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Interactions of the CpxA sensor kinase and cognate CpxR response regulator from *Yersinia pseudotuberculosis*

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Abstract

Background: The CpxA sensor kinase-CpxR response regulator two-component regulatory system is a sentinel of bacterial envelope integrity. Integrating diverse signals, it can alter the expression of a wide array of components that serve to shield the envelope from damage and to promote bacterial survival. In bacterial pathogens such as *Yersinia pseudotuberculosis*, this also extends to pathogenesis. CpxR is thought to dimerize upon phosphorylation by the sensor kinase CpxA. This phosphorylation enables CpxR binding to specific DNA sequences where it acts on gene transcription. As Cpx pathway activation is dependent on protein-protein interactions, we performed an interaction analysis of CpxR and CpxA from *Y. pseudotuberculosis*.

Results: CpxR full-length and truncated versions that either contained or lacked a putative internal linker were all assessed for their ability to homodimerize and interact with CpxA. Using an adenylate cyclase-based bacterial two hybrid approach, full-length CpxR readily engaged with CpxA. The CpxR N-terminus could also homodimerize with itself and with a full-length CpxR. A second homodimerization assay based upon the λ CI repressor also demonstrated that the CpxR C-terminus could homodimerize. While the linker was not specifically required, it enhanced CpxR homodimerization. Mutagenesis of *cpxR* identified the aspartate at residue 51, putative N-terminal coiled-coil and C-terminal winged-helix-turn-helix domains as mediators of CpxR homodimerization. Scrutiny of CpxA full-length and truncated versions revealed that dimerization involved the N-terminus and an internal dimerization and histidine phosphotransfer domain.

Conclusions: This interaction analysis mapped regions of CpxR and CpxA that were responsible for interactions with self or with each other. When combined with other physiological and biochemical tests both hybrid-based assays can be useful in dissecting molecular contacts that may underpin Cpx pathway activation and repression.

Keywords: BACTH assay, λ CI homodimerization assay, Homodimer, Heterodimer, Linker, Coiled-coil, Winged helix-turn-helix, Phosphorylation

Background

Conditions that threaten the integrity of the bacterial envelope are collectively termed extracytoplasmic stresses (ECS). Bacteria employ a series of ECS-responsive regulatory pathways to control the expression of 'survival' genes whose products act in the periplasm to maintain membrane integrity. This ensures continued bacterial growth even in environments poisoned by harmful ECS. A notable

ECS-responsive pathway is the CpxA-CpxR two-component regulatory system (TCRS) [1,2]. Belonging to the class I histidine kinases [3], CpxA is the integral inner-membrane sensor kinase (SK). Upon activation by ECS sensing – presumably in the form of misfolded proteins – it becomes auto-phosphorylated. Through a phosphotransfer reaction, CpxA transduces this signal through the membrane to activate the cytoplasmic CpxR response regulator (RR). CpxR belongs to the OmpR/PhoB family of winged-helix-turn-helix (wHTH) transcriptional RRs [4]. Phosphorylated CpxR (CpxR ~ P) then binds to the promoters of genes coding for several protein folding and degradation factors that operate in the periplasm. An important

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function of the Cpx pathway is therefore in protein quality control in the bacterial envelope with an emphasis on maintaining outer membrane structural integrity [1,2]. However, as the CpxR regulon may include hundreds of genes [5,6], playing sentinel to the cell envelope must incorporate diverse signals and safeguarding mechanisms.

In Gram negative enteropathogenic *Yersinia pseudotuberculosis*, activation of the Cpx pathway also regulates periplasmic protein folding and degradation factors [7-10]. Additionally, accumulation of CpxR~P down-regulates several prominent *Yersinia* virulence determinants. Most notable is the Ysc-Yop type III secretion system [7,10], the cellular adhesin known as invasins and its transcriptional activator RovA [8,9]. RovA is a global regulator in pathogenic *Yersinia* and is capable of influencing the expression of at least 60 genes [11-13]. Hence, the regulatory influence of CpxR~P – either direct or indirect via effects on *rovA* expression – has potential to be quite widespread in these bacteria.

An activated Cpx pathway may therefore function to restrict *Y. pseudotuberculosis* virulence factor expression during times of ECS when all resources must be dedicated to the expression of survival genes. In the absence of ECS, virulence factor expression can be de-repressed to permit *Y. pseudotuberculosis* to mount a successful host infection [14]. In fact, accumulating evidence in a few Gram-negative pathogens suggests that the Cpx pathway might possess a universal role in virulence factor regulation and bacterial pathogen fitness either by aiding in the establishment of an environmental reservoir or during host infections [2,15-17].

CpxA possesses a modular structure [3,18,19] (Figure 1). Consecutive domains in their N-terminus are responsible for signal input, processing and signal transmission. The latter is defined as the HAMP linker domain by virtue of its presence in Histidine kinases, Adenyl cyclases, Methyl-accepting proteins and Phosphatases [20]. Located in the cytoplasm, the HAMP linker domain most likely regulates the phosphorylation of histidine SKs by transmitting conformational changes originating in the periplasmic ligand-binding domains to the cytoplasmic-located C-terminal kinase catalytic domain [20,21]. The remainder of the protein is composed of an internal dimerization and histidine phosphotransfer (DHP) domain and then a C-terminal histidine kinase catalytic domain [22,23]. Together, these two consecutive domains form the kinase core. The DHP domain mediates dimerization [24], and possesses the conserved phospho-accepting histidine residue and also a phosphatase domain for dephosphorylating CpxR [22,23,25,26]. Lastly, the catalytic domain is essential for kinase activity. It contains a number of conserved motifs necessary for ATP binding [27,28] and probably also for catalysis and phosphotransfer [22].

CpxR contains two structurally conserved domains; a N-terminal receiver domain and a C-terminal effector domain that are joined by a flexible internal linker [29,30] (Figure 2). The N-terminal receiver domains of RRs share several conserved features such as a α_4 - β_5 - α_5 interface and an enrichment in aspartate/serine/threonine amino acids. These residues are thought to cooperate in dimerization as a result of phosphorylation induced conformational changes that in turn propagates the signal [29,30]. A

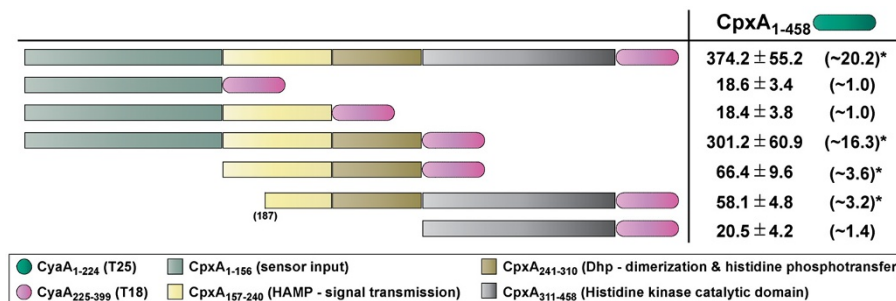


Figure 1 BACTH analysis of CpxA-CpxA interactions. Full-length CpxA₁₋₄₅₈ was translationally fused to the N-terminus of CyaA₁₋₂₂₄ (T25 – dark green shade) creating a CpxA₁₋₄₅₈-T25 hybrid used as the ‘bait’. Full-length CpxA₁₋₄₅₈ was also translationally fused to the N-terminus of CyaA₂₂₅₋₃₉₉ (T18 – magenta shade) giving rise to a ‘prey’ CpxA₁₋₄₅₈-T18 hybrid. Based on CpxA divisions into sensor input (CpxA₁₋₁₅₆, cadet blue shade), HAMP – signal transmission (CpxA₁₅₇₋₂₄₀, soft yellow shade), Dhp – dimerization and histidine phosphorylation (CpxA₂₄₁₋₃₁₀, metallic gold shade) and histidine kinase catalysis (CpxA₃₁₁₋₄₅₈, grey shade) domains, an additional six ‘prey’ CpxA-T18 hybrids were constructed; CpxA₁₋₁₅₆-T18, CpxA₁₋₂₄₀-T18, CpxA₁₋₃₁₀-T18, CpxA₁₅₇₋₃₁₀-T18, CpxA₁₈₇₋₄₅₈-T18 and CpxA₃₁₁₋₄₅₈-T18. BACTH interaction analysis of ‘bait’ and ‘prey’ hybrids was quantified via measurement of β-galactosidase activity and is represented as units/mg dry weight of host *E. coli* BTH101 bacteria (left column; black font). As an internal positive control, we used the provided constructs expressing T18-Zip and T25-Zip that yielded 1547.0 ± 121.2 units of β-galactosidase activity /mg dry weight of bacteria. This was equivalent to ~83.8 fold more enzymatic activity produced compared to bacteria co-expressing only T18 and T25 (18.5 ± 3.9 units of β-galactosidase activity). The fold change in enzymatic activity caused by CpxA-CpxA interactions relative to this negative control is indicated in parentheses to the right. A level of β-galactosidase activity at least 3-fold higher than the negative control was considered to indicate a positive interaction (*). Data is presented as the mean (± standard error of the mean) of at least four independent experiments performed in triplicate.

