Crystal Structure of the Regulatory Domain of MexT, a Transcriptional Activator of the MexEF-OprN Efflux Pump in Pseudomonas aeruginosa

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INTRODUCTION

The opportunistic human pathogen Pseudomonas aeruginosa is clinically important due to its resistance to multiple antibiotics, which can be intrinsic or acquired (Piddock, 2006). In addition to its poor outer membrane permeability, chromosome-encoded efflux pumps are the major determinants of antibiotic resistance in P. aeruginosa (Sobel et al., 2005). The resistance-nodulation-cell division (RND) family efflux pumps are key contributors to the multidrug resistance of P. aeruginosa and are the most common efflux pumps found in Gram-negative bacteria (Piddock, 2006; Poole and Srikumar, 2001). To date, four RND types of efflux pumps have been identified and characterized in P. aeruginosa—MexAB-OprM, MexXY-OprM, MexCD-OprJ, and MexEF-OprN (Hinchliffe et al., 2013; Piddock, 2006). Studies of MexAB-OprM pumps and associated molecular mechanisms have demonstrated their importance for multidrug resistance (Jeong et al., 2016; Kim et al., 2015; Lister et al., 2009; Xu et al., 2012). For example, MexEF-OprN exports several types of antimicrobial compounds such as chloramphenicol, fluoroquinolones, and trimethoprim (Galie et al., 2018). In several laboratory strains and clinical isolates, MexEF-OprN affects the expression of MexAB-OprM, indicating the existence of regulatory interplay...
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between the two efflux pumps (Horna et al., 2018).

The expression level of MexEF-OprN in wild-type strains under laboratory conditions is typically negligible. However, treatment with the fluoroquinolone antibiotic norfloxacin leads to generate certain types of spontaneous mutant strains, such as nfxC-type mutants. And these showed constitutive expression of mexT transcriptionally (Fargier et al., 2012; Hirai et al., 1987). Activation of MexT induces over-expression of MexEF, leading to increased resistance to antibiotics (Kohler et al., 1997). In the P. aeruginosa genome, mexT is located in the upstream of mexEF-oprN operon. MexT reportedly promotes the transcription of the operon (Kohler et al., 1997; 1999). Additionally, the broad regulatory scope including multiple target genes of MexT in P. aeruginosa is being revealed adding weight on its potential as a global regulator (Tian et al., 2009). Indeed, MexT is linked to several genes associated with pathogenesis and the host-pathogen interaction (Fargier et al., 2012; Fetar et al., 2011; Jin et al., 2011; Juarez et al., 2018).

We conducted the experiments to evaluate the molecular mechanism of MexT by understanding its structure. We investigated the mechanism by which MexT regulates the expression of MexEF-OprN by producing the full-length (FL) recombinant protein in Escherichia coli. We analyzed the crystal structure of the MexT regulatory domain (RD), which encompasses putative ligand-binding sites.

MATERIALS AND METHODS

Protein expression and purification

The native MexT RD (residues 95-304) and FL MexT (residues 1-304) were expressed as described previously (Kim et al., 2018). For the selenomethionyl (SeMet)-labeled MexT RD, the host E. coli B834(DE3) was cultured in M9 medium supplemented with L-(+)-selenomethionine, 100 µg/ml ampicillin, and other cofactors. To induce production of SeMet-labeled MexT RD, 0.5 mM isopropyl β-D-1-thiogalactopyranoside (IPTG) was added, followed by incubation at 16°C for 14 h with shaking. The cells were harvested by centrifugation at 1,380g for 10 min and resuspended in 50 ml of lysis buffer (20 mM Tris-hydrochloride [HCl; pH 8.5], 300 mM sodium chloride [NaCl], 5% [v/v] glycerol, and 2 mM 2-mercaptoethanol). The cells were disrupted using a continuous French press (Beijer, Sweden) at a pressure of 23 kpsi and the lysate was clarified using centrifugation at 10,000g for 30 min. MexT was subsequently purified using TALON affinity chromatography and the RD protein was purified using anion-exchange chromatography with a HiTrap Q column (GE Healthcare, USA). We removed the hexahistidine tag using TEV protease, and further purified the proteins using HiTrap Q anion-exchange chromatography. Additionally, we carried out size-exclusion chromatography. The proteins were concentrated using Vivaspin (30 kDa molecular-weight cutoff; Millipore, USA) in buffer containing 20 mM Tris-HCl (pH 8.5), 300 mM NaCl, 5% [v/v] glycerol, and 2 mM 2-mercaptoethanol. The final protein concentration was 12 mg/ml.

