The Mannitol Operon Repressor MtlR Belongs to a New Class of Transcription Regulators in Bacteria

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SUMMARY

Many bacteria express phosphoenolpyruvate-dependent phosphotransferase systems (PTS). The mannitol-specific PTS catalyze the uptake and phosphorylation of D-mannitol. The uptake system comprises several genes encoded in the single operon. The expression of the mannitol operon is regulated by a proposed transcriptional factor, mannitol operon repressor (MtlR) that was first studied in *Escherichia coli*. Here we report the first crystal structures of MtlR from *Vibrio parahaemolyticus* (Vp-MtlR) and its homolog YggD protein from *Shigella flexneri* (Sf-YggD). MtlR and YggD belong to the same protein family (Pfam05068). Although Vp-MtlR and Sf-YggD share low sequence identity (22%), their overall structures are very similar, representing a novel all α-helical fold and indicate similar function. However, their lacking of any known DNA-binding structural motifs and their unfavorable electrostatic properties imply that MtlR/YggD unlikely bind a specific DNA operator directly as proposed earlier. This structural observation is further corroborated by *in vitro* DNA-binding studies of *E. coli* MtlR (Ec-MtlR), which detected no interaction of Ec-MtlR with the well-characterized mannitol operator/promoter region. Therefore MtlR/YggD belongs to a new class of transcription factors in bacteria that may regulate gene expression indirectly as a part of a larger transcriptional complex.
INTRODUCTION

Phosphoenolpyruvate-(PEP)-dependent phosphotransferase systems (PTS) primarily consist of a transmembrane transporter and enzymes responsible for phosphoryl group transfer from PEP to a transporter-bound sugar acceptor (1,2). The PTS are generally substrate-defined and their components may vary across species (2,3). D-Mannitol, or 1,2,3,4,5,6-hexanohexol, is a polyol formed by reduction of mannose or fructose. It is one of the hexitols involved in bacteria catabolism pathways where it serves as a source of fermentable sugar (4,5). The D-mannitol specific PTS was first discovered and sequenced in *Escherichia coli* (6-8). With the consideration that the system was regulated like other hexitol PTS systems, an open reading frame was sought and identified within the mannitol operon (9). Experiments indicated that the loss of the gene (*mtlR*) led to the constitutive expression of the operon, and its gene product MtlR was proposed to be a transcription factor: mannitol operon repressor (MtlR, COG2213)(9). Two putative DNA operator palindromes that might serve as MltR binding sites were identified within the operator-promoter region.

Besides MtlR, a typical mannitol operon also encodes a mannitol-specific PTS system ABC transporter II component (MtlA, COG2213) and a mannitol-1-phosphate 5-dehydrogenase (MtlD, COG0246). The mannitol operon is conserved and has been cloned from many Gram-negative bacterial families, such as *Shigella* (10), *Salmonella* (11), *Yersinia* (12), *Klebsiella* (13,14) and *Vibrio* (15). For example, in the genome sequence of *Vibrio parahaemolyticus* RIMD 2210633(15), a typical mannitol operon, including MtlR, MtlA and MtlD, has been identified on chromosome I. A paralogue gene *yggD* annotated as putative transcriptional regulator was found clustered with *cmta* and *cmtB* on chromosome II of *V. parahaemolyticus*. The *cmta* and *cmtB* genes seem to encode the equivalents of MtlA components according to a recent study (16).

In genomes of many Gram-negative bacteria, a gene named *yggD*, encoding a sequence homolog of MtlR has also been identified. The gene *yggD* is not clustered with *mtlA* and *mtlD* as *mtlR* in a typical mannitol operon. Its gene neighbors vary considerably among organisms, even among strains, and its position provides little insight into its function. In the complete genome sequence of *Shigella flexneri* serotype 2a strain 2457T (10), for example, the *yggD* gene is located about 1 Mbp away from the mannitol operon on the same strand of DNA(10). Its neighboring genes include uncharacterized *yggC* and *yggF*. The latter may encode a fructose-1,6-bisphosphatase II-like protein(10). The function of YggD has never been described. However, based on its sequence similarity to MtlR, it has been assigned to the same protein family, Pfam05068 (mannitol operon repressor) (17).

The protein structural information of mannitol operon components has been limited to the IIA domain of MtlA from *E. coli* (18) and *Salmonella typhimurium* LT2 (PDB code:1XIZ). The enzyme IIA consists of a central β-sheet flanked by α-helices on both sides, which is remarkably different from IIA proteins that belong to other three EIIs groups of PTS (18). To understand the structure, molecular mechanism and regulation of the mannitol operon, we have expressed and purified a number of MtlR and YggD from different bacterial species. Several expression vectors were used to improve protein solubility and reductive methylation was applied to augment crystallization (19).

Here we present the crystal structures of MtlR from *V. parahaemolyticus* (Vp-MtlR) and YggD from *S. flexneri* (Sf-YggD), the first structures of the mannitol operon repressor family. The structures of Vp-MtlR and Sf-YggD were determined at 2.75 Å and 2.50 Å resolution respectively and represent a novel all α-helical fold. Though Vp-MtlR and Sf-YggD share low sequence identity, they are structurally very similar.
and also form similar dimers. The analysis of their structural features and surface electrostatic properties in combination with DNA-binding studies of Ec-MtlR suggests that MtlR/YggD are unlikely to bind DNA directly. We also provide direct biochemical evidence that the MtlR does not bind the regulatory region of the \textit{E. coli} mannitol operon. Therefore, MtlR seems to belong to a new class of transcription factors in bacteria and may regulate gene expression indirectly perhaps as a part of larger transcriptional complex.

