthesis or hydrolysis of different nucleic acid-based metabolites. For example, while GGDEF domains are evolutionary close to adenylate cyclases,¹¹ HD-GYP domains belong to the larger HD superfamily of phosphohydrolases with nucleotide-based substrates such as dGTP or ppGpp.⁸ This highlights the functional complexity and rapid evolution of homologous proteins in bacterial signaling networks and raises the question of whether there are a limited number of universal c-di-GMP binding motifs and/or

protein folds, or individual c-di-GMP receptors have evolved specific modes of recognition to ensure signal isolation.

Taken together, the huge number of homologous but nonredundant c-di-GMP metabolizing proteins per species, the diversity of receptor protein domain architectures and putative domain-domain interactions, the variety of adopted c-di-GMP conformations, and the widespread effects of c-di-GMP-mediated signaling phenomena complicate prediction-based approaches for the characterization of c-di-

Table I. Representative High-Resolution Structures of *c*-di-GMP Metabolizing Enzymes and Protein Receptors

Protein	Organism	Function	Conserved domains	PDB code and nucleotide-bound state	References
C-di-GMP turnover enzymes					
Diguanylate cyclases					
PleD	<i>Caulobacter crescentus</i>	C-di-GMP synthesis; polar differentiation	Two CheY-like REC domains; <u>GGDEF</u> domain	1W25 (c-di-GMP/Mg complex); 2WB4 (c-di-GMP/Mg/BeF ₃ complex); 2V0N (c-di-GMP/Mg/BeF ₃ /GTP- α -S complex)	41,44
WspR	<i>Pseudomonas aeruginosa</i>	C-di-GMP synthesis; biofilm formation and persistence	CheY-like REC domain; <u>GGDEF</u> domain	3BRE (c-di-GMP/Mg complex); 3I5B (GGDEF domain; apoprotein)	39,45
Phosphodiesterases					
YkuI	<i>Bacillus subtilis</i>	C-di-GMP hydrolysis	<u>EAL</u> domain; <u>PAS</u> -like domain	2BAS (apoprotein); 2W27 (c-di-GMP/Ca complex)	46
BlrP1	<i>Klebsiella pneumoniae</i>	Light-regulated c-di-GMP hydrolysis	BLUF sensor domain; <u>EAL</u> domain	3GFX 3GG1 (c-di-GMP/Ca/FMN complex); 3GFY (c-di-GMP/FMN complex); 3GFZ, 3GG0 (c-di-GMP/Mn/FMN complex)	47
Bd1817	<i>Bdellovibrio bacteriovorus</i>	Unknown	<u>HD-GYP</u> domain	3TMS, 3TMB, 3TMC, 3TMD (apoprotein)	48
C-di-GMP protein receptors					
PilZ domain proteins					
VCA0042/P1zD	<i>Vibrio cholerae</i>	Potential role in virulence	YcgR-N* domain; <u>PilZ</u> domain	1YLN (apoprotein); 2RDE (c-di-GMP complex)	28
PP4397	<i>Pseudomonas putida</i>	unknown	YcgR-N* domain; <u>PilZ</u> domain	3KYF (c-di-GMP complex)	31
PA4608	<i>Pseudomonas aeruginosa</i>	Potential role in biofilm formation and/or quorum sensing	<u>PilZ</u> domain	1YWU (apoprotein); 2L74 (c-di-GMP complex)	30,32
Transcription factors					
VpsT	<i>Vibrio cholerae</i>	Global transcription control; biofilm formation	Noncanonical REC domain; HTH DNA-binding domain	3KLN (apoprotein); 3KLO (c-di-GMP complex)	37
Clp	<i>Xanthomonas campestris</i>	Global transcription control; virulence	cNMP-binding domain; HTH DNA-binding domain	3IWZ (apoprotein)	40
Degenerate EAL domains					
FimX	<i>Pseudomonas aeruginosa</i>	Type IV pilus-mediated twitching motility	CheY-like REC domain; PAS domain; <u>GGDEF</u> domain; <u>EAL</u> domain	3HV9 (EAL domain; apoprotein); 3HV8 (EAL domain; c-di-GMP complex); 3HVA (GGDEF domain); 3HVB (GGDEF-EAL module; apoprotein)	35
LapD	<i>Pseudomonas fluorescens</i>	Adhesin maintenance; biofilm formation and dispersal	Periplasmic output domain; HAMP domain; <u>GGDEF</u> domain; <u>EAL</u> domain	3PJV (periplasmic domain); 3PJT, 3PJU (EAL domain; c-di-GMP complex); 3PJW, 3PJX (GGDEF-EAL module; apoprotein)	49
Others					
PNPase	<i>Escherichia coli</i>	O ₂ -dependent RNA degradation	Two RNase PH domains; KH RNA-binding domain; S1 RNA-binding domain	3CDI (apoprotein); 3CDJ (PH1-PH2 module; apoprotein)	50

Conserved domains responsible for c-di-GMP synthesis, degradation, or recognition are underlined.