After TALON affinity chromatography, FL MexT was directly loaded onto a size-exclusion chromatography column (Hi-Load Superdex 16/60 200; GE Healthcare) equilibrated with 20 mM Tris-HCl (pH 8.5), 500 mM NaCl, 10% (v/v) glycerol, and 2 mM 2-mercaptoethanol. The purified proteins were aliquoted and stored frozen at −173°C.

Crystallization and data collection

MexT RD was crystallized using the hanging-drop vapor-diffusion method at 14°C as described previously (Kim et al., 2018). The precipitation solution comprised 0.5 M ammonium sulfate, 1.0 M sodium citrate tribasic dehydrate (pH 5.6), 0.1 M lithium sulfate, and 2 mM ethylenediaminetetraacetic acid (EDTA). Glycerol (30% [v/v], cryoprotectant) was added to the precipitation solution. The X-ray diffraction data were collected in a flash-cooled liquid nitrogen stream at −173°C. SeMet-MexT RD was crystallized under the same crystallization conditions as for native MexT RD. The single-wavelength anomalous diffraction (SAD) dataset was collected from beamline 5C at the Pohang Accelerator Laboratory (Korea) and processed using HKL2000 software (Otwinowski and Minor, 1997).

Structural analysis of MexT RD

The 2.0 Å resolution dataset of the native crystal belongs to space group P2_1_2_1, with unit-cell dimensions of a = 65.7 Å, b = 108.7 Å, and c = 109.2 Å. The initial model was built based on phase information from the SAD dataset for SeMet-substituted protein at 2.3 Å resolution using AutoSol software (Terwilliger et al., 2009). The structure was refined against the 2.0-Å-resolution native dataset using COOT and Phenix.refine softwares (Adams et al., 2010; Emsley et al., 2010).

Size-exclusion chromatography with multi-angle light scattering (SEC-MALS)

The sizes of FL MexT and MexT RD were assayed using SEC-MALS. A high-performance liquid chromatography pump (Agilent, USA) was connected to a Superdex-200 10/300 GL gel filtration column (GE Healthcare) and a MALS instrument (Wyatt Dawn Heleos, USA). The size-exclusion chromatography column was pre-equilibrated with buffer comprising 20 mM Tris-HCl (pH 8.5), 500 mM NaCl, and 2 mM 2-mercaptoethanol for FL MexT; and 20 mM Tris-HCl (pH 8.5), 300 mM NaCl, and 2 mM 2-mercaptoethanol for MexT RD. Bovine serum albumin (2 mg/ml) was used as the standard. MexT RD and FL MexT samples (2 mg/ml) were injected onto the column and eluted at a flow rate of 0.2 ml/min. The datasets were evaluated using the Debye model for fitting static light-scattering data, and refractive index peaks were presented in EASI graphs created using Astra V software (Wyatt Dawn Heleos).