**EXPERIMENTAL PROCEDURES**

**Protein cloning, expression and purification**

A truncated on C-terminus \textit{mtlR} gene from \textit{V. parahaemolyticus} RIMD 2210633 was cloned into pMCSG7 vector (20). The gene codes Vp-MtlR without four C-terminal residues, D173SPF. The protocols for the expression in \textit{E. coli} BL21 (DE3) and purification of the Se-Met-labeled recombinant protein were reported previously (21). The Vp-MtlR expressed well with final protein yield of about 195 mg/L culture.

The full-length \textit{yggD} gene from \textit{S. flexneri} 2a 2457T was initially cloned in pMCSG7 vector. The solubility of Sf-YggD was very low with soluble protein yield approximately 5 mg/L culture. To improve the protein production, the pMCSG19 expression vector was utilized (22), which bears a fusion construct with maltose binding protein (MBP) followed by TVMV protease cleavage site, His\textsubscript{6}-tag and TEV protease cleavage site fused into N-terminus of the target protein. The expression and purification protocols of Se-Met-labeled recombinant Sf-YggD were the same as for Vp-MtlR (21). With the use of pMCSG19 vector, the Sf-YggD yield increased to 62 mg/L culture.

The expression constructs for Ec-MtlR, FruR1 (coding for the first 57 amino acids of fructose repressor (FruR)), and FruR2 (the full length protein) were cloned into the NdeI and XhoI restrictions sites of pET28. The resulting constructs consisting of an N-terminal His\textsubscript{6}-tag and a thrombin cleavage site were expressed in BL21(DE3) and purified as reported earlier (23). The gene for full-length Ec-MtlR was also cloned in pMCSG7 vector. The protein was expressed and purified as reported previously (21).

**Reductive methylation of Sf-YggD protein**

A reductive methylation of lysine residues of Sf-YggD was performed to improve protein crystallization properties (19). During reductive methylation, the protein is modified through addition of two methyl groups to Lys residues. The concentration of the protein was first adjusted to about 10 mg/ml. Dimethylamine-borane complex (ABC, 20 µL of 1M stock) and formaldehyde (40 µL of 1M stock) were added per 1 mL of the protein solution. The mixture was incubated for 2 hours at 4°C. The procedure was then repeated for the second time. After that, only ABC (10 µL per 1 mL protein solution) was added and the mixture was incubated overnight. On the second day, 1 mL of 5 mg/mL glycine and 1M DTT to a final concentration of 5 mM were added and the reaction mixture was incubated on ice for 2 hours. The protein was finally dialyzed against crystallization buffer and concentrated.

**Limited proteolysis of Ec-MtlR**

Ec-MtlR was first treated with trypsin and chymotrypsin and separated on a SDS-PAGE gel, which shows apparent reduction of molecular mass. The digest was then redone with 100µg of protein, stopped after 6 hours with 1 mM PMSF and analyzed by mass spectrometry. The mass of 20,359.6 Da corresponds to a fragment of Ec-MtlR truncated of the first 14 N-terminal amino acids (theoretical mass 20,351 Da).

**Size exclusion chromatography**

Size exclusion chromatography was performed on a Superdex-200 10/300 GL column (General Electric Health Systems) using AKTA\textsuperscript{TM} Xpress. The column was pre-equilibrated with crystallization buffer (20 mM HEPES pH 8.0, 250 mM NaCl, 2 mM DTT) and calibrated with premixed protein standards, including ribonuclease A (13.7 kDa), chymotrypsinogen A (25 kDa),
ovalbumin (43 kDa), albumin (67 kDa), aldolase (158 kDa), catalase (232 kDa), and Blue Dextran (2,000 kDa). For each run, a 5 ml protein sample at a 10 mg/ml concentration was injected into the column. The chromatography was carried out at 4°C at a flow rate of 2 mL/min. The calibration curve of K\textsubscript{av} versus log molecular weight was prepared using the equation K\textsubscript{av} = V\textsubscript{e} – V\textsubscript{o}/V\textsubscript{t} – V\textsubscript{o}, where V\textsubscript{e} = elution volume for the protein, V\textsubscript{o} = column void volume, and V\textsubscript{t} = total bed volume.

**Protein crystallization**

The Se-Met labeled Vp-MtlR, unmethylated and methylated Sf-YggD proteins were screened for crystallization conditions with the help of the Mosquito robot (TTP Labtech) using the sitting drop vapor diffusing technique in 96-well CrystalQuick plate (Greiner). For each condition, 0.4 µl of protein (79 mg/ml for Vp-MtlR and 66 mg/ml for Sf-YggD) and 0.4 µl of crystallization formulation were mixed; the mixture was equilibrated against 135 µl of the reservoir in the well. Crystallization screens used were Index (Hampton Research), Wizard (Emerald Biosystems), and SM4 (Nextal) at both 4°C and 18°C. Diffraction quality crystals of Vp-MtlR appeared under the condition of 0.2M tri-ammonium citrate, pH7.0, 20%(w/v) PEG3350. The crystallization condition of methylated Sf-YggD crystals was 0.1M sodium acetate, 0.1M MES pH6.5, 30%(w/v) PEG2000MME. Prior to data collection, crystals were treated with cryoprotectant complemented from crystallization buffer. For Vp-MtlR, 25%(v/v) glycerol was added. For methylated Sf-YggD, 20% (v/v) glycerol was added while and PEG2000 MME was reduced to 20%. No crystal was produced from unmethylated Sf-YggD protein. Extensive screenings of crystallization conditions for Ec-MtlR and its proteolytic fragment were also performed and no crystal was obtained.

