McbR/YncC: Implications for the Mechanism of Ligand and DNA Binding by a Bacterial GntR Transcriptional Regulator Involved in Biofilm Formation

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ABSTRACT: MqsR-controlled colanic acid and biofilm regulator (McbR, also known as YncC) is the protein product of a highly induced gene in early Escherichia coli biofilm development and has been regarded as an attractive target for blocking biofilm formation. This protein acts as a repressor for genes involved in exopolysaccharide production and an activator for genes involved in stress response. To better understand the role of McbR in governing the switch from exponential growth to the biofilm state, we determined the crystal structure of McbR to 2.1 Å. The structure reveals McbR to be a member of the FadR C-terminal domain (FCD) family of the GntR superfamily of transcriptional regulators (this family was named after the first identified member, GntR, a transcriptional repressor of the gluconate operon of Bacillus subtilis). Previous to this study, only six of the predicted 2800 members of this family had been structurally characterized. Here, we identify the residues that constitute the McbR effector and DNA binding sites. In addition, comparison of McbR with other members of the FCD domain family shows that this family of proteins adopts highly distinct oligomerization interfaces, which has implications for DNA binding and regulation.

Biofilms are complex communities of bacteria that are encased in an extracellular matrix and adhere to almost any surface. Due to properties of the biofilm, these bacterial communities are extremely tolerant to antibiotics and are often able to evade host defenses.1 Furthermore, it is estimated that 60–80% of human infections are caused by biofilms, explaining why much research is focused on elucidating the genetic basis of biofilm formation and proliferation.2,3 One regulator of biofilm formation is McbR/YncC (hereafter referred to as McbR), a transcription factor predicted to belong to the GntR family of DNA binding proteins. In Escherichia coli, deletion of mcbR results in the overproduction of colanic acid,4 a constituent of the biofilm exopolysaccharide (EPS) matrix composed of glucose, galactose, fucose, and glucuronic acid in the ratio 1:2:2:1.5 As a consequence, mcbR deletion results in a mucoidy phenotype and a reduction in biofilm formation. In E. coli, gene array studies coupled with electrophoretic mobility shift assays (EMSAs) showed that McbR binds the ybtM promoter, a gene encoding a putative periplasmic protein whose function is currently unknown.4 A subsequent study using DNA footprinting experiments showed that McbR from Salmonella typhimurium and E. coli binds the yciG promoters from both species.6

The GntR superfamily (Pfam PF00392), to which McbR belongs, is one of the largest families of transcriptional regulators, with more than 8500 members (Figure 1A).7 Members of this family contain an N-terminal DNA binding winged helix-turn-helix (wHTH) domain and a C-terminal effector binding/oligomerization domain. In contrast to the wHTH domain, which is structurally conserved in the GntR family, the C-terminal domain is highly variable. Detailed bioinformatics studies have led to the definition of at least 7 families (AraR, DevA, FCD, HutC, MocR, PlmA, and YtrA), which are classified by the effector binding domain topology and secondary structure (Figure 1A).7 The majority of GntR regulators belong to the FadR C-terminal domain family (FCD, Pfam PF07729; Figure 1A). The effector molecule that regulates the activity of GntR transcriptional regulators is often a product/substrate in the metabolic pathway that the particular GntR transcription factor controls. However, there are examples where this is not the case, and difficulties in identifying the endogenous ligands for this family have limited

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our understanding of how these regulators function in vivo.7 What is known is that effector binding in the C-terminal effector binding domain alters, via a poorly understood structural mechanism(s), the conformations and/or relative orientations of the N-terminal wHTH domains. This, in turn, inhibits DNA binding.7 This limited understanding is due to the dearth of structural data available for the GntR superfamily, especially those of the FCD family. Here, we describe the structure of McbR from E. coli to 2.1 Å resolution. We show that McbR belongs to the FCD family of transcriptional regulators and identify the residues that mediate DNA binding. We also identify the residues that constitute its effector binding site, which are highly conserved in mcbR homologs. Finally, a comparison of the currently available structures of FCD transcriptional regulators reveals different effector binding interfaces at the wHTH domains,8–10 suggesting that this family of proteins undergoes distinct conformational rearrangements upon ligand binding.