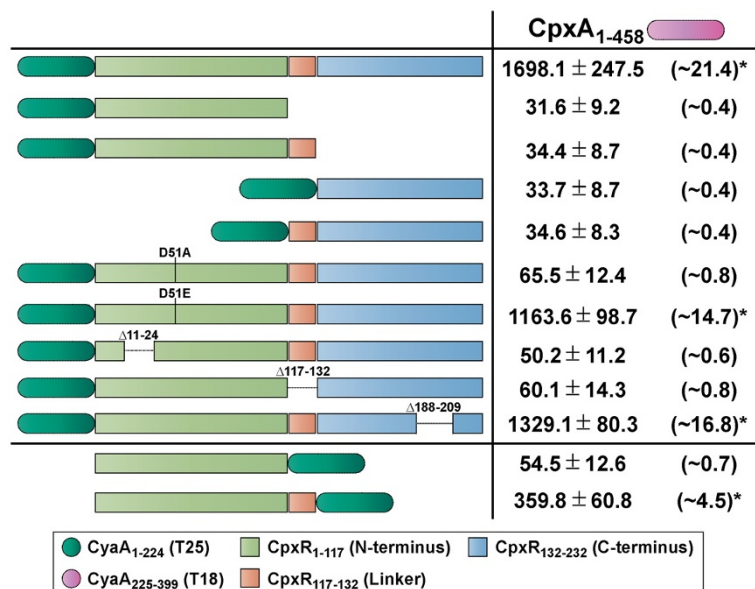


Figure 2 Regions of CpxR interacting with CpxA as monitored by BACTH analysis. Full-length CpxA₁₋₄₅₈ was translationally fused to the N-terminus of CyaA₂₂₅₋₃₉₉ (T18 – magenta shade) giving rise to a ‘bait’ CpxA₁₋₄₅₈-T18 hybrid. Full-length CpxR₁₋₂₃₂ was translationally fused to the C-terminus of CyaA₁₋₂₂₄ (T25 – dark green shade) giving rise to a T25-CpxR₁₋₂₃₂ ‘prey’ hybrid. CpxR was also divided into N-terminal (pistachio green), internal linker (orange) and C-terminal (sky blue) domains. Based on these divisions, additional ‘prey’ T25 hybrids were similarly constructed that consisted of only the N-terminus without linker (T25-CpxR₁₋₁₁₇) or with linker (T25-CpxR₁₋₁₃₂) and the C-terminus without linker (T25-CpxR₁₃₂₋₂₃₂) or with linker (T25-CpxR₁₇₋₂₃₂). CpxR₁₋₁₃₂ was also fused to the N-terminus of T25 giving the CpxR₁₋₁₃₂-T25 hybrid. Full-length or near full-length CpxR ‘prey’ T25 constructs were also generated consisting of T25-CpxR_{D51A} or T25-CpxR_{D51E} lacking the phosphorylated aspartate residue as well as T25-CpxR_{Δ11-24}, T25-CpxR_{Δ117-132} and T25-CpxR_{Δ188-209} lacking the putative N-terminal coiled-coil, internal linker and C-terminal winged helix-turn-helix regions respectively. BACTH interaction analysis of ‘bait’ and ‘prey’ hybrids was quantified via measurement of β-galactosidase activity and is represented as units/mg dry weight of host *E. coli* BTH101 bacteria (left column; black font). The internal positive control based upon constructs expressing T18-Zip and T25-Zip yielded 1603.6 ± 146.4 units of β-galactosidase activity/mg dry weight of bacteria. This was equivalent to ~20.2 fold more enzymatic activity produced compared to bacteria co-expressing only T18 and T25 (79.2 ± 6.2 units of β-galactosidase activity). The fold change in enzymatic activity caused by CpxA-CpxR interactions relative to this negative control is indicated in parentheses to the right. The asterisks (*) indicates a positive interaction. Data is presented as the mean (± standard error of the mean) of at least four independent experiments performed in triplicate.

common wHTH motif defines the C-terminal effector domain. This motif is responsible for binding to target DNA in order to regulate transcription output [29,30]. The internal linker region that tethers together the receiver and effector domain might as well contribute in signal propagation, perhaps by promoting necessary molecular interactions [29,30].

Significantly, TCRs are a potential target for the development of novel anti-bacterial therapeutics [31], which are urgently needed to overcome the current public health crisis associated with common antibiotic resistances among prominent bacterial pathogens [32]. Given the emerging theme of CpxA-CpxR-dependent virulence gene regulation in clinically important bacteria [2,15-17], this pathway might well be an attractive target for chemical modulation. With this in mind, the present study is primarily aimed at establishing practical and reliable assays to identify the molecular interactions of CpxA and CpxR from pathogenic *Y. pseudotuberculosis*. Not only would this guide further *in vivo* mutagenesis studies to explore

the molecular determinants involved in CpxA-CpxR signal transduction and virulence gene control in *Y. pseudotuberculosis*, this knowledge should also benefit studies aimed at targeting the Cpx pathway using chemical manipulation.

Methods

Bacterial strains, plasmids and growth conditions

Bacterial strains and plasmids are listed in Supplementary Table S1 and available for online download as Additional file 1. Unless indicated, all bacteria were routinely cultured in Luria-Bertani (LB) broth or agar at 37°C for *E. coli* or 26°C for *Y. pseudotuberculosis*. When required, antibiotics at the following final concentrations were used: ampicillin (Ap; 100 µg/ml), kanamycin (Km; 25 µg/ml), chloramphenicol (Cm; 25 µg/ml), tetracycline (Tc; 10 µg/ml).

Molecular methods

DNA fragments were PCR amplified from YPIII/pIB102 (parental *Y. pseudotuberculosis*) using Easy-A High

Fidelity PCR cloning enzyme (Agilent technologies, Santa Clara, California, USA) and the oligonucleotides pairs synthesized by Sigma-Aldrich Sweden AB (Stockholm, Sweden) and are listed in Supplementary Table S2 (available for online download as Additional file 2). Amplified DNA fragments were cloned into pTZ57T/R (Fermentas, Vilnius, Lithuania) and correct sequence were verified by Eurofins MWG Operon (Ebersberg, Germany). Confirmed sequences were then re-cloned as a translational fusion into the XbaI/EcoRI site of the pKT25, pKNT25, pUT18, or pUT18C vectors for BACTH analysis or the BglII/KpnI site of the pKWY2428 vector for λ cl homodimerization analysis. Routine maintenance of individual clones was performed in *E. coli* DH5.