**DNA-binding experiments of Ec-MtlR**

**EMSA (electrophoretic mobility shift assay):** 80 fmol of 5'-32P-labeled DNA-oligonucleotides were incubated with a 4 to 10 fold excess of protein in TMD-buffer (20 mM Tris-HCl, pH 8.0, 5 mM MgCl\textsubscript{2}, 1 mM DTT). BSA was added in a ratio of 100:1 to the protein under investigation to prevent unspecific binding. The samples were incubated for 30 min and separated on a vertical polyacrylamide gel. An X-ray film was exposed for 16 hours at -70°C. Additional EMSA was carried out on a PhastSystem (GE) using PhastGel Gradient 4-15% gels (GE), see Supplementary Data.

**Surface plasmon resonance:** An IASys single channel resonant mirror biosensor (Fisons Applied Sensor Technology, Cambridge, U.K.) (George, 1995, Rubio, 1997) was used to measure the affinity of Ec-MtlR constructs to DNA. The 5´-biotin labeled oligonucleotides mtlr1_for (5´-CTCGGGCTTCCAGCCTGCG) and mtlr2_for (5´-ACATAAGAAGGGGTGTTTTATGT) and their complementary strands were obtained from Microsynth, dissolved and annealed according to the standard procedure. A 100-200 µg excess of streptavidin (Pierce, Illinois, U.S.A.) was coupled to a sensing cuvette manufactured with a biotin layer and the biotinylated dsDNA was then coupled to the immobilized streptavidin. All binding reactions were performed in PBS, pH 7.2, 25% glycerol and a range of additional NaCl concentrations between 100 mM and 300 mM were tested to optimize the binding.

**A SELEX-based method for measuring DNA-binding specificities:** A SELEX based binding-site selection assay was developed for the identification of protein-DNA binding specificities. Wells of Nunc-Immunoplates were coated with 400 µl of 1 mg/ml concentrations of the proteins under test and incubated for between 1 and 8 h. The wells were washed with Tris-HCl buffer (pH 7.4) prior to the binding reaction. 50 µl of binding reaction buffer containing 200 ng Poly(dIdC)-Poly (dIdC) and 0.2 ng of the random oligonucleotide R76 were added to each test well. The oligonucleotide R76 comprised a random 26 base sequence flanked by two 25 nucleotide PCR primers as follows: 5´-
CAGGTCAGTTCAGCGGATCCTGTCG(A/G/C/T)26GAGGCGAATTC-AGTGCAACTGCAGC.

Plates were incubated for 4 h with shaking. After the binding reaction, wells were washed thoroughly with buffer E and the bound DNA eluted with 250 μl of 500 mM NaCl. 2 μl aliquots of the eluate were amplified by PCR. The amplified DNA was then rebound to the protein from which it was eluted and the whole binding, elution and amplification procedure was repeated 5 times to facilitate greater accuracy in the selection of DNA sequences. The final amplification of the selected DNA sequences was performed using Eppendorf Taq polymerase, which produces 3'-A overhangs and facilitates cloning into the pGEMT Easy vector system (Promega). The blue/white screening was used to select clones containing inserts and the selected DNA was sequenced.

X-ray diffraction data collection

Diffraction data were collected at 100 K at the 19ID and 19BM beamlines of the Structural Biology Center at the Advanced Photon Source at Argonne National Laboratory using the program SBCcollect. A multiple-wavelength anomalous diffraction (MAD) data set and a single-wavelength anomalous diffraction (SAD) data set were collected at the wavelengths near selenium absorption peaks from Se-Met labeled protein crystals of Vp-MtlR and Sf-YggD, respectively. All data were processed and scaled with HKL3000 suite(24) (Table 1).

Structure determination and refinement

The Vp-MtlR and Sf-YggD crystal structures were determined using MAD and SAD methods, respectively. For both structures, Se sites were first located using the program SHELXD(25) and they were used for initial phasing with the program MLPHARE(26). After density modification (DM)(26), partial models (Vp-MtlR:378 residues with 26 side chains placed; Sf-YggD: 392 residues with 17 side chains placed) were obtained from automatic model building trials using the program RESOLVE(27). All the above programs are integrated within the program suite HKL3000(24).