## MATERIALS AND METHODS

### Protein Expression and Purification

Two constructs of wild-type McbR (McbR1–221) (full-length), residues 1–221; McbR10–221, residues 10–221) were subcloned into the pRP1B bacterial expression vector, which contains a N-terminal His8-tag and Tobacco Etch Virus (TEV) cleavage site;11 both constructs were sequenced prior to subsequent experiments. pRP1B-McbR1–221 variants (single-mutant variants: Arg34Ala, Lys38Ala, Thr49Ala, Arg52Ala, Gln70Ala; double-mutant variant: Gln93Ser/Arg139Phe; triple-mutant variant: Arg89Ala/Lys38Ala/Thr49Ala) were generated using the Quick-Change Mutagenesis Kit (Agilent Technologies) using the manufacturer’s protocols; all constructs were verified by sequencing.

WT McbR and McbR variants were expressed in E. coli BL21-Gold (DE3) cells (Agilent). Cells were grown at 37 °C (250 rpm) to an OD600 of ~0.9, at which point the cells were transferred to 4 °C for 1 h. The cells were induced with 0.5 mM IPTG and grown overnight at 18 °C (250 rpm). The cells were then harvested by centrifugation at 6000g. Selenomethionine (SeMet)-labeled McbR10–221 was produced using identical protocols, with the exception that the cells were grown in minimal medium supplemented with vitamins, metals, and amino acids (with selenomethionine substituted for methionine).12

For purification, cell pellets of either WT McbR or McbR variants were resuspended in lysis buffer (50 mM Tris, pH 8.0, 500 mM NaCl, 0.1% Triton X-100, 5 mM imidazole, complete tabs-EDTA free [Roche]) and lysed by high-pressure homogenization (C3 Emulsiflex; Avestin). Following centrifugation (45 000g, 45 min, 4 °C), the supernatant was applied to a HisTrap HP column (GE Healthcare), and McbR was eluted using a 5–500 mM imidazole gradient. McbR was then incubated overnight with TEV protease (50 mM Tris, pH 8.0, 500 mM NaCl, 4 °C). The following day, McbR was further purified using Ni-NTA (Qiagen) to isolate the cleaved protein from the TEV protease (itself His8-tagged) and the cleaved His8-tag. After concentration, McbR was purified in a final step using size-exclusion chromatography (SEC; Superdex 200 26/60; 20 mM Tris, pH 7.8, 100 mM NaCl, 0.5 mM TCEP). To determine the oligomerization state of McbR, the elution volume was compared to that of MW weight standards (Bio-Rad).

### Crystallization, Data Collection, and Processing.

SeMet McbR10–221 was concentrated to 8–10 mg/mL, incubated for 1 h with glycerol (10% (v/v), final concentration), and used immediately for crystallization trials. Microcrystals of SeMet McbR10–221 were obtained in 2 M sodium malonate pH 7.0 (sitting drop vapor diffusion; 25 °C) and used as seed solution to produce crystals in the same conditions suitable for data collection. The crystals were cryoprotected in mother liquor containing 20% (v/v) MPD and immediately frozen in liquid nitrogen. Data for McbR10–221 were collected at the National Synchrotron Light Source, beamline X25, using a Pilatus 6 M detector (Dectris). Anomalous data was collected from a single crystal and phased using single anomalous dispersion (SAD), collecting data at 0.93 Å. Data were processed and scaled using HKL2000,13 The asymmetric unit contains two protein molecules. The anomalous data was phased using HKL2MAP14 (ShetkC/D/E);15–17 14 of the 16 expected selenium sites were identified. Approximately 90% of the structure was built automatically using ARP/wARP.18 Model building and refinement of SeMet McbR was carried out using a high-resolution data set (2.1 Å) collected at 0.9793 Å. Iterative model building and refinement were performed using COOT19 and Phenix.20 The final model was refined with Phenix using TLS. Molprobity was used for model validation.21 Analysis of the dimerization interface was performed using the Protein Interaction Calculator,22 with solvent-accessible surface areas calculated using Naccess.23 Cavity volumes were calculated using POCOSA.24 Data collection and structure refinement statistics are reported in Table 1.