BACTH and the β -galactosidase assay

Pairs of BACTH vectors expressing CyaA T18 and T25 fusions to 'bait' and 'prey' proteins were sequentially transformed into *E. coli* BTH101 using the chemical transformation method [33]. Transformants were selected at 37°C by overnight growth on LB agar with appropriate antibiotics and supplemented with 0.5 mM isopropyl- β -d-thiogalactopyranoside (IPTG) and 40 μ g/ml 5-bromo-4-chloro-3-indolyl- β -d-galactopyranoside (X-Gal). Well isolated single colonies were then selected for inoculation of bacterial cultures that were grown overnight at 30°C in LB broth supplemented appropriate antibiotics. Then 0.025 volumes were sub-cultured in the same fresh media with 0.5 mM IPTG and grown for a further 2 h at 30°C. The remainder of the β -galactosidase assay followed descriptions outlined in the BACTH manual and associated literature (Euromedex, Souffelweyersheim, France) [34]. β -galactosidase activity was represented in Units/mg dry weight bacteria according to the manufacturer's direction. Data are a representative of at least four independent experiments performed in triplicate. Guided by previous analyses [35], we considered a positive interaction only if the β -galactosidase activity level was at least three-fold higher than that measured for the negative control plasmids (expressing T18 and T25 alone).

For western blotting, similarly grown cultures were harvested after late-logarithmic growth. Pellets were solubilized in loading buffer (50 mM Tris-HCl pH 6.8, 2% SDS, 10% Glycerol, 1% β -Mercaptoethanol and 0.02% Bromophenol blue) and fractionated by SDS-PAGE with 12% acrylamide. Protein was then transferred to Immobilon[®]-P PDVF transfer membrane (Millipore Corporation) using a Hoefer semi-dry transfer apparatus (GE Healthcare). Membranes were exposed to rabbit polyclonal antibodies that were a gift from Thomas Silhavy (anti-CpxA), produced by Agrisera AB (Vännäs, Sweden) from purified antigen (anti-CpxR) or purchased from Santa Cruz Biotechnology Inc [(anti-CyaA (b-300)]. These were then detected with an anti-rabbit monoclonal

antibody conjugated with horse radish peroxidase (GE Healthcare) and Pierce ECL 2 western blotting substrate (Thermo Scientific).

λ cl Homodimerization assays

As performed previously [36], assays were carried-out in the *E. coli* JM109 background. In brief, freshly transformed bacteria were grown overnight at 37°C in LB broth with chloramphenicol selection in the presence of 0.1 mM IPTG. Next, 0.01 volumes were sub-cultured in fresh media and grown to mid-logarithmic phase of growth. All cultures were standardized according to an optical density of 600 nm (OD₆₀₀) and 5 μ l suspensions of 10-fold serial dilutions were spotted onto LB agar with chloramphenicol, tetracycline and 0.1 mM IPTG. The ability to grow in the presence of tetracycline was scored following overnight incubation at 37°C for at least four independent experiments.

Additionally, λ cl-CpxR fusions contained within these pelleted bacterial *E. coli* cells were fractionated by SDS-PAGE with 12% acrylamide. Protein was then transferred to Protran[®] nitrocellulose transfer membrane (Whatman GmbH) using a Hoefer semi-dry transfer assembly (GE Healthcare). Proteins of interest were bound with specific rabbit polyclonal anti-CpxR antibodies and then detected with an anti-rabbit monoclonal antibody conjugated with horse radish peroxidase and a homemade chemiluminescent solution. As a loading control, we utilized specific rabbit polyclonal anti-CAT antibodies (Sigma-Aldrich) to detect levels of chloramphenicol acetyltransferase encoded by the *cat* gene located on the vector pKWY2428.

Results

Dimerization of the CpxA sensor kinase using the BACTH assay

This study had the goal to identify a cost-effective, reliable and convenient protein-protein interaction assay for the characterization of regulatory deficient mutants of CpxA and CpxR from *Y. pseudotuberculosis*. The BACTH system – involving reconstitution of *Bordetella pertussis* adenylate cyclase (CyaA) T18 and T25 domains – is reported to be ideal for analyzing the interactions among membrane-anchored bacterial proteins in their natural environment [34,35], including diverse SKs [37-41]. Since CpxA is expected to dimerize [1,42], we first analyzed the ability of CpxA from *Y. pseudotuberculosis* to interact with itself in the BACTH assay. A series of constructs composed of N- and C-terminal T18 and T25 translational fusions with CpxA were established. Pairs of plasmids expressing a 'bait' and 'prey' hybrid were transformed into *E. coli* BTH101. This strain contains a *lacZ* reporter gene under CAP-cAMP control; an interaction between bait and prey will reconstitute CyaA activity leading to

accumulation of cAMP that, together with CAP, will stimulate transcription of the *lacZ* reporter gene that is indirectly measurable via an elevation in β -galactosidase activity. Serving as bait, full-length CpxA translationally fused to the N-terminus of T25 (CpxA₁₋₄₅₈-T25) could clearly bind to the prey hybrids CpxA₁₋₄₅₈-T18 and C-terminal truncated CpxA₁₋₃₁₀-T18, as a result producing β -galactosidase activity that was ~20.2 and ~16.3 fold higher than the negative control (Figure 1). Modest increases in β -galactosidase activity of ~3.6 and ~3.2 fold were also indicative of a slight interaction between bait and the respective prey hybrids CpxA₁₅₇₋₃₁₀-T18 and CpxA₁₈₇₋₄₅₈-T18 (Figure 1). Since the DHP domain (residues 241–310) is a common feature of all four interacting hybrids, these data demonstrate its importance to CpxA homodimerization. Corroborating this, the remaining three hybrids (CpxA₁₋₁₅₆-T18, CpxA₁₋₂₄₀-T18 and CpxA₃₁₁₋₄₅₈-T18) all lacking the DHP domain produced low reporter activity indicative of poor homodimerization capacity (Figure 1). Although CpxA homodimerization was easily measurable via BACTH analysis, production of β -galactosidase activity was notably lesser than that achieved by a strain having the T25-Zip/T18-Zip positive control plasmids, where accumulated activity was ~83.8 fold more than background (data not shown).

It is possible that low reporter activity might also be due to unstable fusion product. To investigate this, we made numerous attempts to measure protein levels in each fusion bearing strain using western blotting with either anti-CyaA (b-300) or anti-CpxA antibodies. However, neither antibody was successful in detecting accumulated steady state levels of any specific T25 or T18 fusion, even for those fusion-bearing strains that displayed high levels of reporter activity and the positive control (data not shown). Hence, for those three strains where low reporter activity was recorded, we cannot definitively conclude that this was due to genuinely poor interactions.

An N-terminal region of CpxR interacts with CpxA

Our next goal was to examine if the BACTH assay could support CpxR-CpxA interaction studies. In this analysis, we used the full-length CpxA₁₋₄₅₈-T18 hybrid as bait that in Figure 1 resulted in high reporter activity and must therefore be stably produced. A corresponding prey construct that is composed of T25 C-terminally fused to full-length CpxR (T25-CpxR₁₋₂₃₂) was co-transformed into *E. coli* BTH101. An interaction between bait and prey yielded β -galactosidase activity that was ~21.4 fold higher than the negative control and equivalent to the positive control (~20.2 fold increase) (Figure 2, data not shown).

To identify the region of CpxR engaging with CpxA, we were guided by a recent study by Tapparel and

colleagues [43]. It was suggested that the C-terminal domain effector activity of *E. coli* CpxR can become constitutive by removing the N-terminal regulatory domain appended via an internal flexible linker. We therefore established a series of prey constructs with T25 that were C-terminally fused to truncated CpxR derivatives composed of only the N-terminus without internal linker (T25-CpxR₁₋₁₁₇) or with internal linker (T25-CpxR₁₋₁₃₂) and the C-terminus without internal linker (T25-CpxR₁₃₂₋₂₃₂) or with internal linker (T25-CpxR₁₁₇₋₂₃₂). No interaction between any of these prey hybrids and the CpxA₁₋₄₅₈-T18 bait was observed (Figure 2). Attempts to detect any accumulated T25-CpxR variant in cell lysates by western blotting with either anti-CyaA (b-300) or anti-CpxR antibodies were again unsuccessful, even for the full-length CpxR fusion that did interact with CpxA (data not shown). On the other hand, altering the orientation of the T25 fusion by creating an alternative prey hybrid of CpxR₁₋₁₃₂-T25 did permit modest reporter activity (~4.5 fold) indicative of an interaction with the CpxA₁₋₄₅₈-T18 bait (Figure 2). Thus, this suggested that the CpxR N-terminus might be more important for CpxA binding.