After cycles of model buildings using the program COOT(28), the models were refined using the program REFMAC(29) (Table 1). After final refinements, electron density calculated at 1.0 σ is well connected for main chains of three Vp-MtlR monomers. There were density breaks between M127 and D138 for the fourth Vp-MtlR monomer. In case of Sf-YggD, electron density breaks were observed only at the loop region from residue F137 to residue K142 in three out of four monomers. Electron densities were observed for methyl groups of 22 out 32 lysine residues in the methylated Sf-YggD and these methyl groups were built into the structural model.
RESULTS

Expression, purification, crystallization and structure determination

The recombinant proteins Vp-MtlR and Sf-YggD were over-expressed in the E. coli BL21(DE3) strain and purified to homogeneity using His6-tag specific immobilized metal affinity chromatography (IMAC) columns as described previously(21). Sf-YggD was further subjected to reductive methylation of lysine residues to improve crystallization(19). The X-ray diffraction data of Vp-MtlR (2.75Å) and Sf-YggD (2.50Å) were both collected near the selenium absorption K-edge of their Se-Met-labeled protein crystals (Table I) with their structures solved by MAD and SAD techniques, respectively (Table I). The Ec-MtlR and its proteolytic fragment failed to crystallize.

Structures of Vp-MtlR and Sf-YggD Monomers

There are four monomers in one asymmetric unit of both Vp-MtlR and Sf-YggD crystals (Figs. 1A and 2A). Though different in their molecular packing, the structures of the MtlR and the YggD monomers are highly similar. Each monomer assumes an all α-helical fold rather than the predicted α/β structure of E. coli MtlR(9). The Sf-YggD monomer has nine α-helices arranged in three layers (Fig. 2). In Vp-MtlR the α8 helix is replaced by a flexible loop (Figs. 1B and 2C). Except the short α7 helix within the loop between α6 and α8, all helices that are adjacent in amino acid sequence are anti-parallel. The first layer includes α4, α5, α6 and α8 as well as the short helix α7. The four helices α4, α5, α6 and α8 are anti-parallel to each other with a relative rotation about 20° to each other. The second layer or the central helical sheet includes α2, α3 and α9. The α2 and α9 helices are parallel to each other. They are the longest and the most prominent helices of the structure. The second layer has a rotation of about 50° relative to the first layer. The packing between two adjacent individual α-helices within a helical layer or between two helical layers are largely representative of two typical helix-helix packing modes with the ridges of one helix fitting into grooves of the other and vice versa(30). A single N-terminal helix (α1) is stacked onto the second layer, interacting with α2 and α9. The interactions between these three helices are similar to what observed in a coiled-coil helical structure though there are only two layers of hydrophobic interactions in both Vp-MtlR and Sf-YggD. Furthermore, the majority of interhelical interactions in both monomers are predominantly hydrophobic in nature, suggesting the high stability of this all α-helical fold.

There is no significant conformational variation between four MtlR monomers in Vp-MtlR. A pairwise superposition of four MtlR monomers results in rmsd (root mean square deviation) values within a range of 0.66 to 1.20 Å. The primary variable region is the loop between α7 and α9, which has poorly defined electron density in three out of four monomers. The loop also influences conformation changes in the N-terminal region of the α9 helix. A similar pairwise superposition of four YggD monomers results in much smaller rmsd values (0.40 to 0.61 Å). The only variation was found in the disordered or partially disordered loop region between α8 and α9. Multiple sequence alignment (Fig. 3) suggests that the region between α7 and α9 is the most variable part in MtlR and YggD. A typical superimposition of MtlR and YggD monomers gives an rmsd of 1.60 Å with 140 residues from each monomer aligned (Fig. 2C). The primary difference between the two structures is the replacement of the α8 helix in Sf-YggD by a flexible loop in Vp-MtlR as discussed earlier. In addition, the α3 helix shows a small conformational variation; α3 is an edge helix and is involved in dimerization of MtlR and YggD.

MtlR and YggD Dimers and Tetramers

Analysis of crystal packing and interactions across monomer/monomer interfaces suggests that the most likely biological unit of Vp-MtlR and Sf-YggD is a dimer. In Vp-MtlR, monomers A and B, and monomers C and D form two dimers (AB-
dimer and CD-dimer) related through pseudo 2-fold axis with total buried surface areas of 1585 Å² and 1554 Å², respectively (Fig. 1). In Sf-YggD, monomers B and C form a similar BC-dimer with a total buried surface area of 1735 Å² (Figs. 2A and B) while monomer A and monomer D form dimers with their symmetry-related mates, respectively. The resulting buried surface areas of these two symmetric dimers are 1764 Å² and 1783 Å², respectively. These values are consistent with the "standard-size" protein interfaces in which the total area buried by the components in the protein/protein interface is 1600 (±400) Å² (31).

Both MtlR and YggD dimers are formed by the contacts mainly between the α2, α3 and α4 helices of two monomers (Figs. 1B and 2B) with α3 and α4 providing the most important hydrophobic interactions. As a result of the dimerization, an extended five-layered helical structure forms with the central layer contributed primarily by 6 (MtlR) or 8 (YggD) α-helices (Figs. 1B and 2B). The dimerization interface is highly hydrophobic and the residues involved in the interactions across the interface include I42(MtlR)/L41(YggD), F43/F42, V44/V43, A49/I49, V53/A53, V54/V54, L57/L57, P63/P63, V69/V69, K72/R72, L73/L73, F75/Y75, G76/A76, L77/L77 and Y85/Y85 (Fig. 4A). A total of 8 aromatic residues from two monomers are involved in the interaction across the interface in each case. It is remarkable that these residues contributing to the dimer interface are either identical (69%) or highly conserved (31%) in the family though the sequence identity between Vp-MtlR and Sf-YggD is only 22% (Fig. 3). Multiple sequence alignment shows the region including α3 and α4 helices being the most conserved across different species. Additionally, neither specific hydrogen bond nor salt bridge is observed across dimer interface.