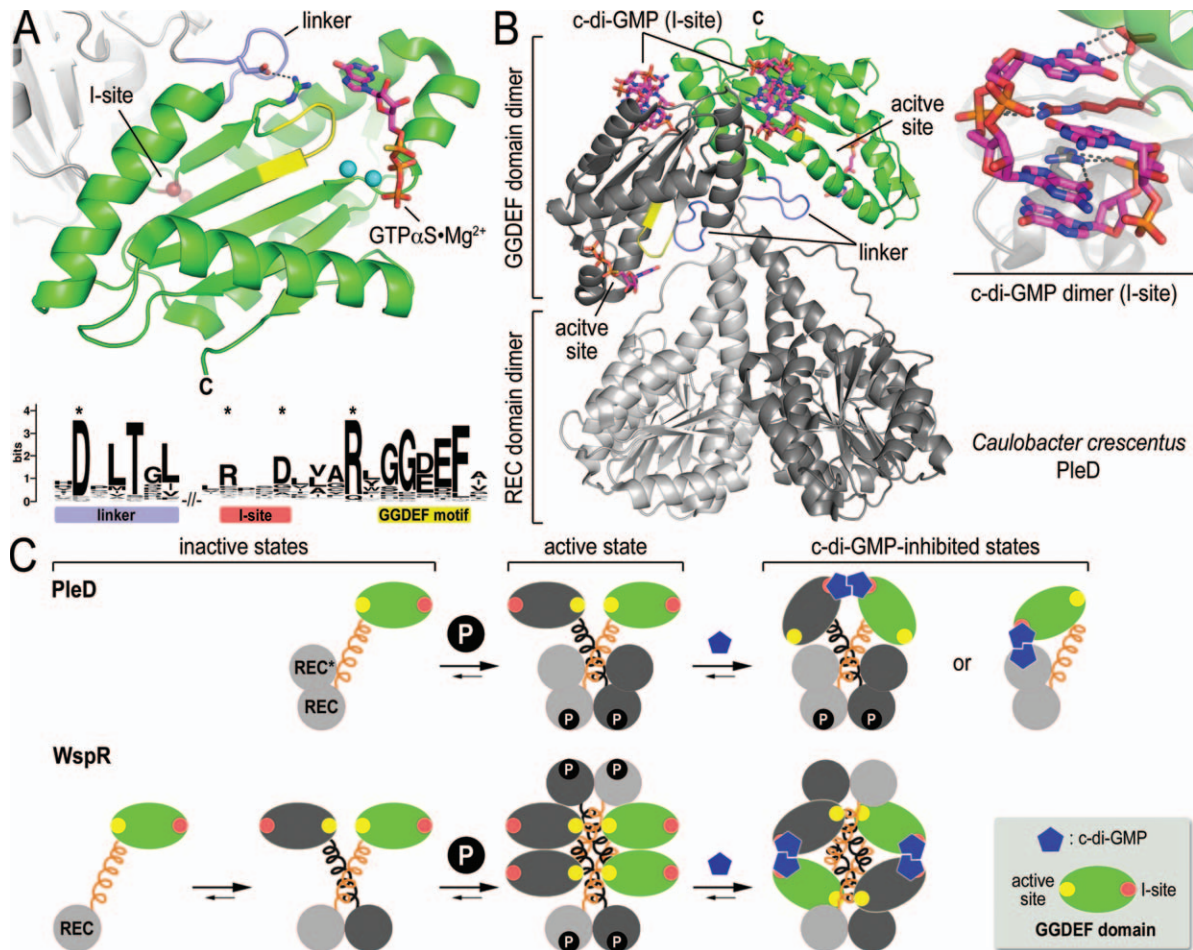


Figure 4. GGDEF domains and diguanylate cyclases. (A) Prototypical GGDEF domain structure. The PleD GGDEF domain bound to GTP- α -S is shown (PDB code: 2V0N).⁴⁴ The GGDEF motif is colored yellow. The position of the I-site is highlighted as small spheres. The sequence logo highlights several conserved motifs extending in to the linker upstream of the GGDEF domain fold. (B) Product-inhibited structure of full-length PleD. The structure of a PleD dimer activated by BeF $_3^-$ bound to one REC domain is shown. The enzyme is inhibited by a stacked c-di-GMP dimer that occupies the I-site on the GGDEF domains (inset). (C) Models for the regulatory cycle for PleD and WspR. The models highlight conserved and unique features that control enzymatic function of the two proteins.

GMP signaling pathways and underscore the need for high-resolution structure-function studies on individual signaling targets (Table I).

The Making and Breaking of c-di-GMP

Structure and regulation of diguanylate cyclases

The predicted adenylate cyclase fold of diguanylate cyclases¹¹ was first confirmed experimentally when the structures of a GGDEF domain-containing protein, PleD from *C. crescentus*, were determined.^{41,44} PleD consists of two CheY-like phosphoreceiver (REC) domains and a C-terminal GGDEF domain, responsible for GTP to c-di-GMP conversion (Fig. 4). The structures revealed conserved residues within the GGDEF domain that are involved in nucleotide and magnesium binding, as well as residues within

the N-terminal REC domain, responsible for phosphate coordination.

More importantly, structural studies on PleD and a related but not identical diguanylate cyclase from *P. aeruginosa*, WspR, elucidated many features controlling enzymatic activity.^{39,44,45,51,52} In particular, dimerization of the GGDEF domains emerged as a key requirement and a regulated step in catalysis. Although WspR contains a single N-terminal REC domain, in both cyclases the response receiver modules were proposed to oligomerize upon phosphorylation by upstream histidine kinases and thus establish the catalytically competent state. In both proteins, an extended helix of the response regulator domain proximal to the GGDEF module contributes to the oligomerization interface, with the catalytic domain resting at the tip of these stalk-like protrusions. In support of the dimerization requirement, a

catalytically active diguanylate cyclase was created by replacing all regulatory subunits by a short leucine zipper motif.⁴⁵

A structurally peculiar feature is the conformation of the loop preceding the GGDEF domain [Fig. 4(A)]. In the majority of GGDEF domain-containing proteins, this loop contains a conserved aspartate residue that forms a hydrogen bond with an equally conserved arginine located just upstream of the GGDEF motif.⁴⁹ The conformation of this loop is invariable in structures of GGDEF domain-containing proteins determined so far, and the specific arrangement may help to coordinate nucleotide binding given the close proximity of the arginine residue to the guanine moiety of the active site-bound nucleotide.⁴⁴ Probably equally important, the particular loop conformation and underlying stabilizing interactions may restrict the freedom of the GGDEF domain relative to the adjacent regulatory modules. This likely applies to the majority of diguanylate cyclase proteins, given the prevalent presence of helical secondary structure elements predicted to precede GGDEF domains in bacteria. As a result of such inherent restricted flexibility of the interdomain linkage, conformational changes in the regulatory modules that change the angle or orientation of the stalk-like motifs will have an impact on the GGDEF domain interdistance and hence catalytic activity. Interestingly, this feature is not only limited to active diguanylate cyclases but is also found in degenerate GGDEF domain-containing proteins that function as c-di-GMP receptors.⁴⁹

The structures of PleD also revealed another very common motif, an inhibitory c-di-GMP binding site (I-site) distinct from the active site and located at the base of the beta-hairpin that displays the conserved GGDEF motif in active diguanylate cyclases [Fig. 4(A,B)].⁴¹ The primary I-site is comprised of a conserved RxxD motif, where the arginine forms both hydrogen bonds, as well as π -stacking interactions with two intercalated dinucleotide molecules. The full c-di-GMP binding I-site requires secondary interactions donated by other parts of the proteins. There are at least two distinct ways that the I-site can be complemented by adjacent subunits or structural modules. In the first structure of PleD, a response regulator domain within one molecule contributes the secondary I-site residues—mainly an arginine side chain—to complete the I-site—c-di-GMP interactions. In the structure of BeF₃⁻-activated PleD and of WspR, this second arginine is located at the back of the GGDEF domain, not overlapping with the primary I-site motif.^{39,41,44,45}

Functionally, the I-site establishes a mode of feedback inhibition. Two nonmutually exclusive molecular mechanisms for feedback regulation have been proposed: allosteric inhibition or sequestration of the active site.^{39,41,44,45,53} The former model is

based on molecular dynamics simulations of the nucleotide-bound and unbound state, while the latter is apparent in several structures of c-di-GMP-bound diguanylate cyclases, including that of the leucine-zipper artificial construct.⁴⁵ In general, experimental data indicate that the main effect of c-di-GMP binding to the I-site is a restriction of conformational changes paired with sequestration of the active site dimer in a split state. Considering the apparent restricted flexibility of the linker segment connecting the regulatory and GGDEF modules, it will be interesting to obtain the structure of a GGDEF domain-containing protein *en route* to nucleotide cyclization, which has not been achieved to date. Interestingly, a recent study proposed a second mode of substrate inhibition based on the structural analysis of an isolated GGDEF domain bound to c-di-GMP at the active site.⁴² The putative inhibitory site overlaps significantly with the predicted active site and is conserved in almost all GGDEF domain sequences. It is therefore plausible that this structure could in fact depict the substrate-bound state right after nucleotide cyclization rather than a distinct mode of feedback inhibition.

Comparative analysis of the structures of PleD and WspR indicated that several of the regulatory mechanisms described above are strictly conserved, while others appear to have flexibility in their implementation [Fig. 4(C)]. In particular, the mechanistic details concerning dimerized GGDEF domains in both the active and product-inhibited states appear virtually identical. In contrast to PleD's canonical regulation via REC domain phosphorylation and subsequent dimerization, however, experimental evidence supports the notion that WspR employs a noncanonical dimer-tetramer transition to establish the active state.^{39,45}

An interesting question is whether or not substrate-inhibited diguanylate cyclases can be reactivated by phosphodiesterases as part of their cellular regulation, a phenomenon that has been demonstrated *in vitro*.³⁹ Such a post-transcriptional mechanism could be controlled by targeted spatial distribution of c-di-GMP synthesis and degradation enzymes, and may provide a fast and localized signaling response. Taken together, the structure–function studies on catalytically active diguanylate cyclases demonstrate sophisticated regulation at multiple levels, suggesting that their activity can be finely tuned in the bacterial cytosol.