To explore this further, we utilized a CpxR_{D51A} variant that is non-phosphorylatable and is therefore essentially inactive ('locked off') [9]. Additionally, another non-phosphorylatable variant, CpxR_{D51E}, was generated on the basis that it may cause a constitutively 'locked-on' CpxR phenotype, as is often the case for related RRs [44-46]. In parallel, we deleted *in silico* predicted structural elements of CpxR including the putative N-terminal-located coiled-coil domain (CpxR _{Δ 11-24}), a C-terminal-located wHTH domain (CpxR _{Δ 188-209}) as well as the flexible internal linker that separates the N- and C-terminal CpxR domains (CpxR _{Δ 117-132}) [43]. Again, the same CpxA₁₋₄₅₈-T18 bait construct was used in combination with the newly generated prey constructs T25-CpxR_{D51A}, T25-CpxR_{D51E}, T25-CpxR _{Δ 11-24}, T25-CpxR _{Δ 117-132} and T25-CpxR _{Δ 188-209}. Enzymatic assays revealed that CpxR could interact well with CpxA, but only if its N-terminus remained intact, such as with the fully functional CpxR_{D51E} variant (~14.7 fold increase in reporter activity) or the CpxR _{Δ 188-209} variant (~16.8 fold) (Figure 2). Conversely, disruption of CpxR N-terminus, as occurred with the CpxR_{D51A}, CpxR _{Δ 11-24} and CpxR _{Δ 117-132} variants, only permitted low reporter activity in these fusion-bearing strains (Figure 2). Although western blotting could not determine whether any of these T25-CpxR fusions were actually produced and stable, these data corroborate earlier *in silico* predictions that the CpxR N-terminus is probably involved in crosstalk with CpxA [47,48]. They also suggest that BACTH is a valid approach to further investigate this aspect of Cpx pathway activation.

The extreme C-terminus of CpxA is dispensable for interactions with CpxR

A reciprocal experiment was next conducted in an effort to identify the region of CpxA necessary for engaging CpxR. We selected the T25-CpxR₁₋₂₃₂ construct as bait, since it induced high reporter activity in a strain co-expressing CpxA₁₋₄₅₈-T18 (see Figure 2). We also used the same prey constructs incorporating the CpxA truncated variants fused to the N-terminus of T18 and described in Figure 1. *E. coli* BTH101 expressing both bait T25-CpxR₁₋₂₃₂ and prey CpxA₁₋₄₅₈-T18 hybrids produced β-galactosidase levels that were ~24.2 fold higher than the negative control, and only slightly lower than the T25-Zip/T18-Zip positive control (~32.8 fold elevation) (Figure 3, data not shown). The CpxA₁₋₃₁₀-T18 hybrid could also interact with bait T25-CpxR₁₋₂₃₂ as indicated by ~12.1 fold higher reporter activity in the host strain (Figure 3). No remaining fusion combinations induced any significant reporter activity in BTH101. Once again, we could not confirm by western blotting whether this negligible reporter activity was due to poor interactions or unstable CpxA₁₋₁₅₆-T18, CpxA₁₋₂₄₀-T18, CpxA₁₅₇₋₃₁₀-T18, CpxA₁₈₇₋₄₅₈-T18 and CpxA₃₁₁₋₄₅₈-T18 protein (data not shown). However, since the CpxA₁₅₇₋₃₁₀-T18 and CpxA₁₈₇₋₄₅₈-T18 fusions must at least be modestly produced (see Figure 1), this data suggests that the DHP domain in cooperation with additional upstream N-terminal CpxA sequence might be sufficient for engaging CpxR, corroborating reports focused on other SK-RR cognate pairs [47-49]. It also suggests that

sequence in the extreme C-terminus of CpxA is probably not required for interacting with CpxR, at least via the BACTH assay.

BACTH analysis of CpxR-CpxR interactions

Next, we explored the suitability of the BACTH assay for dissecting CpxR dimerization [50]. In the first instance, a 'bait' construct consisting of full-length CpxR fused to the C-terminus of T25 (T25-CpxR₁₋₂₃₂) was transformed with a 'prey' construct of full-length CpxR fused to the N-terminus of T18 (CpxR₁₋₂₃₂-T18). However, the detected β-galactosidase activity barely increased above the negative control value (Figure 4). As with all our other BACTH analysis, neither fusion was detected by western blotting (data not shown) making it difficult to conclude whether the low reporter activity represents the absence of an interaction or merely unstable protein. Curiously, co-expression of full-length CpxR fusions still did not stimulate transcription of the *lacZ* reporter even after the orientation of the T18 and T25 fusions were swapped (N-terminus vs C-terminus) (data not shown).

Given a report that the C-terminal domain effector activity of *E. coli* CpxR can be unlocked by physically removing the N-terminal regulatory domain [43], we entertained the notion that our full-length CpxR fusions might be locked in a static structural confirmation that prevents measurable dimerization. To investigate this, we employed a series of prey constructs with T18 N-terminally fused to truncated CpxR derivatives composed

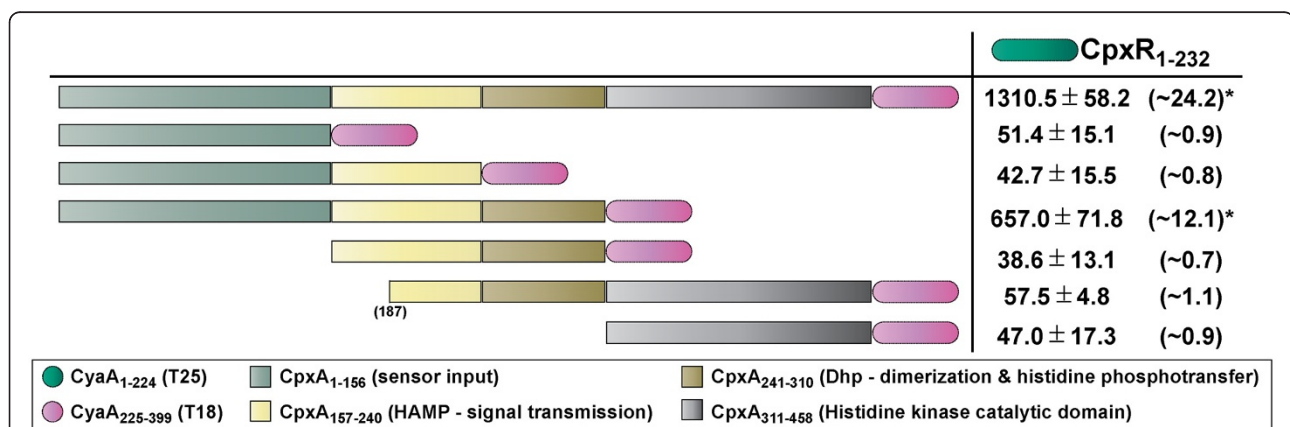


Figure 3 Regions of CpxA interacting with CpxR as monitored by BACTH analysis. Full-length CpxR₁₋₂₃₂ was translationally fused to the C-terminus of CyaA₁₋₂₂₄ (T25 – dark green shade) creating a T25-CpxR₁₋₂₃₂ hybrid used as the 'bait'. As in Figure 1, the same seven CpxA variants translationally fused to the N-terminus of CyaA₂₂₅₋₃₉₉ (T18 – magenta shade) were used as 'prey' hybrids, that is CpxA₁₋₄₅₈-T18, CpxA₁₋₁₅₆-T18, CpxA₁₋₂₄₀-T18, CpxA₁₋₃₁₀-T18, CpxA₁₅₇₋₃₁₀-T18, CpxA₁₈₇₋₄₅₈-T18 and CpxA₃₁₁₋₄₅₈-T18. BACTH interaction analysis of 'bait' and 'prey' hybrids was quantified via measurement of β-galactosidase activity and is represented as units/mg dry weight of host *E. coli* BTH101 bacteria (left column; black font). The internal positive control based upon the constructs expressing T18-Zip and T25-Zip yielded 1778.1 ± 120.3 units of β-galactosidase activity/mg dry weight of bacteria. This was ~32.8 fold more enzymatic activity produced compared to bacteria co-expressing only T18 and T25 (54.2 ± 9.0 units of β-galactosidase activity). The fold change in enzymatic activity caused by CpxR-CpxA interactions relative to this negative control is indicated in parentheses to the right. The asterisks (*) indicates a positive interaction. Data is presented as the mean (± standard error of the mean) of at least four independent experiments performed in triplicate.