In the Sf-YggD crystal, all contacts between dimers seem insignificant suggesting that no higher oligomers can be formed. The observation agrees with the results of size-exclusion chromatography. The elution profiles of Sf-YggD showed a single major peak (Fig. 5A) with apparent molecular weight of 47.8 kDa (data not shown). The calculated molecular weight of the Sf-YggD monomer from its amino acid sequence is 19.5 kDa (including 3 N-terminal vector derived residues, SNA) and estimated molecular weights for the dimer and trimer are 39.0 and 58.5 kDa respectively. Therefore it appears that Sf-YggD is a dimer in the solution. Its slightly larger apparent molecular weight obtained from size-exclusion chromatography might be a result of its elongated shape.

The calculated molecular weight of the Vp-MtlR monomer is 19.7 kDa and is very similar to Sf-YggD. However, its elution volume from size-exclusion chromatography was significantly different to that of Sf-YggD and its peak profile was much broader (Fig. 5A). The Vp-MtlR molecular weight was estimated to be 90.4 kDa, more than 4 times that of the Vp-MtlR monomer (78.8 kDa). Therefore it appears that Vp-MtlR can form a tetramer, possibly through the interaction between the AB-dimer and the CD-dimer as shown in our structure (Fig. 1A). In fact, the two Vp-MtlR dimers interact much more extensively than in case of Sf-YggD, resulting a total buried interface of area as high as 2135 Å². The dimer-dimer contact is also the primary contact between any two dimers in the crystal structure of Vp-MtlR. Across the dimer-dimer interface, the flexible loops between α7 and α9 form two short anti-parallel strands with five main chain-main chain hydrogen bonds (Fig. 4B). In addition to these contacts there are also hydrogen bonds contributed by side chains across the interface and a number of hydrophobic interactions (Fig. 4B). However, the tetramer can’t be very stable because a tetramer with the AB-dimer/CD-dimer interactions would lead to polymerization of Vp-MtlR dimers as is observed in the crystal. In solution, at high protein concentration, the tetramer is likely to exist as a well as other oligomeric forms of Vp-MtlR in an equilibrium with a dimer. This may explain the broadening of elution peak of Vp-MtlR and its shift towards high molecular weight (Fig. 5A).

Similar results were obtained for the Ec-MtlR (unpublished data). The results from size
exclusion chromatography, native PAGE, dynamic light scattering and analytical ultracentrifugation showed a polydisperse state of MtlR with multimers formed in a concentration-dependent manner. This polydispersity may be the primary reason why Ec-MtlR fails our crystallization trials. Our data show that the MtlR (as well as YggD) dimer is a basic biological unit but proteins in different species and under different conditions can form higher level aggregates.

**MtlR/YggD Represent a Unique Protein Fold**

A structural similarity search using the DALI server (www.ebi.ac.uk/dali) (32) resulted in no significant hits. The 4 top structural homologues from a search by using Sf-YggD as a search template, for example, generally represent a 4-helix bundle with one additional short helical turn and one 3-turn α-helix with Z score: 5.3-6.7, rmsd: 3.4-4.3, number of aligned residues: 96-103 and sequence identity: 5-11%. They include proteins of unknown function from bacteria and the C-terminal domain of kanamycin nucleotidyltransferase from *Haemophilus influenzae* (KNTase) (PDB: 1KNY). A part of the C-terminal domain of KNTase involves nucleotide binding (33). Although part of the MtlR structure can be aligned with the C-terminal domain of KNTase-like proteins, the 8- or 9-helix layered MtlR and YggD monomers represent a novel protein fold.

**DNA-binding studies of Ec-MtlR**

The mannitol operon has been best described in *E. coli* and the effect of Ec-MtlR on the transcription of the operon has been studied (Fig. 5B) (9). However, the *in vitro* binding of MtlR to the operator/promoter region has not been established so far. Similarly to the results described by Figge (9), we were able to show an inhibitory effect of MtlR on the uptake of mannitol into *E. coli* by plating on MacConkey agar plates. *In vitro* studies using several different techniques were undertaken to identify the DNA binding site of MtlR within the mannitol operon. In experiments with an optical biosensor (IAsys)(34,35), we tried to prove the hypothesis that MtlR may bind to two palindromic sequence stretches identified on the *mtlR* promoter (Fig. 5B).