Structure and regulation of EAL domain-containing phosphodiesterases

Many of the regulatory principles that establish phosphodiesterase activity have been uncovered using biochemical and biophysical approaches.^{54,55} Informed by then unpublished EAL domain structures determined by a structural genomics

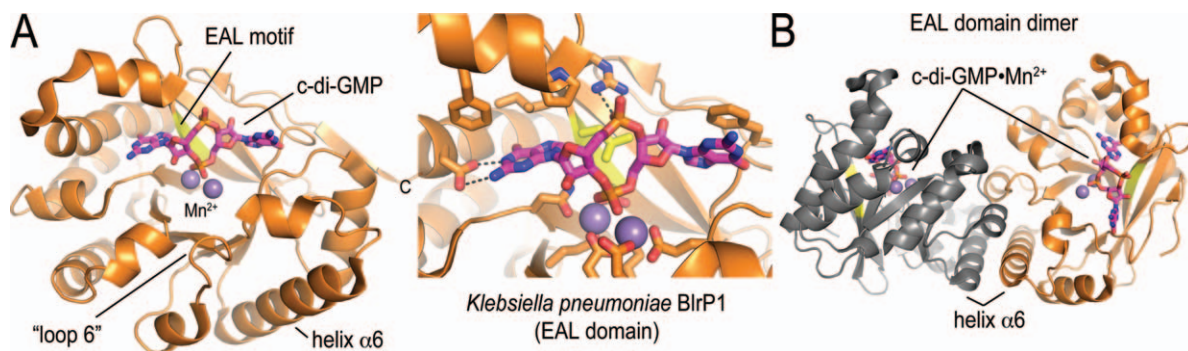


Figure 5. Structure of a canonical EAL domain. (A) EAL domain fold. The EAL domain was extracted for the full-length crystal structure of the light-regulated phosphodiesterase BlrP1 (PDB code: 3GFZ).⁴⁷ Metal ion and c-di-GMP binding sites are shown. (B) EAL domain dimer. Several structures of EAL domains show a dimeric assembly that is likely relevant for c-di-GMP binding and establishing the catalytically competent state.

consortium, a modeling approach and its experimental validation revealed several conserved motifs that were important for c-di-GMP cleavage.⁵⁵ These findings could be used to identify catalytically active EAL domains based on their sequence, which was later confirmed and extended by an independent crystallographic study.⁵⁶ Another insight came from the identification of a conserved loop in active EAL domains.⁵⁴ While the initial study established its crucial role for catalysis, crystal structures revealed its location at a EAL domain-dimer interface and assigned its functions as being important for metal coordination [Fig. 5(A)].^{46,47} This loop was referred to as loop 6 in SadR/RocR,⁵⁴ β5-α5 loop in the light-regulated phosphodiesterase BlrP1,⁴⁷ and the switch loop in LapD.⁴⁹

The first published crystal structures of an EAL domain-containing protein, YkuI from *Bacillus subtilis*, documented the triose-phosphate isomerase (TIM)-barrel fold as the building block of EAL-domain containing c-di-GMP-specific phosphodiesterases [Fig. 5(A)].⁴⁶ Structures were determined both for the apo- and c-di-GMP-bound states, which revealed the nucleotide binding site and residues crucial for catalysis. The conserved EAL motif is presented on a central strand and faces the active site, which in turn is conserved in its location to canonical TIM-barrel-containing proteins.^{46,57} Calcium, a known inhibitor of EAL domain-containing phosphodiesterases,^{2,58,59} was co-crystallized with c-di-GMP, resulting in a bound calcium ion at the active site that marked a conserved metal-binding motif.⁴⁶ Later, two independent structural studies of active phosphodiesterases, those of *Klebsiella pneumoniae* BlrP1 and *Thiobacillus denitrificans* TDB1265, established a two-metal ion mechanism for c-di-GMP catabolizing enzymes [Fig. 5(A)].^{47,56}

At the same time, nucleotide binding to the EAL domain fails to trigger any major structural rearrangements except for changes of rotamer conformation in the residues that are involved in c-di-GMP

coordination.⁴⁶ The low extent of conformational changes upon c-di-GMP-binding has been replicated in several other cases, including those of *T. denitrificans* TDB1265 and the inactive EAL domain-containing protein FimX from *P. aeruginosa*.^{35,56} Yet, an interdependence of c-di-GMP binding and EAL domain dimerization was noted in LapD, a receptor for the dinucleotide,⁴⁹ suggesting that even subtle changes can have a profound effect on the stability of intermolecular interactions.

In several structures of EAL domain-containing proteins, dimerization occurs between equivalent structural motifs within the EAL domain, which include helix α6 and loop 6 mentioned above [Fig. 5(B)].^{46,47,49,56} In addition, the overall quaternary structure of otherwise very different EAL domain-containing proteins can be conserved, as demonstrated in the comparison between *K. pneumoniae* BlrP1 and the unrelated protein YkuI from *B. subtilis*.⁴⁷ Both proteins were crystallized in a dimeric state, with the former containing a light-sensing BLUF domain in place of the PAS domain of YkuI. While YkuI was inactive in its purified form,⁴⁶ the structure of BlrP1 highlighted light-induced, allosteric regulation across the dimer interface as a likely mechanism of activation.^{47,60}

In addition to proteins that contain either an EAL or GGDEF domain, dual-domain proteins with a GGDEF-EAL domain module are also widely encoded in bacterial genomes.⁸ While in the majority of the cases studied to date either one or both domains are inactive,^{22,35,58} there is an increasing number of dual-activity enzymes being identified.^{61–63} Yet, their regulation and the functional consequences of coupled opposing activities are poorly understood.

The first structure of a HD-GYP domain

HD-GYP domain-containing proteins were first identified and implicated in c-di-GMP turnover in the plant pathogen *Xanthomonas campestris*.^{17,64} Similar to EAL

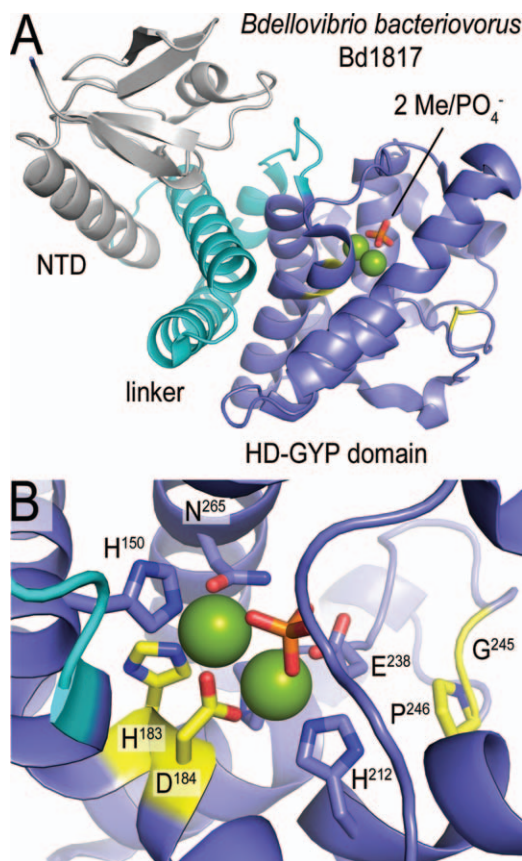


Figure 6. Structure of an HD-GYP domain fold. (A) Overall fold. The modular nature of a full-length HD-GYP domain-containing protein is shown (PDB code: 3TM8).⁴⁸ (B) Close-up view of the putative nucleotide binding site.