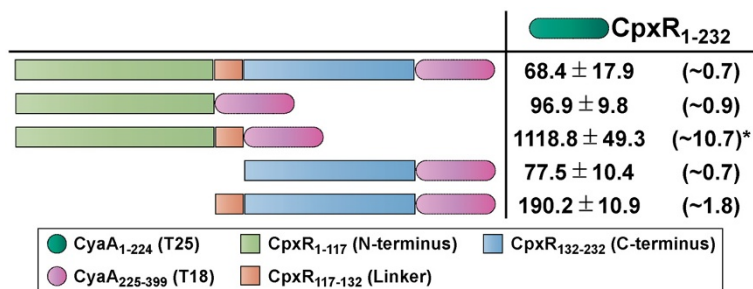


Figure 4 BACTH analysis of CpxR-CpxR interactions. Full-length CpxR₁₋₂₃₂ was translationally fused to the C-terminus of CyaA₁₋₂₂₄ (T25 – dark green shade) creating a T25-CpxR₁₋₂₃₂ hybrid used as the ‘bait’. Full-length CpxR₁₋₂₃₂ was translationally fused to the N-terminus of CyaA₂₂₅₋₃₉₉ (T18 – magenta shade) giving rise to a CpxR₁₋₂₃₂-T18 ‘prey’ hybrid. Based upon divisions of CpxR into N-terminal (pistachio green), internal linker (orange) and C-terminal (sky blue) domains, additional ‘prey’ T18 hybrids were constructed that consisted of only the N-terminus without linker (CpxR₁₋₁₁₇-T18) or with linker (CpxR₁₋₁₃₂-T18) and the C-terminus without linker (CpxR₁₃₂₋₂₃₂-T18) or with linker (CpxR₁₁₇₋₂₃₂-T18). BACTH interaction analysis of ‘bait’ and ‘prey’ hybrids was quantified via measurement of β-galactosidase activity and is represented as units/mg dry weight of host *E. coli* BTH101 bacteria (left column; black font). The internal positive control based upon the constructs expressing T18-Zip and T25-Zip yielded 1521.9 ± 150.6 units of β-galactosidase activity/mg dry weight of bacteria. This has ~14.6 fold more enzymatic activity than bacteria co-expressing only T18 and T25 (104.6 ± 12.9 units of β-galactosidase activity). The fold change in enzymatic activity caused by CpxR-CpxR interactions relative to this negative control is indicated in parentheses to the right. The asterisks (*) indicates a positive interaction. Data is presented as the mean (± standard error of the mean) of at least four independent experiments performed in triplicate.

of only the N-terminus without (CpxR₁₋₁₁₇-T18) or with internal linker (CpxR₁₋₁₃₂-T18) and the C-terminus without internal linker (CpxR₁₃₂₋₂₃₂-T18) or with internal linker (CpxR₁₁₇₋₂₃₂-T18). These constructs were related to those already described in Figure 2 and are also utilized in subsequent aspects of this study. These truncated CpxR-T18 prey constructs were transformed into *E. coli* BTH101 containing the full-length T25-CpxR₁₋₂₃₂ bait construct. On the basis of a ~10.7 fold elevation from baseline β-galactosidase activity, T25-CpxR₁₋₂₃₂ formed a strong interaction with CpxR₁₋₁₃₂-T18 (Figure 4). On this occasion, the strength of this union mirrored the well established dimerization of the Zip protein used as a control (~14.6 fold elevation from baseline β-galactosidase activity) (Figure 4, data not shown). In contrast, no other interaction of truncated CpxR with full-length CpxR could be detected, even after the orientation (N-terminus vs C-terminus) or combination (‘bait’ vs ‘prey’) of T18 and T25 domains were altered (Figure 4, data not shown). This also means that full-length CpxR-CpxR₁₋₁₃₂ interaction observed in the BACTH assay is conditional, being heavily influenced by the orientation and combination of the T18 and T25 fusions.

N-terminal mediated CpxR dimerization

Having the goal to better define internal interacting regions of CpxR, additional BACTH analyses were performed based upon the pair-wise transformation of *E. coli* BTH101 using CpxR₁₋₁₃₂-T18 as the bait with variable prey constructs of T25 N-terminally fused to truncated CpxR derivatives (CpxR₁₋₁₁₇-T25, CpxR₁₋₁₃₂-T25, CpxR₁₃₂₋₂₃₂-T25 and CpxR₁₁₇₋₂₃₂-T25). β-galactosidase levels were ~10.1 and ~12.9 fold higher in

bacteria co-expressing CpxR₁₋₁₃₂-T18/CpxR₁₋₁₁₇-T25 and CpxR₁₋₁₃₂-T18/CpxR₁₋₁₃₂-T25 bait/prey pairs respectively (Figure 5A). This compared to the positive control that yielded β-galactosidase levels that were ~17.3 fold higher. Hence, N-terminal CpxR with linker could interact with itself or with N-terminal CpxR lacking the linker. Significantly, it did not matter if the T18 and T25 domains were switched in either orientation (N-terminus vs C-terminus) or combination (bait vs prey), the β-galactosidase levels were still elevated (data not shown). These positive interactions indicate all CpxR N-terminal fusions are stably produced. To explore whether the internal linker contributes to the interactions of the CpxR N-terminus, parallel BACTH studies with the same prey constructs as used in Figure 5A were performed with a T18 bait construct fused at the N-terminus to truncated CpxR composed of only the N-terminus without linker (CpxR₁₋₁₁₇-T25). Elevated β-galactosidase activity indicated that the CpxR N-terminus alone could still interact with itself and with N-terminal CpxR containing linker (Figure 5B). Moreover, these interactions were consistently observed regardless of the T18/T25 fusion combinations used (data not shown). Consistently however, the highest reporter activity was achieved for the CpxR₁₋₁₃₂-T18/CpxR₁₋₁₃₂-T25 interaction pair i.e.: when the linker was present (Figure 5). These data therefore reveal that the linker is not essential for interactions of the CpxR-N terminus but may enhance them. On the other hand, we routinely observed low reporter output by strains harboring co-expressing CpxR N-terminal and C-terminal fusions (Figure 5) or between C-terminal fusions alone (data not shown). While this may be indicative that the

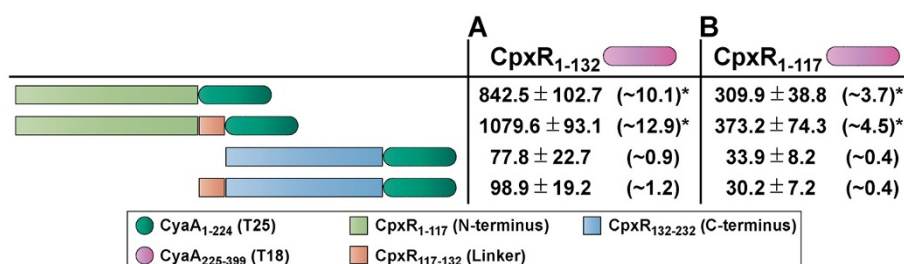


Figure 5 N-terminal CpxR dimerization in BACTH assays is enhanced by inclusion of the internal CpxR linker. The N-terminal domain of CpxR either with linker (A) or without linker (B) were translationally fused to the N-terminus of CyaA₂₂₅₋₃₉₉ (T18 – magenta shade) creating the ‘bait’ CpxR₁₋₁₃₂-T18 and CpxR₁₋₁₁₇-T18 hybrids respectively. As ‘prey’ hybrids, the CpxR N-terminus without linker (CpxR₁₋₁₁₇-T25) or with linker (CpxR₁₋₁₃₂-T25) and the C-terminus without linker (CpxR₁₃₂₋₂₃₂-T25) or with linker (CpxR₁₁₇₋₂₃₂-T25) were fused to the N-terminus of CyaA₁₋₂₂₄ (T25 – dark green). Interactions between ‘bait’ and ‘prey’ hybrids were again quantified via measurement of β-galactosidase activity (left columns; black font). Measurement of the interaction between T18-Zip and T25-Zip yielded 1449.1 ± 113.2 units of β-galactosidase activity was ~17.3 fold more than the enzymatic activity produced by negative-control bacteria co-expressing only T18 and T25 (83.8 ± 16.3 β-galactosidase activity units). The fold change in enzymatic activity caused by CpxR-CpxR interactions relative to this negative control is indicated in parentheses to the right. The asterisks (*) indicates a positive interaction. Data is presented as the mean (± standard error of the mean) of at least four independent experiments performed in triplicate.