Two different DNA sequences from the mannitol operator site were investigated in electrophoretic mobility shift assay studies with Ec-MtlR. They were named as MtlR1, MtlR2, for operator1 and operator2 respectively (Fig. 5B). MtlR1 consists of sequence 5’-CTCGGGCTTCCAGCTGCG and MtlR2 consists of sequence 5’-ACATAAGAAGGGTGTATTTATGT, operator1 and 2 were created by PCR with *mtlR_pro1f* (5’-TATGACGAAGGCATAACATGC) as forward primer and the reverse oligonucleotides of the MtlR1-box (*mtlR1_rev*) and the MtlR2-box (*mtlR2_rev*) as reverse primer. No interaction could be detected between the MtlR protein and the two palindromic sequence stretches on the promoter (MtlR1 and MtlR2) (Figs. 5C and D). In order to exclude the possibility of another binding site on the operator-promoter region the whole region from the first CRP binding site to the two palindromic sequence sites MtlR1 and MtlR2 was investigated in gel shift experiments. Again no binding of MtlR to the DNA could be detected. Different buffer conditions, in the presence and absence of mannitol and various protein concentrations were tested without success. The binding of the global regulatory protein, fructose repressor (FruR) to the *mtlR*-operon was used as a positive control in these experiments (36).

We have also investigated the DNA-binding properties of MtlR using SELEX based approach. This allows the identification of the DNA sequence, which binds to the protein of interest. The proteins tested were the well characterized sequence specific Fructose Repressor (FruR), the structure specific HMG-box protein Cmb1, MtlR and a BSA control. Binding of the random-sequence oligonucleotide R76 as well as to specific binding sequences could be detected for...
DISCUSSION

The MtlR has been experimentally studied and proposed to be a transcriptional regulator (9). Its gene \textit{mtlR} is one of the conserved components of the mannitol operon across many bacterial species. The presence of two putative palindromes within the operator-promoter region implied possible MltR binding sites in the early studies of the \textit{E. coli} mannitol operon(9). Many bacterial genomes code for the MtlR sequence homolog named YggD. However, the function of YggD has never been experimentally determined and the \textit{yggD} gene location in chromosomes provides little clues about its function. In this study, we have established that Ec-MtlR does not bind regulatory regions of mannitol operon and we have determined the crystal structures of both MtlR and YggD. It is apparent that MtlR and YggD resemble each other in both their monomer and dimer structures, implying potentially similar functions.

Although assigned to the same protein sequence family (Pfam05068), MtlR and YggD are largely different in their molecular sizes. Generally, MtlRs are longer having about 195 residues while YggDs are shorter with about 170 amino acids. The primary difference between MtlR and YggD seems the presence of additional 20 plus N-terminal residues in MtlR. For example, MtlRs of \textit{Shigella} and phylogenetically indistinguishable \textit{E. coli} are 26 residues longer than their YggDs (Fig. 3). However, this rule is not observed by all MtlRs. In most \textit{Vibrio} species MtlRs have 176 residues with exceptions like \textit{V. cholerae} 623-39, which has an MtlR of 193 residues (17). Therefore, these extra N-terminal residues in many MtlRs are unlikely determinants of the mannitol operon repressor’s function. Rather, it could be a species-specific factor. Structurally, the extra N-terminal sequence could lead to an elongated \(\alpha\) helix (Fig. 2B) and a loop linked to a possible short helix at the very N-terminus, based on secondary structure prediction (http://cubic.bioc.columbia.edu) (37). The prediction seems to be consistent with the results of limited proteolysis of Ec-MtlR, which showed the 14 N-terminal residues removed when Ec-MtlR was treated with trypsin or chymotrypsin (see Experimental Procedures).

Another line of evidence suggesting that MtlR and YggD may function similarly is that MtlR and YggD do not necessarily coexist in one species while we believe that the mannitol-specific PTS is a common and vital molecular system in all bacterial species. For example, only the \textit{yggD} gene was detected in \textit{S. dysenteriae} Sd197 meanwhile only the \textit{mtlR} gene was reported in \textit{S. sonnei} Ss046 (38). In \textit{S. dysenteriae} Sd197, the \textit{yggD} gene is clustered with \textit{cmtA} and \textit{cmtB}, the two genes that are the frequent neighbors to \textit{yggD}. They are generally annotated as cryptic mannitol-specific PTS enzyme:IIB/IIC components and IIA component, respectively, which sequentially corresponds to the components of MtlA. A recent solution structure study of \textit{E. coli} K12 CmtB revealed that the overall structure of CmtB is highly similar to that of the IIA domain of MtlA(16). Though the exact phosphoryl transfer mechanism of CmtB could be different due to the position change of the catalytic arginine residue, the function of CmtB may be similar to that of IIA domain of MtlA. This information further suggests that YggD could play similar role to MtlR in the regulation of mannitol-specific PTS.

With the structures of Vp-MtlR and Sf-YggD determined we analyzed the presence of potential DNA-binding motifs in these proteins. It is especially intriguing that they both form homodimers, which is expected for prokaryotic transcription factors binding to palindromic operator sequences within promoter region (9). However, no known DNA-binding motif is found in both structures, that could correspond to the DNA double helix spacing in major or minor grooves (39,40). Although MtlR/YggD are all \(\alpha\)-
helical there is no helix-turn-helix (HTH) motif or other related structural motif typically found in transcriptional regulators (40). The surface of MtlR/YggD dimer is rather flat without the protruding structural element that could specifically recognize bases of DNA in the major or minor groove. The lack of an apparent DNA-binding motif also seems to be consistent with the results from the DALI structural homolog search mentioned previously. Even low score hits did not include DNA-binding proteins.