![Figure 1. McbR classification and structure. (A) Flowchart illustrating the classification of McbR within the GntR superfamily. (B) McbR monomer with all secondary structural elements annotated. The N-terminal winged helix-turn-helix (wHTH) domain is shown in light blue, and the C-terminal FCD domain is shown in teal; the wing loop of the wHTH domain is labeled. McbR residues 10–220 were observed in the electron density maps for subunit A.]
Table 1. Crystallographic Data Collection and Refinement Statistics

| Crystal Data          |   |
|-----------------------|---|
| space group           | P6₁ |
| no. McbR/ASU          | 2  |
| a, c (Å)              | a = 107.6, c = 72.7 |

| Data Collection       |   |
|-----------------------|---|
| wavelength (Å)        | 0.9793 0.9300 |
| unique reflections    | 28120 50871 |
| resolution (Å)        | 50.0−2.1 (2.14−2.10) 50.0−2.15 (2.19−2.15) |
| mean 1/σ             | 18.1 (3.0) 18.4 (2.5) |
| completeness (%)      | 100.0 (100.0) 99.3 (98.7) |
| redundancy (%)        | 9.8 (10.0) 5.1 (5.2) |
| Rmerge (%)            | 8.3 (55.8) 6.1 (45.4) |

| Refinement            |   |
|-----------------------|---|
| Rwork (%)             | 16.8 |
| Rfree (%)             | 20.8 |
| no. non-hydrogen atoms | 3214 |
| no. water molecules   | 163 |
| Ave. B-factor (Å²)    | 43.6 |
| rmsd bond length (Å)  | 0.008 |
| rmsd bond angle (deg) | 1.003 |
| Ramachandran Plot     | 99.8 |
| allowed (%)           | 100.0 |
| disallowed (%)        | 0 |
| PDB code              | 4PF |

“Highest resolution shell data are shown in parentheses. b5% of the reflections used for Rfree. cRmerge = \sum_{i=1}^{\text{all}}|I(hkl)|/\sum_{i=1}^{\text{all}}I(hkl) where I(hkl) is the ith observation of a symmetry equivalent reflection hkl. Reported values for the 0.93 Å data set are for unmerged Friedel pairs.

Electrophoretic Mobility Shift Assay. The E. coli yciG promoter (PyciGECO) was used for DNA binding studies. Following synthesis of the individual oligonucleotides (IDT Technologies; each oligonucleotide includes a 3’ biotin label; Table 2), the complementary oligonucleotides were combined, heated at 95 °C, and then cooled at 1 °C/min to a final temperature of 25 °C. For EMSA experiments, 1 pmol of protein was added to the biotin-labeled DNA (PyciGECO, 100 fmol). All reactions were carried out in binding buffer (10 mM Tris, pH 7.5, 50 mM KCl, 1 mM DTT) in the presence of a poly(dI−dC) DNA probe (50 ng/µL) to prevent nonspecific binding. For the unlabeled competitor EMSA control, a 100-fold excess of unlabeled PyciGECO DNA was added. All binding reactions were incubated at room temperature for 20 min. Samples were then loaded onto a 6% DNA retardation gel (Invitrogen) and subjected to electrophoresis at 4 °C for 75 min at 100 V in 0.5-fold TBE buffer (45 mM Tris, pH 8.3, 45 mM Boric acid, 1 mM EDTA). The DNA was transferred to a nylon membrane at 390 mA for 30 min, followed by UV cross-linking at 302 nm by placing the membrane face-down on a UV illuminator for 15 min. Chemiluminescence was performed using the LightShift Chemiluminescent EMSA Kit (Pierce), and the samples were detected using a CCD imager (Typhoon 9410 Imager).