and GGDEF domains, they occur as single-domain proteins or as modules in multidomain proteins.^{7,8} Furthermore, they may couple directly to GGDEF domains via direct protein-protein interactions.⁶⁵

The first structural model for a HD-GYP domain was revealed recently based on a crystallographic study on *Bdellovibrio bacteriovorus* Bd1817, an inactive protein that lacks the active site tyrosine (Fig. 6).⁴⁸ The HD-GYP domain is composed of seven helices that form a compact fold, roughly resembling other proteins of the HD superfamily [Fig. 6(A)]. In addition to the HD-GYP domain, this protein also contains an N-terminal, less-characterized domain that is connected to the degenerate phosphodiesterase domain via a compact three-helix linker. Importantly, bound metal and phosphate ions mark the active site, which is located within the C-terminal domain of Bd1817 and is lined by the conserved HD-GYP motif [Fig. 6(B)]. Modeling suggested that the fold may accommodate *c*-di-GMP in an extended conformation similar to that observed bound to EAL domains. More detailed mechanistic insights would be aided by structures of active and/or *c*-di-GMP-bound HD-GYP domain-containing proteins.

C-di-GMP sensors

PilZ homology domains: The first identified *c*-di-GMP receptors. Comparative sequence analyses identified PilZ homology domains as widespread protein modules with a phylogenetic distribution pattern similar to those of GGDEF and EAL domain-containing proteins.²⁶ In addition, many of the proteins predicted to contain PilZ domains were also suggested to participate in *c*-di-GMP mediated signal transduction based on available biochemical, genetic and functional data. For example, a PilZ homology domain was detected in *G. xylinus* cellulose synthase, where allosteric regulation by *c*-di-GMP was initially described, as well as in YcgR of *E. coli*, which controls motility in a way opposite to that of a *c*-di-GMP specific phosphodiesterase.^{26,66,67}

High-resolution structural data are now available for several PilZ domain-containing proteins in their apo- and/or *c*-di-GMP-bound form.^{28,30–32,68} The core of the isolated module consists of a six-stranded antiparallel β -barrel, which is commonly found in hydrophobic ligand-binding proteins. Sequence alignment across protein family members showed low sequence conservation except for a few interspersed residues responsible for nucleotide docking.²⁸ *C*-di-GMP is primarily coordinated by two consensus motifs—RxxxR and D/NxSxxG—located at the extreme N-terminus and the β 2/ β 3 hairpin, respectively.²⁸ The nucleotide binds as a single molecule in *cis* conformation (in VCA0042/PlzD of *V. cholerae*)²⁸ or as an intercalated dimer (in PP4397 and PA4608 of *Pseudomonas* sp.)^{30–32,68} through typical H-bonding and arginine-mediated π - π stacking interactions.

This similar mode of nucleotide recognition among PilZ domain-containing proteins is in contrast to the markedly diverse modes of conformational switching observed in the various structures. In PA4608, which is a single domain protein of *P. aeruginosa*, nucleotide recognition does not affect the monomeric oligomerization state but is instead accompanied by significant rearrangements in the N- and C-terminal regions peripheral to the core β -barrel.^{30,32,68} In apo-PA4608, the N-terminal RxxxR motif, also known as the *c*-di-GMP “switch,” is unstructured and a C-terminal helix alternatively covers and exposes the hydrophobic surface of the nucleotide-binding pocket. In the presence of *c*-di-GMP, the “switch” motif wraps around the ligand and in turn ejects the C-terminal lid, creating a highly negative unilateral surface proposed to act in downstream signal transduction.^{32,68}

In contrast to PA4608, two other model PilZ domain-containing proteins, PP4397 of *Pseudomonas putida* and VCA0042/PlzD of *V. cholerae*, are found to adopt a dimeric conformation in their nucleotide-free states. *C*-di-GMP binding is accompanied by

dimer-to-monomer transition (for PP4397) or by conformational reorganization within the dimer (for VCA0042/PilZ), underscoring once again the diversity in signaling output.^{28,31} The two proteins are structurally homologous to the YcgR motility switch of *E. coli* and contain an additional N-terminal YcgR-N* domain with an overall similar fold to that of the C-terminal PilZ module. In both proteins the c-di-GMP-binding RxxxR motif forms a hinge-like connector between the two domains. While in VCA0042 this “switch” is unstructured in the apo-state, but wraps around the ligand to dramatically change the relative orientation of the two domains in the presence of c-di-GMP, in apo-PP4397 it forms a short helix participating in dimerization contacts, which become restructured upon ligand recognition and in turn contributes to dimer disassembly.

The significant effects on conformation following c-di-GMP binding to YcgR homologs were also employed to design the first c-di-GMP biosensor.⁶⁹ The fusion of full-length *E. coli* YcgR between the established FRET pair of cyan and yellow fluorescent proteins⁷⁰ was successfully applied to monitor cellular c-di-GMP distribution during the cell cycle of *C. crescentus* and *P. aeruginosa*. The imaging study also confirmed for the first time spatiotemporal control of c-di-GMP signal transduction, contrary to the more intuitive model of intracellular diffusion due to the second messenger’s small size and high solubility.

Taken together, the bioinformatic and structural studies on PilZ domain-containing proteins have revealed a plethora of protein domain architectures and mechanisms for molecular read-out. This complexity of c-di-GMP signaling mechanisms further underscores the need for high-resolution structure–function analyses on individual targets in addition to more limited, homology-based models.

Degenerate c-di-GMP turnover domains. The ensemble of bioinformatic and structural studies on c-di-GMP turnover domains indicates that many of these exist in a catalytically inactive form. However, knockout, overexpression, or mutational studies of proteins containing such “degenerate” domains have strong effects on biofilm formation or virulence. In the current model, these proteins function at the post-translational level to ensure c-di-GMP signal relay through direct protein-protein interactions.

Interplay between the PilZ and FimX proteins. As discussed above, PilZ homology domains were identified as the first bonafide c-di-GMP binding modules. Nevertheless, the PilZ protein itself, originally identified as necessary for biofilm-related Type IV pilus (T4P) biogenesis in *P. aeruginosa*,⁷¹ is incompetent for direct c-di-GMP recognition. Studies on PilZ/PA2960 orthologs from two *Xanthomonas* species revealed important struc-

tural differences from c-di-GMP-binding PilZ homology domains, including lack of the signature RxxxR and D/NxSxxG motifs responsible for ligand coordination.^{72,73} Even so, the PilZ protein has retained its role in c-di-GMP signal transduction, acting as an adaptor between the c-di-GMP receptor protein FimX and the T4P assembly ATPase PilB.⁷²

FimX is a multidomain protein with an N-terminal CheY-like REC domain followed by a PAS domain of unknown sensitivity, and a GGDEF-EAL tandem domain module. Similarly to PilZ, it was originally identified as involved in the T4P-mediated twitching motility of *P. aeruginosa* and was found to localize to a single pole of the bacterial cell together with the pilus biosynthetic machinery.⁷⁴