C-terminus is not involved in CpxR dimerization *per se*, the data is inconclusive because we cannot confirm stable production of the C-terminal fusions, either by western blotting, or inferred by high reporter output from fusion-bearing strains. Nevertheless, the presented data definitively supports using the BACTH assay for studying dimerization of the CpxR N-terminus when produced in isolation.

Key structural features of CpxR mediate homodimerization

Since the CpxR N-terminus dimerizes with full-length CpxR (Figure 4), we designed another experiment in an effort to highlight structural regions within the full-length CpxR that may mediate specific interactions with the N-terminal region. As in Figure 4 and 5, CpxR₁₋₁₃₂-T18 was used as the ‘bait’ fusion. Additionally, we capitalized on the availability of variable ‘prey’ constructs previously used in Figure 2, which were composed of T25 fused to the N-terminal end of the larger CpxR constructs giving rise to T25-CpxR_{D51A}, T25-CpxR_{D51E}, T25-CpxR_{Δ11-24}, T25-CpxR_{Δ117-132} and T25-CpxR_{Δ188-209}. Native full-length CpxR fused to T25 (T25-CpxR₁₋₂₃₂) served as a control. In *E. coli* BTH101, the interacting hybrids CpxR₁₋₁₃₂-T18 and T25-CpxR₁₋₂₃₂ induced reporter activity that was ~18.9 fold higher than background (Figure 6). Only one other bait/prey combination – CpxR₁₋₁₃₂-T18 with the potentially ‘locked-on’ CpxR_{D51E} variant mimicking a phosphorylated state (T25-CpxR_{D51E}) – induced higher reporter activity of ~4.9 fold (Figure 6). Although we are unable to confirm the stable production of the non-phosphorylated and inactive CpxR_{D51A} variant (T25-CpxR_{D51A}), the low reporter activity of this fusion-bearing strain could indicate that interactions with N-terminal and full-length CpxR derivatives requires phosphorylation

at position D₅₁. The ability to interact was also lost upon generating Δ11-24, Δ117-132 and Δ188-209 deletions in *cpxR*. Hence, dimerization might also require CpxR to be inherently flexible, garnered by structural and/or functional elements encoded along the protein’s entire length. At least one of these regions encompassing residues 188 to 209 might be one such segment, since this fusion must be capable of being stably produced (see Figure 2).

Homodimerization of CpxR as determined by reconstitution of the λcl repressor

In parallel, we also employed a homodimerization assay based upon the reconstitution of the Lamda cI repressor (λcl) protein. This assay was effective in studying homodimerization of the YycG SK from *Staphylococcus aureus* [51] and the YycF RR from *B. subtilis* [36]. It is also an attractive basis for developing a high throughput screening system for the discovery of novel anti-bacterials targeting homodimerization of TCRS constituents [51]. Various alleles coding for different CpxR derivatives were translationally fused to the C-terminus of the first 131 residues of λcl (λcl₁₋₁₃₁-CpxR_n) expressed from plasmid pKWY2428 [36]. Reconstitution of an active repressor dimer can occur through the homodimerization of CpxR. In these situations where the fused CpxR protein does homodimerize, reconstitution of λcl₁₋₁₃₁ DNA binding activity specifically represses transcription from the *tet* resistance gene promoter present on the λcl₁₋₁₃₁-containing plasmid. This confers tetracycline sensitivity to the host bacterial strain *E. coli* JM109 [36]. An absence of homodimerization however will confer tetracycline resistance (Tet^R) to the host strain. Contrary to our BACTH analysis, full-length CpxR (λcl₁₋₁₃₁-CpxR₂₋₂₃₂) and the two C-terminal CpxR variants with linker (λcl₁₋₁₃₁-CpxR₁₁₇₋₂₃₂) or without linker (λcl₁₋₁₃₁-CpxR₁₃₃₋₂₃₂) were all able to

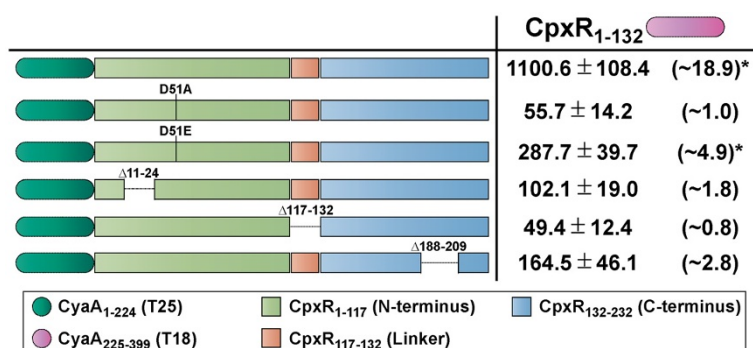


Figure 6 Regions of CpxR interacting with N-terminal CpxR as monitored by BACTH analysis. The N-terminal domain of CpxR either with linker was translationally fused to the N-terminus of CyaA₂₂₅₋₃₉₉ (T18 – magenta shade) creating the ‘bait’ CpxR₁₋₁₃₂-T18 hybrid. As already used in Figure 2, a selection of full-length or near full-length CpxR variants translationally fused to the C-terminus of CyaA₁₋₂₂₄ (T25 – dark green shade) were used as ‘prey’ hybrids; that is T25-CpxR_{D51A} and T25-CpxR_{D51E} lacking the phosphorylated aspartate residue as well as T25-CpxR_{Δ11-24}, T25-CpxR_{Δ117-132} and T25-CpxR_{Δ188-209} lacking the putative N-terminal coiled-coil, internal linker and C-terminal winged helix-turn-helix regions respectively. We also employed the T25-CpxR₁₋₂₃₂ hybrid containing wild type CpxR sequence. BACTH interaction analysis of ‘bait’ and ‘prey’ hybrids was quantified via measurement of β-galactosidase activity and is represented as units/mg dry weight of host *E. coli* BTH101 bacteria (left column; black font). The internal positive control based upon constructs expressing T18-Zip and T25-Zip yielded 1495.1 ± 237.3 units of β-galactosidase activity/mg dry weight of bacteria. This was on average ~25.6 fold more enzymatic activity produced compared to bacteria co-expressing only T18 and T25 (58.3 ± 14.1 units of β-galactosidase activity). The fold change in enzymatic activity caused by CpxR-CpxR interactions relative to this negative control is indicated in parentheses to the right. The asterisks (*) indicates a positive interaction. Data is presented as the mean (± standard error of the mean) of at least four independent experiments performed in triplicate.

homodimerize as determined by heightened sensitivity of JM109 (reduced growth) on tetracycline supplemented LB agar (Figure 7A). The extent of this growth restriction mirrored that achieved for the RR YycF C-terminal region encompassing residues 120 to 235 fused to λC_{I-131} and expressed from the plasmid pKWY-YycF(120C) [36] (Figure 7A). Interestingly, dimerization and growth suppression by the λC_{I-131}-CpxR₁₃₃₋₂₃₂ fusion occurred despite steady-state protein levels being quite minimal (Figure 7B). In further contradiction to our BACTH analysis (see Figure 5), we were unable to observe any growth restriction of JM109 when expressing λC_{I-131}-CpxR₂₋₁₁₆ or λC_{I-131}-CpxR₂₋₁₃₂ (Figure 7A), even though protein accumulated to high levels (especially in the case of λC_{I-131}-CpxR₂₋₁₃₂) (Figure 7B). This extensive growth was consistent with the negative control expressing only λC_{I-131} from the plasmid pKWY2428 [36] (Figure 7A). Hence, N-terminal CpxR with or without the internal linker could not homodimerize when fused at the C-terminus of λC_{I-131}, although independent fusions to T25 and T18 domains still support homodimerization (see Figure 5 and 6). Thus, the λC_I homodimerization assay is a tool especially suitable for studying dimerization of the CpxR C-terminus when produced in isolation.