Mucoidy Assay. WT mcbR and two mutated mcbR variants (mcbR-E93S-R139F and mcbR-R89A-E93A-R139A) were subcloned from pRP1B-mcbR, pRP1B-mcbR-E93S-R139F, and pRP1B-mcbR-R89A-E93A-R139A into the KpnI and SacI sites of pBS(Kan) via using primers in Table 3 so that their expression is under control of a lac promoter (instead of a T7 promoter in the former plasmids). The resulting plasmids are pBS(Kan)-mcbR, pBS(Kan)-mcbR-E93S-R139F, and pBS(Kan)-mcbR-R89A-E93A-R139A. After verifying these plasmids by DNA sequencing, they were introduced into E. coli MGI65S ΔmcbR ΔKm via electroporation, and the transformed clones were plated on LB agar supplemented with 50 µg/mL kanamycin and 0.2% (w/v) glucose. At least three independent colonies were streaked on LB agar supplemented with 50 µg/mL kanamycin and 1 mM IPTG to test the mucoidy of each strain. Cells were incubated at 37 °C for 12 h.

## RESULTS AND DISCUSSION

McbR Is a Member of the VanR Subfamily of GntR Transcriptional Regulators. Two constructs of McbR were screened for their ability to form diffraction-quality crystals: MbcR−221 and MbcR−221. The latter is missing the first 9 amino acids, which were predicted to be disordered (PSIPRED, IUPRED). Only MbcR−221 formed crystals suitable for structure determination and is referred to hereafter as McbR. The crystal structure of McbR was determined by single-wavelength anomalous dispersion (SAD) using SeMet-labeled protein, and the atomic model was refined to 2.1 Å resolution (Table 1 and Figures 1B and 2A). Two molecules of McbR are present in the asymmetric unit and are related by a nearly perfect 2-fold axis (179.8°); superposition using the C-terminal domain is composition-quality crystals: McbR−221 and McbR−221. This is consistent with the observation that McbR is predominantly a dimer in solution (Figure 2B). McbR, like other members of the GntR family, consists of an N-terminal wHTH domain (residues 10−76; residues 38−46 in the second subunit were not modeled due to a lack of clear electron density) and a C-terminal all-α-helical effector binding domain (residues 77−219; Figure 1B). The wHTH domain is composed of three α-helices (α1−α3) and three β-strands (β1−β3), which form a small β-sheet that is a defining characteristic of the wHTH fold. The C-terminal domain is composed of six α-helices (α4−α9). The secondary structure elements and topology of the C-terminal domain places McbR in the FadR C-terminal domain (FCD) family of GntR transcriptional regulators (Figure 1A).

McbR Dimerization Interface Is Extensive and Includes Both the N- and C-Terminal Domains. The FCD family contains more than 400 distinct species from archaea to eukaryota. A structural homology search using Dali identified only 6 other structures that have a

Table 2. PCR Primers Used To Generate PyciGECO for EMSA Experiments

| Primer name      | sequence (5’−3’)           |
|------------------|----------------------------|
| PyciGECO-Fb      | AATTGTTTATATATCCAGAATGTTCTCCAAATTATTTTTCCCTCTAT |
| PyciGECO-Rb      | ATAGAGGGAAAAATATTTTTGAGAGACATTGATTAAACAATT |

“F indicates forward primer, and R indicates reverse primer. bPrimer contains a 3’ biotin label.
high degree of similarity to McbR (Z-score > 7 using only the FCD domain; Table 4 and Figure 3). These represent the only other members of the FCD family with known structures. The dimerization interface mediated by the C-terminal FCD domain is topologically conserved within the FCD family and is composed of the first helix in the FCD domain (α4 in McbR) and the N-terminal half of the kinked fourth helix (α7 in McbR). In McbR, the FCD dimerization interface buries 1655 Å² of solvent-accessible surface area, which is 70% of the buried surface area (BSA) for the entire McbR dimer. The hydrophobic core of the FCD dimerization interface is formed by residues Ile85, Ile88, Leu92, Met148, Ile150, Leu151, Met154, and Leu158 from both monomers, each of which is completely occluded from solvent (Figure 2C). It is further stabilized by polar and salt bridge interactions, especially a bidentate hydrogen bond between Gln157A and Gln157B and a bidentate salt bridge between Arg161A and Glu153B (A or B subscript indicates that the residue is from subunit A or B, respectively; Figure 2D).