Although initially described as an active phosphodiesterase,⁷⁵ FimX was subsequently shown to lack significant c-di-GMP catabolizing activity based on homology modeling, enzymatic assays, and direct structural evidence.^{35,55} In particular, the crystal structures of the EAL domain showed that while the overall TIM-barrel fold of conventional phosphodiesterases is preserved, the protein lacks key residues for metal ion coordination required for catalytic activity.^{35,76} In addition, low-resolution SAXS (small angle X-ray scattering) data indicated that the full-length protein likely exists as a dimer through interactions between its N-terminal REC/PAS domains, while the EAL domains remain separated at the distant ends of the extended quaternary structure. Such an assembly excludes EAL domain dimerization through the functionally important “switch” loop discussed above.^{46,47,49,54}

Similarly to the EAL domain, the N-terminal REC domain and the GGDEF module of FimX have also evolved away from their canonical sensory and enzymatic activities, respectively: the REC domain lacks the conserved phosphoreceiver aspartate residue of archetypal response regulators and both the active and inhibitory sites on the GGDEF module show significant sequence deviation from the consensus motifs required for c-di-GMP synthesis or regulatory binding.³⁵

Regardless of the lack of catalytic and phosphoreceiver activities, the REC, GGDEF, and EAL domains of FimX have important roles in c-di-GMP signal relay. The second messenger was shown to bind the degenerate phosphodiesterase domain with high affinity, and the crystal structure of the c-di-GMP-bound EAL domain revealed a conserved mode of protein-nucleotide interactions.³⁵ Furthermore, while no major changes in the overall quaternary structure were observed between the apo-protein and its nucleotide-bound form, hydrogen/deuterium exchange experiments revealed that binding of c-di-GMP to the EAL domain triggers a long-range conformational change in the N-terminal REC domain and the adjacent linker.⁷⁷

Interestingly, residues in this region together with intact signature motifs in the GGDEF and EAL domains of FimX—G³⁴⁶DSIF and E⁴⁷⁵VL, respectively—have been shown to be crucial for proper unipolar protein localization.^{74,75,77} It has also been proposed that the C-terminal EAL domain directly binds PilZ, and therefore indirectly the T4P biosynthetic machinery, and that this interaction is stabilized by c-di-GMP.⁷²

Collectively, these data indicate that the various structural modules of FimX have evolved an important and interdependent role in the nucleotide signal relay. Important questions still remain, such as the biological significance of the long-range conformational changes upon nucleotide recognition, and the identities of additional binding partners which may be responsible for regulation of c-di-GMP-dependent T4P assembly and twitching motility.

Inside-out c-di-GMP signaling in *Pseudomonas fluorescens*: A role for degenerate domain-containing protein LapD. Similar to FimX, many GGDEF and EAL domain-containing proteins possess both types of domains in the same polypeptide chain, with variation among catalytic activities preserved among protein family members. In addition, several of these function as bacterial transmembrane receptors and contain an additional HAMP domain as a juxtamembrane signal relay module, located N-terminally to the GGDEF-EAL domain tandem.

One such receptor with HAMP-GGDEF-EAL domain organization is LapD of *Pseudomonas fluorescens*, which controls cell adhesion during biofilm formation in response to phosphate availability via direct read-out of cellular c-di-GMP (Fig. 7).^{22,49,78–81} C-di-GMP binding to the degenerate EAL domain in the cytosol is communicated via the HAMP relay module to an N-terminal output domain in the periplasm. In the c-di-GMP bound state, LapG, an otherwise active periplasmic protease, is bound to the output domain of LapD. As a result, cleavage of LapG's substrate, the large adhesin protein LapA, is prevented, leading to biofilm formation.^{22,49,79}

Structural analyses of the cytosolic domains of LapD revealed an autoinhibited conformation in the absence of c-di-GMP. In particular, the EAL domain was found to fold back onto a helical extension of the HAMP domain, dubbed as the S- or signaling-helix.⁴⁹ Interaction of the EAL domain with the S-helix are mostly hydrophobic and extend over the conserved “switch” loop crucial for canonical dimerization and metal ion coordination in active phosphodiesterases. In the closed conformation, the GGDEF domain restricts access of c-di-GMP to the EAL domain though the ligand-binding pocket remains only partially occluded. Increased c-di-GMP levels were

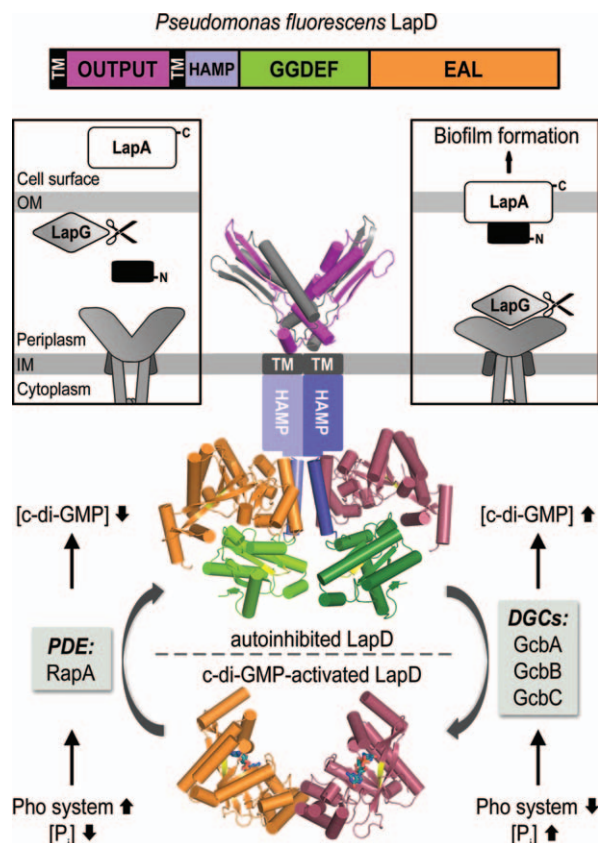


Figure 7. Model for LapD-mediated signaling. The composite structural model highlights the proposed conformational changes of the transmembrane, c-di-GMP effector LapD.⁴⁹ Switching in the intracellular domains regulates the conformation of its periplasmic output domain, which differentially interacts with a protease, LapG, that in turn controls the stability of a cell surface adhesin, and hence biofilm formation. Intracellular regulators as well as environmental signals are well-established for this system.^{80,81}

therefore proposed to compete out the inhibitory interactions and lead to opening of the cytosolic tandem domain module. Disrupted EAL domain-S-helix interactions are likely compensated by canonical EAL domain dimerization—as observed in crystal structures of the nucleotide-bound isolated module⁴⁹—as well as molecular motions of the HAMP domain helices securing signal transduction to the periplasmic output domain.^{22,49}

The isolated output domain was shown to be poised for LapG sequestration, as demonstrated in direct protein binding studies. Interestingly, the distal tips of the V-shaped dimer are formed by a conserved GWxQ motif, and mutation of the tryptophan residue alone is sufficient to abolish LapG-output domain binding *in vitro*, as well as biofilm phenotype rescue in a $\Delta lapD$ genetic background *in vivo*.⁴⁹ Overall, these data indicate that the periplasmic output domain, and in particular the conserved

tryptophan residues at the distal ends of the V-shaped module, act as a molecular caliper for LapG recognition. The observation that the isolated output domain can bind LapG is consistent with a model in which the nucleotide-free intracellular domains hold the receptor in an autoinhibited conformation that relaxes into a LapG-binding state upon c-di-GMP activation.⁴⁹

Bioinformatic analyses based on the structural model for regulation of the LapD-LapG signaling system identified homologous circuits in a variety of free-living and pathogenic species. Importantly, LapG homologs are likely to have different substrates in systems for which no clear target adhesins can be identified. In particular, proteins that contain regions homologous to the cleavage site in LapG's substrate adhesin include RTX-like bacterial toxins and the majority of such proteins are encoded in close proximity to *lapD* and *lapG* homologous genes.⁴⁹ This indicates that c-di-GMP-dependent inside-out signaling systems might have evolved a common mechanism to control periplasmic proteolysis for toxin secretion or biofilm dispersal depending on bacterial physiology and adaptational needs.