To further explore λC_{I-131}-CpxR homodimerization, we analyzed essentially full-length CpxR derivatives carrying the specific deletions of the N-terminal coiled-coil (λC_{I-131}-CpxR_{Δ11-24}), internal linker (λC_{I-131}-CpxR_{Δ117-132}) and C-terminal wHTH domain (λC_{I-131}-CpxR_{Δ188-209}). In addition, we examined fusions harboring site-directed mutations in the N-terminal phosphorylation

site (λC_{I-131}-CpxR_{D51A} and λC_{I-131}-CpxR_{D51E}). Importantly, we could confirm that all of these fusion proteins accumulated in the cytoplasm of JM109 cells (Figure 7B). Clearly however, CpxR₁₈₈₋₂₀₉ exhibits a major defect in dimerization capacity (Figure 7A); a finding also observed in the BACTH assay (see Figure 6). This dimerization defect was also conferred to the fusion expressing only the C-terminus (λC_{I-131}-CpxR₁₁₇₋₂₃₂, Δ188-209) (Figure 7A), although this is less conclusive because this fusion was poorly expressed (Figure 7B). On the other hand, non-phosphorylated CpxR or CpxR lacking the N-terminal coiled-coil domain or internal linker, could all still homodimerize (Figure 7A). Once again, this data is at odds with our BACTH analysis, where a CpxR_{D51A}, CpxR_{Δ11-24} or a CpxR_{Δ117-132} mutation inhibited subsequent CpxR interactions (see Figure 6).

Discussion

TCRSs play a significant role in the regulation of physiological processes in the bacterial cell ranging from the control of single genes to diverse multi-cellular behavior. This occurs by means of transcriptional, post transcriptional and post translational regulation whereby TCRSs exploit a variety of protein-protein and protein-DNA interactions [3,42,52]. In this study, we attempted to gain some insight into the protein-protein interactions that are the hallmark of CpxA-CpxR signal transduction.

BACTH analysis revealed that full-length CpxR and the truncated N-terminal derivatives were able to interact together, albeit in a manner conditionally dependent upon the direction and orientation of the T18 and T25

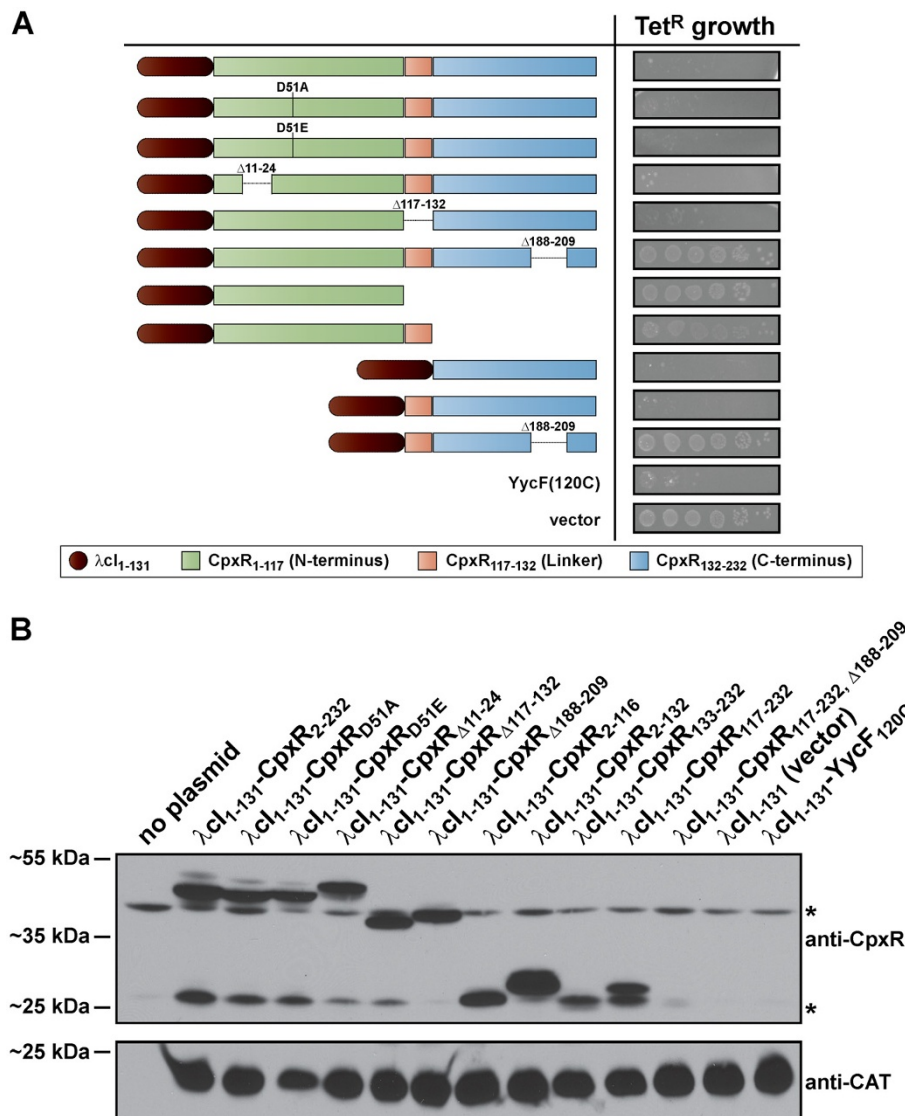


Figure 7 Reconstitution of the λclI repressor function through CpxR homodimerization. Fusion of CpxR_n variants to the C-terminus of λcl₁₋₁₃₁ (auburn brown shade) were generated. When expressed in *E. coli* JM109, growth was assessed by spotting 5 μl of 10-fold serially diluted (10⁰, 10⁻¹, 10⁻², 10⁻³, 10⁻⁴ and 10⁻⁵) exponentially grown cultures onto LB agar containing 10 μg/ml tetracycline (Tet) and 0.1 mM IPTG (A). The assay was controlled through the expression of inactive λcl₁₋₁₃₁ alone [vector; pKWY2428] or the dimerization-competent fusion λcl₁₋₁₃₁-YycF₁₂₀₋₂₃₅ [YycF(120C); pKWY-YycF(120C)] [36]. Protein lysates were also fractionated on 12% acrylamide SDS-PAGE and analysed by western blotting (B). Fusions of λcl₁₋₁₃₁ to the N-terminus of CpxR_n variants were detected with rabbit polyclonal anti-CpxR antiserum. Samples were also probed with antiserum raised in rabbit and specific for chloramphenicol acetyltransferase (anti-CAT) to confirm the loading of an equal quantity of protein in each lane. The asterisks (*) highlight unknown protein bands that cross-react non-specifically with antibodies in the sera or, in some cases may represent a degradation product of the recombinant fusion proteins. Shown to the left is the approximate mobility of molecular weight standards (PageRuler™ Plus Prestained Protein Ladder, Thermo Scientific).

fusions. Interactions of full-length CpxR were also observed in our λclI homodimerization analysis, but we were unable to validate dimerization of N-terminal truncates via this method. Interactions dependent upon the positioning of the fused CyaA domains might well indicate that the CpxR-CpxR interaction occurs as a head-to-head symmetrical dimer mediated by a N-terminal

α₄-β₅-α₅ interface highly conserved among OmpR-PhoB RR family members [30,42]. Indeed, structural studies of full-length and isolated N-terminal receiver domains of various RRs belonging to this OmpR-PhoB family suggest head-to-head dimerization is common and functionally important [53-56]. Hence, further studies are required to better understand how the α₄-β₅-α₅ interface

of CpxR determines CpxR dimerization orientation and contributes to subsequent DNA binding and gene transcription control.