DNase I footprinting assay

A DNA fragment of 630 bp (the mexT-mexT intergenic region [230 bp] + 200 bp upstream + 200 bp downstream) was amplified using polymerase chain reaction (PCR) from P. aeruginosa chromosomal DNA. For this, we used the following primers: 5'-6-fluorescein amidite-GGCAGGGACACATCAGCCACCCTGC-3' and 5'-GGCGCAGCTCCACCAGCTCCGGGGGCC-3'. The PCR product was purified from agarose gel electro-
phoresis. We mixed 400 ng of DNA and 200 ng of purified FL MexT for 30 min at room temperature. Next, 0.04 units of DNase I (Fermentas) was added, and the reaction mixture (10 mM Tris [pH 8.0], 50 mM KCl, 8 mM MgCl₂, 50 ng/µl BSA, 5% glycerol) was incubated for 2 min at room temperature. The reactions were terminated by adding an equal volume of stopping buffer (200 mM NaCl, 1% sodium dodecyl sulfate, 30 mM EDTA) and DNase I was inactivated by heating at 75°C for 10 min. The DNA fragments were recovered via phenol extraction and alcohol precipitation and fragment-sequences were analyzed using an ABI3730xl DNA Analyzer (Applied Biosystems, USA). The size distribution was scanned using Peak Scanner software (ver. 1.0; Applied Biosystems).

RESULTS AND DISCUSSION

Overall structure of the MexT regulatory domain

The FL MexT (residues 1-304) and its RD (residues 95-304) were produced in E. coli and purified until homogeneity was attained. FL MexT and MexT RD were homotetrameric and homodimeric in solution, respectively, as determined using SEC-MALS (Fig. 1).

To obtain phase information, we acquired the diffraction dataset using the crystal of the SeMet-labeled protein, which was crystallized in the presence of a high concentration of ammonium sulfate. The structure was determined via the SAD method using the anomalous signals from SeMet, which was refined to the 2.0-Å-resolution native dataset with an R factor of 0.21 and Rfree of 0.25 (Table 1). The obtained crystals belonged to the spacegroup P2₁2₁2₁, and the asymmetric unit contained four protomers or two homodimers (Fig. 2).

Its overall structure shows the features, typical of the RDs of LysR-type transcriptional regulators (LTTRs). The extensive dimeric interaction in the crystal structure is consistent with the results of SEC-MALS (Fig. 1).

Structure of RD protomers

MexT RD protomers are composed of 10 β-strands and 7 α-helices, as in the typical LTTR RD (Jo et al., 2015). Each protomer can be divided into two subdomains—RD-I (residues 95-172, 274-304) and RD-II (residues 173-273). The folding pattern of the two subdomains was similar to Rossmann-fold topology; i.e., a central β-sheet bound by helices and loops. The RD-I subdomain comprised three α helices (α1, α2, and α7) and five β-strands (β1, β2, β3, β4, and β10), whereas the RD-II subdomain comprised four α helices (α3, α4, α5, and α6) and five β-strands (β5, β6, β7, β8, and β9). The RD-I and RD-II subdomains were connected by two loops—one connects β4 of RD-I to β5 of RD-II, and the other connects β9 of RD-II to β10 of RD-I (Fig. 2).

Table 1. Statistics for X-ray data collection and refinement

|                  | Native MexT RD | Se-Met MexT RD |
|------------------|----------------|----------------|
| Data collection  | PAL 5C         | PAL 5C         |
| Wavelength (Å)   | 0.97960        | 0.97940        |
| Space group      | P2₁2₁2₁        | P2₁2₁2₁        |
| Cell dimensions  | a = 65.7 Å, b = 108.7 Å, c = 109.2 Å | a = 65.9 Å, b = 108.7 Å, c = 109.3 Å |
| α, β, γ (°)      | 90, 90, 90     | 90, 90, 90     |
| Resolution (Å)   | 50-2.00 (2.03-2.00) | 50-2.30 (2.34-2.30) |
| Rmerge           | 0.042 (0.50)   | 0.10 (0.83)    |
| Rmerge/σ(I)      | 0.210.25       | 0.210.25       |
| No. of reflections | 50,630        | 6,365          |
| Wilson B-factor  | 22.18          | 23.76 (2.91)   |
| RMSD              | 0.003          | 0.67           |
| Ramachandran plot | Favor (°) 97.0 | Allowed (%) 3.0 |
|                  | Outliers (%)   | 0.0            |
| PDB ID           | 6L33           | 6L33           |

RMSD, root mean square deviation.

Table 1. Statistics for X-ray data collection and refinement

*Values in parentheses are for the highest-resolution shell.