Moreover, both Vp-MtlR and Sf-YggD are very acidic proteins with calculated pls of 4.46 and 4.33, respectively. They are both highly negatively charged with the same net charges of -28e per dimer. The net charges per amino acid for two molecules are -0.083 and -0.080, respectively. The surface electrostatic potential shows the overwhelming negative potentials of the surfaces of both MtlR and YggD dimers; there are no major positive charge patches present on the surface (Fig. 6). The electronic dipole moments of Vp-MtlR and Sf-YggD dimers are calculated to be 90 e*A and 71.56 e*A, respectively. The calculation was performed by using the “Web Server to Calculate Dipole Moments of Proteins”, (http://bioportal.weizmann.ac.il/dipol/dipint.html) (41,42). The electrostatic properties including net charge and dipole moment are believed to be important to a protein’s ability to bind DNA (43,44). It was found that these bulk electrostatic properties of DNA-binding proteins are significantly different from those of non-binding proteins (44). Based on these calculations, both Vp-MtlR and Sf-YggD, are not favored as DNA-binding proteins due to the net negative charges and small dipole moments.

It has been reported that the mannitol operon of E. coli is subject to activation by the cyclic AMP receptor protein (CRP) (45). There are 5 unusual CRP binding sequences in the promoter region and all of them have been proved to bind CRP in vitro in DNA band mobility shift experiments (Fig. 5B) (45). Additionally, the fructose repressor, FruR also binds to the regulatory region of the mannitol operon as one common regulatory element of almost every major carbon metabolism pathway (36). The FruR recognition element was indicated to be between the –35 and –10 promoter region of the mannitol operon (Fig. 5B). The mannitol repressor MtlR negatively regulates the expression of the mannitol operon and it was proposed to block expression through direct interaction with DNA the two palindromic sequences located within mtlR promoter (9). However, thus far there is no corroborating experimental evidence that MtlR binds directly to the operator regulatory region, (9). Our in vitro DNA-binding experiments clearly showed there was no binding of MtlR to the promoter region, even at high protein concentrations. A search for a different DNA regulatory sequence has also been unsuccessful. Considering the multiple binding sites for transcriptional factors in the mannitol operon, five for CRP, one for FurR, two for RNA-polymerase and potentially one or two for MtlR (or other regulatory protein), the operon control was regarded as one of the most complex found in bacteria (45). However, with the structures of Vp-MtlR and Sf-YggD determined and their properties tested, it is doubtful whether MtlR and/or YggD can bind DNA directly and regulate transcription in chorus or in competition with other DNA-binding regulators.

It is noticed that the mannitol uptake PTS of many Gram-positive bacteria such as Bacillus (3,46), Streptococcus (47), and Clostridium(48), have quite different mannitol operon repressors. They are similar to antiterminators and assigned to a different protein family (Pfam00874). For example, B. stearothermophilus MtlR is composed of 697 amino acids and contains a HTH DNA-binding motif and two antiterminator-like PTS regulatory domains (49). This MtlR can bind to the promoter region and likely control the expression of the mannitol operon by blocking RNA-polymerase binding to the promoter (3). Moreover, the activity of MtlR itself can be
regulated through phosphorylation by PTS components (3,50).

It is possible that there is yet another transcriptional factor (TFX) that recognizes mtlR operator1 and 2 sequences and links expression from mtlR operon to other cellular function or external signal. We propose that the mannitol repressor MtlR and possibly YggD are a part of a transcriptional complex that controls the mannitol operon expression of *E. coli, Shigella, Vibrio* and other Gram-negative bacteria. Instead of directly binding to DNA, they may interact directly or indirectly with other DNA-binding protein(s), resulting in the repression of mannitol operon transcription. Therefore, the true function of MtlR may be a competition with RNA polymerase in the initiation complex involving several proteins including CPR, FurR and TFX. This could represent a transcription initiation complex comparable to simple eukaryotic initiation complexes but functioning in bacteria.

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Figure Legends

Figure 1. Crystal packing of Vp-MtlR monomers and Vp-MtlR dimer. (A) The four Vp-MtlR monomers in one asymmetric unit. The monomers A and B form a dimer, and monomers C and D form a dimer. Both dimers are of 2-fold pseudo-symmetry. For clarity, only the front helices of the AB-dimer are labeled. The AB-dimer and CD dimer form extensive interactions across their interface and potentially form a tetramer as discussed in the text. (B) A ribbon drawing of a Vp-MtlR dimer (AB-dimer). The orientation of the dimer is related to its position in Fig.1A by a rotation about 90° around the horizontal axis. All helices of the novel layered α-helical fold are labeled. The α8 helix is missing in Vp-MtlR in comparison to the structure of Sf-YggD shown in Fig.2. This figure and other ribbon diagram figures in this paper were prepared using the program Pymol (http://www.pymol.org).

Figure 2. Crystal packing of Sf-YggD monomers and Sf-YggD dimer. (A) The four Sf-YggD monomers in one asymmetric unit. The monomers B and C form a pseudo 2-fold dimer while the monomers A and D form dimers with their 2-fold symmetry-related monomers, respectively. For clarity only front helices of the BC-dimer are labeled. The interaction between monomers A and B is different from the interaction between monomers C and D, by skipping one turn of the helix. They represent two types of primary dimer-dimer contacts found in the structure. (B) A ribbon drawing of a Sf-YggD dimer (BC-dimer). The orientation of the dimer is related to its position in Fig.2A by a rotation about 90° around the horizontal axis. The α8 helix is unique to the YggD structure. (C) A structural alignment of Vp-MtlR (dark green) and Sf-YggD (cyan) monomers.