In McbR, the wHTH domains also interact, extending the dimerization interface beyond that typically observed in the FCD subfamily of GntR regulators. The wHTH interface buries 740 Å² of BSA, for a total of 2395 Å² buried between the two McbR monomers. Although the FCD domains are related by a near perfect 2-fold axis centered on Gln157A/B, the wHTH are not. Instead, they are related by a rotation of ∼172°. Thus,
Figure 3. Quaternary structures of FCD family. FCD family members whose structures have been determined are shown, with one monomer depicted in teal and one in gold. Metals bound to the FCD domains are depicted as magenta spheres or sticks. The dimerization helices (α4 and α7 in McbR) are colored green and orange. The corresponding quaternary structures are depicted as cartoons, with the N-terminal domains shown as triangles and the C-terminal domains as spheres. (A) Head-to-head dimerization in which both the wHTH domains and the FCD domains contribute to the dimerization interface. (B) Dimerization in which the wHTH domains do not interact with either one another or the FCD domains. (C) Head-to-head dimerization in which only the FCD domains contribute to the dimerization. (D) Domain swapping dimerization in which the wHTH domain of one monomer reaches across the FCD domain to interact with the FCD domain of the second monomer. (E) Same as panel D except that the wHTH domains are inverted with respect to one another.

whereas Leu56\textsubscript{A} is buried in the wHTH interface, the corresponding residue (Leu56\textsubscript{B}) is located at the interface periphery (Figure 2E). The wHTH dimerization is composed largely of polar interactions (i.e., a hydrogen bond between Ser60\textsubscript{A/B} and Arg57\textsubscript{A/B} and a few hydrophobic interactions (Leu14\textsubscript{A} and Leu65\textsubscript{A}); however, unlike the residues at the FCD interface, none of the wHTH interface residues become extensively buried upon complex formation (Figure 2E). Finally, Asn62\textsubscript{A} (wHTH domain) hydrogen bonds with Glu153\textsubscript{B} (FCD domain); this is the only noncovalent interaction connecting the two different domains from the distinct subunits in the dimer.

Comparison of McbR with the other members of the FCD subfamily reveals that while the FCD dimerization interface is conserved within the family, the relative orientation of the wHTH and FCD domains is not. This gives rise to distinct differences in the orientations of the wHTH domains and, in some cases, distinct quaternary structures (Figure 3). This is why the FCD family member identified to be most similar to McbR using the DALI structural homology search database changes depending on whether the search is performed with the McbR FCD domain alone (PSS454, PDB ID: 3C7J) or full-length McbR (Reut\_B4629, PDB ID: 3IHU; Table 4). As expected, the FCD proteins identified as most different from full-length McbR are FadR and CGL2915. These are also both members of the FadR subfamily, but they have an additional helix between the wHTH and the FCD ligand binding domains (Figure 4). The presence of this helix leads to domain swapped quaternary structures, in which the wHTH domain of subunit A across the dimerization interface to make contacts with the FCD domain of subunit B. This domain swapping is not observed in the VanR subclass of FCD regulators, and instead, in these proteins, the wHTH and FCD domains of the same subunit are more intimately associated.

The Structure of McbR Is Predominantly in a Ligand-Bound Conformation. The C-terminal FCD domains are composed of either 6 (VanR subclass) or 7 (FadR subclass) α-helices that form an antiparallel helical bundle. McbR, which has 6 helices, is a member of the VanR subclass (Figure 1A). The FCD domains have a large cavity in the center of this helical bundle, which is the location of the ligand binding site. The structure of this cavity is identical between both FCD domains in McbR, as the FCD domains superimpose with a root-mean-squared deviation (rmsd) of only 0.24 Å (Figure 6B). Although the helical topology is conserved among FCD domains, the sequence conservation among FCD family members, especially the residues that line the ligand binding cavities, is very low, likely reflecting their distinct ligand specificities (Figure 4A).