Use of the λ CI assay indicated that the CpxR C-terminus could readily dimerize; a finding not observed during our BACTH analysis. Nevertheless, a previous study has demonstrated that solitary *trans*-expression of C-terminal CpxR is constitutively active and capable of transcriptional control of an array of target genes in *E. coli* [43]. Moreover, the degree of CpxR activation was independent of the internal linker [43]. This data was explained through an expectation that the C-terminal effector domain can alone dimerize to activate its target genes; a notion we can now experimentally verify via λ CI homodimerization analysis. Incidentally, this analysis also demonstrated that the linker was not required for this dimerization process as was previously assumed [43]. In addition to the ability to interact with itself, constitutive activation of the CpxR C-terminus implies that this region must engage one of the six subunits of the RNA polymerase holoenzyme in order to specifically modify its transcription output from targeted genes. Direct interaction with the RNA polymerase through the C-terminus is well established for other RRs of the wHTH family; prototype OmpR interacts with the α subunit [57-63] and PhoB the σ^{70} subunit [64-66]. Hence, it is of considerable interest to determine how the C terminus of CpxR coordinates dimerization, DNA binding and modulation of RNA polymerase activity in the pathogenic *Yersinia*.

The studies performed to elucidate the role of the phospho-modified residue (D51) were intriguing. The non-phosphorylated and functionally inactive CpxR_{D51A} variant could not interact with CpxA or other CpxR variants via BACTH analysis. The caveat here concerns our inability to confirm stable production of CpxR_{D51A} fused to either T18 or T25. Where production was confirmed in the λ CI homodimerization assay, λ CI₁₋₁₃₁-CpxR_{D51A} did dimerize. Clearly therefore, this aspect requires more investigation, especially given the suggestion that phosphorylation promotes homodimer formation within the OmpR/PhoB subfamily [50]. Intriguingly, the potentially 'locked-on' CpxR_{D51E} always maintained the ability to interact. This suggests that CpxR_{D51A} and CpxR_{D51E} are functionally distinct, the glutamate having the ability to mimic phosphorylation at residue 51 to maintain transcriptional activity. We have confirmed that CpxR_{D51E} does maintain function in *Y. pseudotuberculosis* (EJT *et al.*, unpublished), while purified CpxR_{D51A} is not phosphorylated *in vitro* by small phospho donors that in turn dramatically lessens target DNA binding ability [9]. It is evident therefore that conformational change induced by phosphorylation at D₅₁ permits certain inter- and intra-molecular interactions. Based on these data, we

anticipate that performing a thorough pair-wise molecular and biochemical characterization of the CpxR_{D51A} and CpxR_{D51E} variants will benefit our understanding of the complexities of CpxR-mediated transcriptional output mechanisms.

Our investigation into the role of the CpxR linker region (residues 117–132) in inter- and intra-molecular interactions was inconclusive. Both BACTH and λ CI homodimerization studies revealed the linker to be non-essential for homodimerization of any CpxR truncated variant. However, the near full-length CpxR Δ ₁₁₇₋₁₃₂ variant lost the ability to engage with the CpxR N-terminus and with full-length CpxA via BACTH analysis, although it still efficiently dimerized in λ CI-based studies. Hence, it seems that the linker has a function in facilitating protein-protein interactions, but this ability is context dependent. Recent studies of other members of the OmpR/PhoB subfamily are starting to shed some light on internal linker function [67-69]. Collectively, these demonstrate how linker sequence composition and length can influence RR interdomain interactions, DNA binding and subsequent transcriptional output. Presumably, the linker optimizes N- and C-terminal domain function by imparting physical separation as well as offering inherent flexibility that assists in domain conformational changes as a consequence of phosphorylation and/or DNA binding. Given this potential for influencing RR function, a detailed investigation of the CpxR linker is warranted.

The C-terminal wHTH DNA binding domain is also likely to contribute to CpxR homodimerization. While we and others have shown that truncated CpxR can dimerize even in the complete absence of the C-terminus (this study) [50], restricted removal of only residues 188 to 209 encompassing the wHTH prevents subsequent dimerization of CpxR Δ ₁₈₈₋₂₀₉ in both BACTH and λ CI homodimerization assays. While not tested here, the role of the wHTH domain in CpxR dimer formation might well relate to the effects of binding to target DNA. After all, it is established that DNA binding by non-phosphorylated OmpR is a mechanism to enhance phosphorylation and dimerization capacity [70,71]. Hence, the wHTH motif may play a dual role in both RR target DNA binding and dimerization. On the basis of our own DNA binding assays, where non-phosphorylated native CpxR [8], but not the non-phosphorylated CpxR_{D51A} mutant [9] only very weakly bound to target DNA, it is conceivable that initial binding can dramatically enhance phosphorylation and then dimerization.

Although both the BACTH and λ CI homodimerization assay were highly reproducible, they were often contradictory in their ability to predict interactions. This should not mean that these assays are unreliable or not physiologically relevant. However, it does highlight the

need to perform in parallel other complementing experimental analyses to ensure relevancy of the phenotypic data. No doubt, creation of hybrid proteins fused with structurally distinct CyaA or λ CI tags could diminish inherent protein flexibility or stability, curbing the subsequent ability to measure authentic protein-protein interactions. Indeed, a good correlation between protein instability and the absence of interactions in a BACTH assay has already been stated [37]. Despite our numerous attempts, we could not detect any T18 or T25 fusions, including those reliably inducing high reporter activities, such as the positive control. However, we could infer from numerous positive interaction data sets involving various T18 and T25 fusion combinations that the majority of our fusions must, to some extent, be stably produced. The notable exceptions to this deduction are the fusions involving CpxR_{D51A} and its C-terminal domain when in isolation. We are therefore confident that BACTH analysis is useful for CpxR N-terminal interaction studies, but it may have limited use for similar studies of the C-terminus. Critically, the majority of our λ CI fusions were produced and stable. Hence, product instability cannot account for the failure of λ CI₁₋₁₃₁-CpxR₂₋₁₁₆ and λ CI₁₋₁₃₁-CpxR₂₋₁₃₂ fusions to homodimerize. Moreover, as homodimerization of similar CpxR N-terminal fusions occurred via BACTH analysis, poor production alone cannot explain the contradictory results using the two reporter systems. This can be further illustrated by observations with the stably produced full-length CpxR; it could dimerize to itself only when fused to λ CI, but via BACTH analysis could only interact with CpxA or the CpxR N-terminus. Thus, it is apparent that individual domains appended to either CpxA_n or CpxR_n variants can alter their structural context in unique ways that significantly influence protein-protein interactions, making no one assay fully self-sufficient. By necessity therefore, future work will analyze the significance of these CpxR and CpxA interaction motifs by means of other sensitive interaction techniques in concert with analysis of pertinent *in vivo* phenotypes.

Conclusions

We have shown here that the BACTH and λ CI assays are two simple and complementary approaches amenable to identify the independent modular homo- and hetero-interactions of CpxA and CpxR. Notably, the former is ideally suited to analyze the molecular context of CpxR interactions involving the N-terminus, while the latter can aid in dissecting interactions of the CpxR C-terminus. Moreover, when combined with *in vivo* assays designed to assess bacterial survival and virulence, a better understanding of the roles of the CpxA-CpxR TCRS in *Y. pseudotuberculosis* will ensue.

Additional files

Additional file 1: Table S1. Bacteria strains and plasmids used in this study.

Additional file 2: Table S2. Oligonucleotides used in this study.

Abbreviations

ECS: Extracytoplasmic stress; Cpx: Conjugative pilus expression; RR: Response regulator; SK: Sensor kinase; TCRS: Two-component regulatory system; wHTH: winged helix-turn-helix; LB: Luria-Bertani; IPTG: Isopropyl- β -D-thiogalactopyranoside; X-Gal: 5-Bromo-4-chloro-3-indolyl- β -D-galactopyranoside; Ap: Ampicillin; Km: Kanamycin; Cm: Chloramphenicol; Tc: Tetracycline, BACTH, bacterial adenylate cyclase two-hybrid; CAP: Catabolite activator protein; cAMP: Cyclic adenosine monophosphate.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

ET and JM participated in study design, its coordination and performed all experimentation. MF conceived of the study, and participated in its design and coordination and drafted the manuscript. All authors analyzed the data and read and approved the final manuscript.

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