Fig. 1. Molecular size of the purified proteins. (A) FL MexT. (B) MexT RD. Primary γ-axis, molar mass determined using multi-angle light scattering (MALS; black line); secondary γ-axis, concentration. Protein concentration was assessed by measuring the absorbance at 280 nm (red), light scattering (LS; green), and the refractive index (RI; purple). X-axis, elution time from size exclusion chromatography.
The four protomers (chains A-D) in the asymmetric unit are arranged as two of dimers (chains A:B and C:D; Fig. 2). The overall arrangement of the MexT RD dimer was similar to those of other LTTRs. Within crisscrossing protomer dimer, RD-I of one protomer directly interacted with RD-II of the other protomer. The major interactions of two protomers were intermolecular hydrogen bonds between residues in the RD-I
β backbone and the RD-II β7 backbone at the dimeric interface. Hydrophobic interactions were found at the dimeric interface of RD-1 α1 and RD-II α5, and likely stabilize the dimer (Supplementary Fig. S1).

The asymmetric unit contained five sulfate ions, which may originate from the crystallization solution (Fig. 3 and Supplementary Fig. S2). Of the five sulfate ions, three were found at the dimeric interfaces; the sulfate ions interacted with Arg134 and Arg135 near the molecular pseudo-2-fold axis—one in the A:B chain dimer and two in the C:D chain dimer (Fig. 3). The fourth sulfate ion is found in the putative ligand-binding site (Fig. 4A). The fifth sulfate ion is located outside of the A:B chain dimer, interacting with Arg120 and Arg121 of chain A (Supplementary Fig. S2).

The dimeric arrangements of the two dimers were similar (Supplementary Fig. S3), but the relative position of subdomain RD-II within the overall frame showed ~5.0 Å deviation. The orientation of the protomers in the dimer can be used to distinguish active from inactive LTTRs (Choi et al., 2001; Jo et al., 2015). To examine the conformation of the RD dimer, the obtained MexT RD dimer (chains A:B) was superimposed onto the E. coli OxyR RD dimer in the reduced (inactive; PDB code, 1A69) and oxidized (active; PDB code, 1I6A) forms as a prototype LTTR (Choi et al., 2001). The conformation of the MexT RD dimer fit better to that of the oxidized OxyR RD (Fig. 3C and 3D). However, further studies are required to confirm this idea.

**Putative ligand-binding site**

Compared to a typical LTTR E. coli OxyR, MexT exhibits substantial variation in the conformation of the putative ligand-binding sites at the inter-subdomain junction. These structural variations of MexT appear to be beyond the subtle superficial conformational changes resulting from crystal packing contacts. We noted a loop connecting β6 and α4 in RD-II (residues 199-208), which we named the variable (V)-loop (Fig. 4A). A V-loop with two different conformations lines the putative ligand-binding sites, which are not involved in the crystal packing contacts between the symmetry-related molecules (Supplementary Fig. S4). In chains A and B, Leu205 of the V-loop forms a hydrophobic bond with Val222 of RD-II, and with Val130 and Val132 of RD-I of the other protomer, leading to the formation of a cavity at the putative ligand-binding site (Fig. 4A; chains A and B). In chain C, Leu205 in the loop forms hydrophobic bonds with Tyr156 and Tyr138 of RD-I but has no inter-protomer interaction (Fig. 4A; chain C). The electron density of the V-loop of chain D is low and disordered (residues 204-217). However, the conformation of the V-loop of chain D was similar to that of chain C (Fig. 4A; chain D). Thus, our findings suggest that the V-loop regulates the putative ligand-binding sites.

As mentioned above, one sulfate ion is located at the putative ligand-binding site (Fig. 3A). Arg220 and Arg269 ionically interact with the sulfate ion at the putative ligand-binding site of chain B (Fig. 3A). In particular, guanidine group of Arg269 in chain B can make two configurations by flipping. And it is the site where one sulfate ion can bind. Pocket formation by the flipping guanidine group makes space for holding small molecules. The sulfate-binding region extends to the hydrophobic surface region decorated by Tyr138, Phe201, Phe208, and Phe229 (Fig. 4B and Supplementary Fig. S5). Highlighting the space of adjacent sulfate-binding region and hydrophobic patches in the ligand-binding site, our structure suggests that candidate ligands should have a negatively charged (or polar) moiety linked to a hydrophobic moiety.