Recently, it was shown that the majority of FCD family members use three conserved histidines to bind a metal ion in the ligand binding cavity, suggesting that these regulators bind ligands that interact directly with the bound metal (Figure 4A,B). In McbR, these histidines are not conserved and are instead replaced by Arg139, Tyr185, and Ile207 (Figure 4B). Thus, McbR is one of the few FCD family members that does not bind a metal. Because of this, the ligand pocket in McbR is large, with a volume of ~200 Å\textsuperscript{3}, nearly double that of the metal-binding FCD domains (Figure 4C). Although the endogenous ligand for McbR is still unknown, clear unambiguous density for a bound entity was observed in the FCD ligand binding cavities of both monomers of McbR (Figure 4D). None of the protein and crystallization buffer components, or derivatives thereof, fit the density. This is likely because the density is rather undefined, potentially because it is not fully occupied, a phenomenon commonly observed without externally supplied ligands and/or cofactors. Alternatively, the density could correspond to the biologically relevant ligand, as McbR is an E. coli protein and was expressed in E. coli. However, potential ligands, such as glucuronic acid, a component of colanic acid whose metabolism has been shown to be regulated by McbR, did not fit the density. Finally, automated ligand fitting routines, such as the LigandFit program implemented in Phenix, also failed to identify a ligand that satisfactorily fit the density. Because the density did not enable the identity of the ligand to be confidently determined, it has not been modeled.

However, the presence of the density did reveal the identity of the residues that likely define the McbR ligand binding site. Namely, the bound entity is strongly coordinated by two histidines, Arg89 and Arg139, which themselves are organized via a shared salt bridge with Glu93 (Figures 4C,D). Two neighboring asparagine residues, Asn135 and Asn211, also contribute to binding. To investigate whether these residues are important for McbR function, we generated two variants of McbR by mutating the residues that define the entity binding site. Because mutating residues in the interior of a protein can also lead to protein unfolding, we generated two distinct mutants: a double mutant in which Glu93 and Arg139 were
substituted with Ser and Phe, respectively, the structurally homologous residues in FadR (the residue structurally homologous to McbR Arg89 is also an Arg in FadR) and a triple mutant in which all three residues were mutated to alanines (Arg89Ala, Glu93Ala, and Arg139Ala). CD polarimetry demonstrated that both McbR variants are folded, and EMSAs showed they are functional (Figure S1). The mutants were somewhat less thermostable (Δ$T_m$ of −8.6 and −18.6 °C compared to WT for the double and triple mutant, respectively), but this was expected, as the mutations are in the interior of the protein; indeed, this is exactly why two mutants, one in which the residues were mutated to those present in FadR (the double mutant) and one in which the residues were simply mutated alanine (the triple mutant), were tested. McbR deletion from E. coli results in EPS overproduction and elicits a mucoidy phenotype.4 This mucoidy phenotype is substantially reduced upon producing McbR ectopically (Figure 5). However, cells producing McbR with triple mutations (Arg89Ala, Glu93Ala, and Arg139Ala) are mucoid (Figure 5). This observation demonstrates the importance of Arg89, Glu93, and Arg139 in binding the unknown ligand and, more importantly, the physiological relevance of the unknown ligand in affecting EPS production.

Both Arg89 and Arg139 are required for ligand binding, as cells producing McbR with only two mutations (Glu93Ser and Arg139Phe) also remain less mucoid than cells with empty plasmid or cells producing McbR with three mutations (Arg89Ala, Glu93Ala, and Arg139Ala) (Figure 5).

The Conformation of McbR Crystallized Is Likely Incompatible with DNA Binding. The $\alpha$HTH domain is defined by helix $\alpha_2$, a connecting turn, and helix $\alpha_3$ (HTH) and a small loop in the antiparallel $\beta$-sheet (the wing).
whTH domain is slightly more conserved than the FCD domain (5% identity, 13% similar) when comparing the 7 structurally characterized FCD family members, with McbR residues Leu20, Leu24, Leu29, Gly32, Leu35, Leu40, Leu44, Met46, Val51, Arg52, Glu53, Leu55, Leu58, and Leu64 being highly similar (Figure 4A). The conserved hydrophobic residues function to stabilize the whTH domain fold, whereas the two charged residues are located at the whTH dimerization interface (Figure 2E).