Degenerate GGDEF domains as c-di-GMP receptors. Degenerate GGDEF domains capable of c-di-GMP sensing and signal relay through a preserved I-site motif (RxxD) form another class of functionally important c-di-GMP receptors. Examples of proteins with degenerate GGDEF domains that function as c-di-GMP receptors are PelD of *P. aeruginosa*,³⁶ CdgG of *V. cholerae*,³³ the multidomain histidine kinase SgmT of *Myxococcus xanthus*,⁸² MxDa of *Shewanella oneidensis*,⁸³ and PopA of *C. crescentus*.⁸⁴ These proteins play roles in diverse physiological processes including biofilm formation, motility, and cell cycle progression. Insight in the mechanisms of c-di-GMP signal transduction is mostly limited to genetic studies, inferred protein-protein interactions, sequence alignments, or homology modeling on the isolated GGDEF domain. High-resolution structure-function analyses in the context of full-length proteins and macromolecular assemblies would thus complement the genetic and molecular studies already done, and shed light on the target-specific mechanisms of nucleotide signal transduction.

C-di-GMP sensing transcription factors. The global picture of c-di-GMP mediated signaling, where bacteria carefully monitor their environment to control motility, growth, secretion, and biofilm formation in order to rapidly adapt to changing conditions, makes regulators of gene expression likely candidates for c-di-GMP-responsive signal effectors. To date, several protein transcription factors (e.g. FleQ of *P. aeruginosa*, Clp of *Xanthomonas sp.*, VpsT and VpsR of *V. cholerae*, MrkH of *K. pneumoniae*),

as well as two distinct riboswitch classes in messenger RNAs, have been identified to directly sense c-di-GMP and regulate gene expression at the transcription initiation and post-transcriptional levels, respectively.^{34,38,40,43,85–88}

VpsT of *V. cholerae*: A noncanonical response regulator. *V. cholerae*, causative agent of the eponymous diarrheal disease, is an opportunistic pathogen with an impressive arsenal of c-di-GMP turnover domains encoded by its genome (41 GGDEF, 22 EAL, and 9 HD-GYP protein modules). In contrast, only a few and strikingly diverse protein- and RNA-based sensors for the nucleotide have been so far identified.

Initial bioinformatic searches identified five PilZ domain-containing proteins, as well as two distinct riboswitch sequences as putative targets for the second messenger. Nevertheless, the regulatory role for the c-di-GMP-sensitive RNA aptamers is limited to the expression of their immediately downstream open reading frames and quintuple knockout of all PilZ domain-containing proteins fails to disrupt colonial rugosity and biofilm formation,^{33,34} pointing toward additional targets.

In *V. cholerae*, these c-di-GMP-mediated adaptational strategies are strictly dependent on the expression of *Vibrio* polysaccharide (*vps*) genes, which in turn are under the control of two positive transcription regulators, VpsT and VpsR.^{85,89} This led to the hypothesis that the two could serve as direct c-di-GMP sensor-effectors, which was later corroborated by direct experimental evidence and, in the case of VpsT, by high-resolution structural data.^{37,90}

VpsT folds into an N-terminal receiver (REC) domain for nucleotide recognition and a C-terminal helix-turn-helix (HTH) domain for DNA binding.³⁷ It belongs to the FixJ/LuxR/CsgD family of response regulators, which are typically effectors in two-component signal transduction systems and depend on phosphoryl transfer from upstream histidine kinases for signaling input. Interestingly, both VpsT and its homolog in *Escherichia* and *Salmonella spp.*, CsgD, show divergent receiver domains where only half of the residues required for phosphotransfer are conserved and no cognate kinases have been identified to date.

In addition, the crystal structure of VpsT showed that the canonical (α/β)₅-fold of the receiver domain is extended by an additional helix, α_6 , which was identified as a common motif among available structures of LuxR-like response regulators, regardless of their input mode for regulation (Malashkevich *et al.*, unpublished).³⁷ Receiver domain oligomerization also differs significantly from that of canonical response regulators (Fig. 8). Rather than utilizing the α_4 - β_5 - α_5 structural motif for

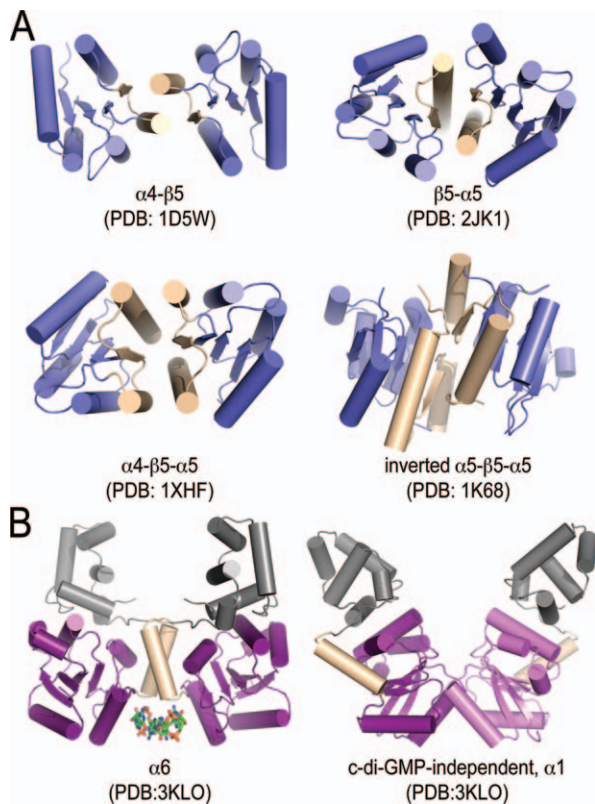


Figure 8. Overview of REC domain dimerization. (A) Canonical REC domains.^{91–94} (B) VpsT dimerization.³⁷

dimerization,⁹⁵ VpsT utilizes two different and non-overlapping dimerization interfaces.