Cinnamaldehyde upregulated the expression of MexEF-OprN. Indeed, cinnamaldehyde has a polar aldehyde moiety and a hydrophobic phenyl ring (Juarez et al., 2017). The polar aldehyde moiety likely interacts with the positively charged pocket, which is composed of Arg220 and Arg269, and the hydrophobic phenyl ring probably participates in the interaction. A molecular model showed that cinnamaldehyde fitted the pocket well (Dallakyan and Olson, 2015) (Supplementary Fig. S6). Molecules with such characteristics may activate MexT and alter its association with DNA.
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Binding of MexT to the promoter region of mexEF-oprN
To elucidate the interaction between MexT with the promoter of mexEF-oprN, a DNase I footprinting assay was performed focusing on the region between mexT and mexE, which spans the promoter region of the mexEF-oprN operon. According to the short fragment DNA size analysis data, MexT binding protected two parts of the DNA sequence (MexT binding sites I and II) that are 17 bp apart. MexT binding site I is located between –112 bp and –89 bp, and the MexT binding site II is between –72 and –58 bp from the transcriptional initiation site of the mexEF-oprN operon, which is located 30 bp upstream of the mexE start codon (Maseda et al., 2010) (Fig. 5).

Next, the two MexT-binding site sequences identified by DNase I footprinting were aligned with the promoter sequences of other genes in the MexT regulon (PA1744, PA1970, PA2486, and PA2759). The MexT regulon was identified with microarray-based transcriptome profiling (Tian et al., 2009), and was further experimentally confirmed by electromobility shift assays and β-galactosidase assays using transcriptional fusion reporters (Maseda et al., 2010; Tian et al., 2009). The sequence alignment showed a palindromic consensus pattern of ATCA(N7)CGAT at each MexT-binding site. Notably, there was partial overlap between the front region ATCA(N5)GTCGAT(N2)ACYAT and the nod box sequence ATC(N5)GAT, indicating a region of degeneracy (marked as N2) in the binding motif bracketed by ATC and GAT. While the sequence ATCA(N7)CGAT is the binding motif of the DNA-binding domain (DBD) dimer of LTTR, our findings show that the DBD dimer of MexT facilitates close contact between the above two regions. Therefore, it is not surprising that the tetrameric conformation of MexT brings the DNA region (55 bp) spanning the two MexT-binding sites into close association.

Since recombinant MexT was not tested with candidate ligands, its binding to MexT binding sites I and II likely represents its inactive status as a mexEF-oprN promoter. The effect of cognate ligand binding on protein–DNA binding is unclear. Importantly, the two MexT consensus sequence positions form an interrupted palindromic sequence compared to those seen in other LTTRs (Maddocks and Oyston, 2008; Muraoka et al., 2003a; 2003b).

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this study.

In conclusion, MexT is a LTTR (Kohler et al., 1997; 1999). The typical tetrameric arrangement of OxyR (Jo et al., 2015) and the atypical tetrameric arrangement of VV2_1132 and HypT have been characterized previously (Jang et al., 2018; Jo et al., 2019). Recognition of ligand binding or stimuli by the RD induces structural changes in transcriptional regulators, which alters their DNA binding (Jo et al., 2015). We elucidated the crystal structure of the RD of MexT of *P. aeruginosa*. Since the RD should contain the region required for sensing the signals in response to the environmental cues to turn on the efflux pumps, our study is important in understanding the molecular nature of the gene expression regulation required for the efflux pump-mediated antibacterial resistance of *P. aeruginosa*. Furthermore, our findings will facilitate the development of drugs with activity against *P. aeruginosa* by promoting the expression of multidrug efflux pumps.

**Note:** Supplementary information is available on the Molecules and Cells website (www.molcells.org).

**Disclosure**

The authors have no potential conflicts of interest to disclose.

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