In McbR, the C-terminal portion of whTH helix α1 contributes to the top of the FCD binding cavity, with Ile26 (helix α1) ~12 Å away from the FCD domain ligand coordinating arginines (Arg89 and Arg139; Figure 6A). Thus, this whTH–FCD interface provides a conduit by which effector binding in the FCD domain can be structurally communicated to whTH DNA binding domain. The conformation and orientation of the McbR whTH domains appear to be incompatible with DNA binding. First, residues 37–48, which comprise helix α2, are disordered in subunit B (Figure 2A). Residues from helix α2 often contribute to DNA recognition, as has been observed for the FCD transcription factor FadR. Second, the two domains in McbR differ not only in their relative orientations to the FCD domain but also in conformation, with an rmsd of 1.2 Å (Figure 6B). This is due to a change in the orientation of the wing between strands β2 and β3.

Implications for McbR Function. Currently, E. coli FadR is the only member of the GntR family whose DNA-bound structure has been determined, revealing that FadR binds the short palindromic consensus sequence 5′-TGGNNNNNCCA-3′. Previously, the E. coli McbR protein was shown to bind upstream of the E. coli yciGFE promoter (PyciGECO). Subsequent DNAseI footprinting identified two distinct DNA sequences within PyciGECO protected by McbR binding. To confirm that McbR binds this operator, we performed EMSA experiments using WT McbR and PyciGECO DNA. As shown in Figure 7, McbR binds and shifts PyciGECO DNA.

Figure 5. Mucoidy level of E. coli MG1655 ΔmcbR ΔKm producing different McbR variants. Each strain was grown on LB agar supplemented with 50 μg/mL kanamycin and 1 mM IPTG at 37°C for 12 h. WT/empty, E. coli MG1655/pBS(Kan); ΔmcbR/empty, E. coli MG1655 ΔmcbR ΔKm/pBS(Kan); ΔmcbR/mcbR, E. coli MG1655 ΔmcbR ΔKm/pBS(Kan)-mcbR ΔmcbR/ΔmcbR ΔKm/pBS(Kan)-mcbR ΔKm/pBS(Kan)-mcbR-E93S-R139F, E. coli MG1655 ΔmcbR ΔKm/pBS(Kan)-mcbR-E93S-R139F, ΔmcbR/R89A-E93A-R139A, E. coli MG1655 ΔmcbR ΔKm/pBS(Kan)-mcbR-R89A-E93A-R139A.

Figure 6. N-Terminal domain of McbR. (A) The pocket of α1 into the FCD domain. Helices α6 and α7 are colored teal (cartoon), and helix α1 is shown in light blue (sticks). (B) Superposition of McbR chain A (light blue/deep teal) and McbR chain B (light green/dark green). Whereas the C-terminal domains superimpose well (deep teal/dark green), the N-terminal domains (light blue/light green) do not.

Figure 7. McbR:PyciGECO EMSA experiments. (A) Superposition of the N-terminal domain of McbR (teal) and E. coli FadR (beige, PDB ID: 1HW2) bound to DNA. Residues making base-specific contacts in FadR and the structurally overlapping residues in McbR are shown as sticks and labeled. (B) EMSA experiments using biotin-labeled PyciGECO and WT McbR (the migration of the DNA alone is shown in the left lane). (C) EMSA experiments using biotin-labeled PyciGECO and either WT McbR or the McbR variants as indicated; the migration of the DNA alone is shown in the left lane. All binding reactions in panels B and C contain the nonspecific poly(dI–dC) probe.
α3 (Thr49). Superposition of the FadR–DNA complex and McbR shows Gln70 as the only residue with a polar side chain in close proximity with the DNA in the β2–β3 wing. Additional basic residues in close proximity to the DNA include Arg34 and Arg52 (Figure 7A). We tested the role of these residues in DNA binding using EMSA experiments performed with the PyciGECO promoter DNA and McbR mutants (we used CD to show that the variants are folded; Figure S1; the Tm’s of the variants are within 3.7 °C of that of WT, which has a Tm of 63.2 °C). The EMSA experiments show that residues Arg34, Lys38, Thr49, and Arg52 are important for DNA binding, as mutating these residues to alanine result in a loss of DNA binding compared to that of WT McbR (Figure 7B). Furthermore, Arg34, Lys38, and Thr49 have the most debilitating effects, suggesting that β1, α2, and α3 play key roles in DNA binding.