The first dimerization interface is independent of nucleotide recognition and involves a hydrophobic groove and polar interactions formed between residues of α_1 , α_5 , and a binding pocket that extends into the canonical phosphorylation site. Point mutations disruptive to this interface were shown to have an overall stimulating effect on VpsT activity as a positive regulator of biofilm formation and a negative regulator of flagellar motility, emphasizing its biological significance.³⁷

The second dimerization interface is specific to the additional helix α_6 and is necessary for and stabilized by c-di-GMP recognition. The nucleotide-binding pocket is formed primarily by four consecutive residues located at the base of α_6 in each protomer and conserved as a W[F/L/M][T/S/P]R motif in closely related homologs. C-di-GMP binds as an intercalated dimer and its purine rings form π -stacking interactions with the side chains of the conserved tryptophans and arginines, while the threonine residues in VpsT form hydrogen bonds with the phosphate moieties of the nucleotide. The biological relevance of this second dimerization interface was confirmed in experiments showing that binding of wild-type VpsT to promoter sequences of a *vps* gene occurred only in the presence of

c-di-GMP, while mutants defective for nucleotide binding or α_6 dimerization were incapable of DNA recognition and had markedly different phenotypes.³⁷

The relative orientation of the DNA-binding and receiver domains was preserved between the crystal structures of apo-VpsT and the c-di-GMP bound protein. Interestingly, if the orientation of the HTH motifs is preserved in solution as well, it would require significant changes in DNA architecture for promoter binding to occur. This, together with the fact that VpsT bound to relatively remote fragments of the studied *vps* gene promoter, led to the hypothesis that promoter binding is accompanied by DNA loop formation, similar to the regulatory mechanisms at play at the *lac*, *gal*, and *ara* operons.^{37,96,97} To determine whether this is actually the case, what particular oligomerization state VpsT adopts at each target promoter, and what the consensus DNA sequence for VpsT recognition is, would nevertheless require further structure-function studies on the required c-di-GMP—VpsT—DNA interactions.

As mentioned above, VpsT is homologous to CsgD from *E. coli* and *Salmonella typhimurium*, a protein whose expression and function are tightly intertwined with various c-di-GMP signaling pathways and are required for the secretion of key biofilm determinants.^{98,99} Similarly to VpsT, CsgD has been postulated to be incompetent for phosphotransfer and to instead depend on small molecule binding for efficient gene expression regulation.^{99,100} Nevertheless, CsgD is unlikely to act as a c-di-GMP sensing transcription factor, as it has a divergent and highly conserved YF[T/S]Q motif at the putative ligand-binding site and direct binding experiments have ruled out c-di-GMP recognition.^{37,101}

A recent study has proposed that CsgD could nevertheless be regulated by phosphorylation, as the protein autophosphorylates in the presence of high concentrations (10mM) of acetyl phosphate *in vitro*.¹⁰¹ While this phosphorylation appears to inhibit protein binding to its cognate DNA, it remains unclear whether this is due to a distinct functional state or is an artifact of nonspecific binding and reduced protein stability. In support of the latter, acetyl phosphate does not affect CsgD function *in vivo* and a loss-of-function asparagine substitution of the conserved phosphoreceiver aspartate (D⁵⁹N) preserves phosphate binding to the protein *in vitro*. In addition, a phosphomimetic aspartate to glutamate substitution (D⁵⁹E) appears to inactivate the protein, but the mutant is indeed characterized by markedly reduced protein stability.¹⁰¹ Based on the crystal structures of homolog VpsT, the phosphoreceiver aspartate (D⁵⁹) is likely proximal to a dimerization interface, disruption of which might explain the mutant's altered stability and function and account for the observed effects *in vitro* and *in vivo*.

Although there are little data available on the regulation of CsgD *in vivo*, it is clear that the FixJ/LuxR/CsgD family of response regulators has evolved an array of diverse regulatory sensitivities. Although far from complete, the structural studies and comparative sequence analyses on VpsT identified several conserved structural features (α_6 -dimerization, ligand-coordinating residues, conserved amino acids connecting the canonical phosphoreceiver site with the nucleotide-binding pocket), which could shed light on the evolution from archetypal two-component phosphotransfer to small molecule recognition and facilitate the prediction of additional second messenger targets.

VpsR of *V. cholerae* and FleQ of *P. aeruginosa*: AAA+ domains as putative c-di-GMP sensors. VpsR of *V. cholerae* is another c-di-GMP-responsive positive regulator of biofilm formation and rugosity, which functions in concert and shares an overall similar regulon with VpsT.^{89,90,102} Interestingly, VpsR is homologous to one of the first identified protein sensors for c-di-GMP, FleQ of *P. aeruginosa*.³⁸ The two proteins contain a relatively divergent N-terminal response receiver domain, followed by a more conserved AAA+ σ^{54} -interaction domain and a C-terminal helix-turn-helix motif.

In FleQ, the N-terminal receiver domain has been proposed to participate in regulatory protein-protein interactions, such as binding of the active ATPase FleN, which is also a known antagonist of FleQ function.^{38,103} Little is known of the structural requirements for c-di-GMP recognition, other than that it occurs independently of the N-terminal receiver domain or ATP hydrolysis. This may suggest that rather than acting as a P-loop NTPase for energy-dependent transcription regulation, the AAA+ domain of FleQ (and possibly VpsR) has evolved an altered nucleotide specificity for c-di-GMP sensing.

When bound to DNA, FleQ acts as a master activator of flagellar gene expression, as well as a repressor for the *pel* exopolysaccharide synthesis operon.^{38,104} C-di-GMP recognition serves as an “off” switch for DNA binding and thus leads to motility inhibition and biofilm formation. Interestingly, c-di-GMP binding to the protein has been shown to have a much greater effect on promoters that FleQ represses (e.g. of the *pel* operon), compared to those it activates (e.g. of flagellar biosynthesis genes).³⁸ The main difference between the two types of promoters is the requirement for the alternative RNA polymerase sigma factor σ^{54} during FleQ-mediated transcription activation, which likely interacts directly with the nucleotide-sensing module.³⁸

CRP-like proteins (Clp): Ditching cAMP sensitivity for c-di-GMP. Clp proteins, homologs of the

cAMP receptor protein (CRP, also known as catabolite activator protein) of *E. coli*, have now been identified as c-di-GMP responsive transcription regulators in several species.^{40,43,87,105}

In *Xanthomonas* sp., Clp regulates approximately 300 genes involved in plant pathogenesis. Interestingly, *clp* deletion mutants show markedly decreased virulence potential, similarly to mutants lacking one or more c-di-GMP-specific phosphodiesterases.^{106,107} This, together with the fact that Clp can bind promoter DNA with high affinity in the absence of any effector, lead to the hypothesis that Clp-dependent virulence gene expression can be inhibited by direct c-di-GMP recognition.⁴³ Such a regulatory mechanism was subsequently validated in at least two different *Xanthomonas* species, and high-resolution structural data for the nucleotide-free Clp of *X. campestris* (XcClp) is now available.^{40,43,87}

XcClp consists of an N-terminal cNMP-binding domain and a C-terminal, HTH-based DNA-binding domain, linked by a dimerization module consisting of a long α -helix. The protein adopts a symmetrical dimeric structure, roughly similar to the structure of *E. coli* CRP in its *EcCRP*-cAMP-DNA complex.^{40,108} Specifically, the structure of ligand-free XcClp is characterized by a decreased gap between the N- and C-terminal domains (“closed” form), defined as intrinsically active for DNA recognition. Detailed structural analysis indicated that regardless of the high sequence identity between XcClp and *EcCRP* (44%), differences of key residues in at least three regions are responsible for a constitutive DNA-binding conformation and altered nucleotide specificity of XcClp.⁴⁰

First, residue S84 of *EcCRP*, which alone positions cAMP through five direct or indirect H-bonds in the *EcCRP*-cAMP complex, is substituted by a larger and negatively charged glutamate (E99), whose side chain is flipped away from the ligand-binding site. This single serine-to-glutamate substitution appears sufficient to render the protein insensitive to cAMP and has been even proposed to serve as a sole marker for cAMP- versus c-di-GMP-ligand specificity.^{40,109,110}