So, how is DNA binding regulated? As stated earlier, the GntR transcription factors are typically regulated by ligands that are metabolic substrates/products/cofactors of the genes that they regulate. In many cases, these genes are often located next to or near the GntR gene itself. McbR was previously shown to bind the promoter of yciGFE and ybiM. While the molecular functions of the protein products of these genes are currently unknown, ybiM has been shown to effect colanic acid production in a McbR-dependent manner, suggesting that colanic acid, or one of its constituents, may be the biologically relevant ligand for McbR. Currently, our results suggest that this is not the case, as none of the components of colanic acid satisfactorily fit the ligand density in the McbR cavity. An examination of the genes near mcbR in the E. coli chromosome shows that they are involved in a variety of biological processes (Table 5); our data again shows that McbR is unlikely to be regulated by these metabolites (methionine, curcumin/dihydrocurcumin, iron, asparagine, and glutathione), as they also did not satisfactorily fit the density. However, sequence similarities between the E. coli and Salmonella McbR do suggest that they likely bind similar, if not identical, ligands. Namely, although the FCD domains of McbR from both organisms are less conserved than their corresponding wHTH domains (FCD domain sequence conservation: 46% identity, 74% similarity), the ligand binding residues are nearly perfectly conserved, including Arg89, Glu93, and Arg139 (Figure 4A); the only differences in the ligand binding pocket are distal from the Arg-Glu-Arg pocket: Ile214 and Leu215 (Thr214 and Thr215 in Salmonella). Because these residues change from hydrophobic (E. coli) to polar (Salmonella), the distal portion of the ligand may be slightly different between the organisms. Once the biologically relevant ligand(s) of McbR have been confidently identified, this ligand, or a derivative thereof, may be able to function as a novel therapeutic to target biofilms.

### ASSOCIATED CONTENT

#### Supporting Information

CD spectra and EMSA of McbR variants (Figure S1). This material is available free of charge via the Internet at http://pubs.acs.org.

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### ABBREVIATIONS

ASU, asymmetric unit; CD, circular dichroism; EPS, exopolysaccharide; EDTA, ethylenediaminetetraacetic acid; EMSA, electrophoretic mobility shift assay; FCD, FadR C-terminal domain; IPTG, isopropyl-β-D-thiogalactopyranoside; MPD, 2-methyl-2,4-pentanediol; PDB, Protein Data Bank; SEC, size-exclusion chromatography; SeMet, selenomethionine; TCEP, tris(2-carboxyethyl)phosphine; TEV, Tobacco Etch Virus; wHTH, winged helix-turn-helix; WT, wild-type

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Table 5. DNA Sequences Surrounding the mcbR (ynC) Gene in Escherichia coli (MG1655)

| gene | other names | gene description |
|------|-------------|------------------|
| yncA | mnaT, b1448 | Methionine N-acyltransferase; l-amo acid N-acyltransferase |
| yncB | curH, b1449 | Curcumin/dihydrocurcumin reductase, NADPH-dependent |
| yncD | b1451 | Predicted inner membrane transporter |
| yncE | b1452 | ATP-binding protein, periplasmic, function unknown |
| yncF | ansP, b1453 | l-asparagine transporter |
| yncG | b1454 | Glutathione S-transferase homologue |
| yncH | b1455 | Conserved protein, function unknown |
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