Two additional regions in XcClp were shown to contain key amino acid substitutions, likely responsible for the constitutive DNA-binding activation. These include the gap region between the N- and C-terminal domains, where a single glutamine (Q175 in *EcClp*) to arginine (R195 in XcClp) substitution serves to reduce the gap dimensions, as well as several amino acid substitutions, which are proximal to the DNA-binding motif and known to result in cAMP-independent transcription when introduced in *EcClp*.^{40,111}

Although no direct structural data are available for a XcClp-c-di-GMP complex, molecular modeling

and mutational analysis identified a putative ligand-binding site on the protein.⁴⁰ In the docked model, c-di-GMP is bound as a single molecule with its purine bases oriented parallel to each other. Similarly to other c-di-GMP binding motifs, the nucleotide is stabilized mainly by side-chain interactions with aspartate (D70, D170) and arginine (R150, R154) residues, including hydrogen bonds, salt bridges and π - π stacking interactions.⁴⁰ The binding pocket in XcClp is distinct from the one for cAMP recognition in EcCRP but is nevertheless found in a region known to strongly affect the DNA-binding strength of the latter.¹¹¹

The structural and functional analyses on XcClp indicate that the protein has evolved an altered nucleotide- and DNA-binding propensity, where c-di-GMP recognition serves to block constitutive DNA binding and rapidly switch off virulence gene expression.

C-di-GMP dependent RNA processing: A role for *E. coli* polynucleotide phosphorylase (PNPase). The role of c-di-GMP in regulating RNA transactions was initially highlighted by the discovery of c-di-GMP-sensitive riboswitches in messenger RNAs of various species (for review see Ref. ¹¹²). In addition, a recent study showed that c-di-GMP can also regulate RNA turnover by direct binding to PNPase, an important 3' to 5' exonuclease of *E. coli*.¹¹³ The enzyme was found to associate in a megadalton-sized RNA “degradosome” complex with two direct O₂ sensors, the diguanylate cyclase DosC and the c-di-GMP-specific phosphodiesterase DosP. The two enzymes are activated in anaerobic and aerobic conditions, respectively, offering a spatially restricted mechanism for regulation of available c-di-GMP levels and PNPase-dependent RNA processing in response to environmental oxygen stimuli. While high-resolution structural data for the c-di-GMP-free PNPase are already available in the literature and the PDB database,⁵⁰ further structure-function analyses are necessary to explain its c-di-GMP-dependent mode of regulation.

Eukaryotic c-di-GMP receptors

Although c-di-GMP appears to have ubiquitous effects on the survival and virulence strategies adopted by most bacterial species, enzymes for its synthesis or degradation are not found in any other domain of life. In the context of infectious diseases, the presence of c-di-GMP would thus serve as a unique beacon for microbial pathogenesis and recent findings suggest that the mammalian immune system has developed dedicated machinery for c-di-GMP recognition and subsequent pathogen elimination.

It is important to note that c-di-GMP's immunostimulatory effects in mammals also hold promise for the development of novel adjuvants, vaccines, or

other immunotherapeutics. Indeed, one early report demonstrated that exogenous c-di-GMP can inhibit the growth of human colon cancer cells without cytotoxic effects against non-cancerous cells in culture.¹¹⁴ More recently, studies from two different groups indicated that pretreatment of mice with c-di-GMP prior to challenge with *K. pneumoniae* or *Staphylococcus aureus* resulted in suppressed pathogen burdens,^{115,116} and that the immune response in mice was enhanced when an intranasal influenza vaccine was supplemented with c-di-GMP.¹¹⁷ Deciphering the mechanisms of c-di-GMP recognition and signaling effects in mammals can thus prove instrumental for the prevention and treatment of a variety of human diseases, which further underscores the need for diligent structure–function analyses.

STING: A mammalian receptor for c-di-GMP.

The presence of c-di-GMP is known to elicit host Type I interferon response, which is typically induced by viral and bacterial pathogens and contributes to diverse outcomes toward successful pathogen elimination *in vivo*.^{118,119} Interestingly, while the transcription response to c-di-GMP in mammalian cells is virtually identical to the one triggered by cytosolic presence of pathogen DNA, c-di-GMP recognition was found to be independent from known DNA or RNA cytosolic sensors.¹¹⁸

These puzzling effects were recently resolved by the discovery that STING (Stimulator of Interferon Genes) serving as an essential signaling adaptor between cytosolic DNA detection and interferon gene induction, can moonlight as a direct target for c-di-GMP stimulation.¹²⁰ Bioinformatic analyses indicate that STING is a transmembrane protein with several membrane-spanning segments at the N-terminus and a globular carboxy-terminal domain. Mutational analysis confirmed that residues important for c-di-GMP binding by the C-terminal domain are separate from those important for bacterial DNA recognition. Interestingly, competitive binding studies indicated that STING is also a likely sensor for another RNA dinucleotide established as a prokaryote-specific second messenger, c-di-AMP.

In terms of structural classification, STING does not share homology with other immunosensory proteins in mammals and thus appears as a stand-alone representative of a novel class of c-di-GMP receptors. Surprisingly, part of the protein is homologous to the diadenylate cyclase Dac1 of *Listeria monocytogenes*, an intracellular pathogen known to elicit STING-dependent Type I interferon response through efflux of cyclic dinucleotides.^{120–122} This raises the interesting hypothesis that the mammalian immune system has hijacked bacterial c-di-NMP signaling elements to evolve an effective and highly specific system for pathogen recognition.

Apart from STING, initial binding studies have pointed toward several additional mammalian proteins as putative c-di-GMP receptors. Examples include p21/Ras small GTPases, cytoskeleton remodeling protein coronin 1A, RNA-capping and processing proteins RNMT and cyclophilin H, and others.^{123,124} Although potentially quite interesting, these candidates will require further structural and functional studies to confirm c-di-GMP binding specificity and physiologically relevant effects and to distinguish specific binding from potentially promiscuous affinity for RNA-based ligands.

Outlook

According to recent World Health Organization reports bacterial infections causing gastroenteric diseases, obstructive pulmonary conditions, and AIDS-derived opportunistic infections comprise some of the leading mortality causes worldwide. Given the long battle that medicine has fought against virulent microbes, such a grim outlook inevitably raises concerns about the effectiveness of conventional prevention and treatment, and underscores the pressing need for alternative therapeutic approaches.

Traditional antibiotics are designed to kill susceptible bacteria and thus select for naturally occurring drug-resistant forms. This selective enhancement of resistant bacteria is especially rampant in multiple or prolonged exposures, often associated with hospital settings or chronic diseases. Mechanistic studies of nonessential virulence determinants, such as enzymes and protein sensors involved in c-di-GMP signal transduction and biofilm formation, could thus provide targets for the design of novel anti-infectives that are aimed at jamming bacterial communication and dispersal of the pathogenic crowds without associated risks of evolutionary pressure and resistance development. We highlighted here recent advances in the structure–function studies on such important c-di-GMP signal transduction proteins highlighting the functional complexity and idiosyncracies of individual targets. It is important to note, that the general lack of redundancy between c-di-GMP signaling mechanisms across or even within species holds the additional promise for selective targeting of parasitic and virulent microbes while preserving beneficial symbiotic bacteria. It remains to be seen whether the rapidly developing field of structural biology dedicated to c-di-GMP signaling research will be able to provide the necessary molecular blueprints and mechanistic insight for the design of such therapeutics.

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