Figure 3. Multiple Sequences Alignment of MtlR and YggD. The identical residues are highlighted in red and the similar residues in cyan. The residues forming the hydrophobic core of MtlR and YggD are marked with *. The residues contributing to their dimerization are marked with ^. The helices of Vp-MtlR and Sf-YggD structures are underlined with blue coils. The first 14 N-terminal amino acids of Ec-MtlR were removed with limited proteolysis. The proteins used in the alignment include E. coli MtlR (Ec-MtlR), Vp-MtlR I (from chromosome I), Sf-YggD, E. coli YggD (Ec-YggD) and Vp-MtlR II (from chromosome II). The S. flexneri MtlR (Sf-MtlR) and S. sonnei MtlR I (Ss-MtlR) are identical to Ec-MtlR in primary sequences. The S. dysenteriae MtlR (Sd-YggD) is identical to Sf-YggD.

Figure 4. The dimer interface and potential tetramer interface of Vp-MtlR. (A) Stereo ribbon drawing of the dimeric interface of the AB-dimer as shown in Fig.1. The predominant hydrophobic residues involved in the dimer formation are drawn in sticks and colored differently according to monomers. Since the dimer is of a pseudo 2-fold symmetry, only those residues from one monomer are labeled. (B) Stereo ribbon drawing of the potential tetrameric interface, which corresponds to the interface between the monomers B and C and represents the interaction between the AB-dimer and the CD-dimer as shown in Fig.1A. The residues from two dimers involved in the dimer-dimer interactions are drawn in sticks and colored in different codes. The two flexible loops between α7 and α9 helices form a pair of anti-parallel strands with five mainchain-mainchain hydrogen bonds.

Figure 5. Size exclusion chromatography and DNA-binding studies. (A) Size exclusion chromatographies of Vp-MtlR and Sf-YggD. The elution peak of Sf-YggD is sharp and its apparent molecular size was estimated to be 47.8 kDa, slightly larger than a Sf-YggD dimer. The Vp-MtlR peak is broad and the eluted Vp-MtlR was estimated to be 90.4 kDa, roughly a tetramer size. A possible mixture of oligomers is proposed as discussed in the text. (B) A schematic drawing of the mannitol
operator/promoter region and the four constructs used in EMSA experiments (operator1, operator2, MtlR1 and MtlR2). (C) Electrophoretic mobility shift assay with DNA fragments corresponding to the two palindromic boxes MtlR1-box and MtlR2-box and (D) the whole operator region.

Figure 6. The electrostatic potential surface representations of Vp-MtlR and Sf-YggD. (A) The Vp-YggD dimer on the top is in an orientation similar to the orientation of the AB dimer on the left panel of Figure 1A. The dimer on the bottom is rotated about 90 degrees along the horizontal axis. (B) The Sf-YggD dimer on the top is in an orientation similar to the orientation of the BC dimer on the top panel of Figure 1B. The dimer on the bottom is rotated about 90 degrees along the horizontal axis. The approximate distances between two ridges from both Vp-MtlR and Sf-YggD dimers are about 50Å as marked. The figures were prepared with the program GRASP(51).
### Table 1: Crystallographic statistics

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<th>Vp-MtlR</th>
<th>Sf-YggD</th>
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<td>Space group</td>
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<tr>
<td>(Å$^2$), Mainchain/Sidechain</td>
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$^1$ Including Bijvoet pairs; $^2$(Last resolution bin); $^3$ Peak and Inflection
Figure 1.
Figure 3.

Ec-Mt1R  gi|1790029  MVDQAGDTRPNNLSDMQATMEQTOFQERVNLERNAGKTVRSLFILTAVEILTEAVNLVLOV  64
Vp-Mt1R I gi|637397307  MADQINESIIERNSAPSVGFFVATVDFESIDGLIQR  42

Sf-YggD  gi|30042499  MATLTEDDVLEQLELDAQDNLFSQMKTAHSSLQGRIQFLPSL  41

Ec-YggD  gi|1789297  MATLTEDDVLEQLELDAQDNLFSQMKTAHSSLQGRIQFLPSL  41
Vp-Mt1R II gi|637400663  MSREDLEKIEDECNATAPIQSNKILK1KALLPS  39

FRK-DDYAVYAVFVPLLGLGVLRLKLIGGLVINKQYEDAEELLMALREELNHGNEYATTDDEILGPPFELHCVAAL---PP  149
FRK-DDNFAVQSVGGLDGVVRLKLLEGGLVLDDYHIDEDLKLKMNLSAASDYETFDDNILEPIKKHLVKKMVQ  129

FVNDNEEIVYEAVKPLAQQGPLLDDIVALRLLYALGKMDKWLYADITHFSQYWHY1NQDETPFGADDITWDFSVNVSITRNN---A  126

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PPQFEP---ADSSLYAMQIQRYQAQVSTYVLSLTELISIKSLKKAFOK  195
LEVNBDDIDILDBFQGDQGQITISQGILSAIVEICNELGDKSF  176

TLYDALKAKFADFAWAREFSGM/VFTAIITAVTTTLKLT  168
MLYDALKAKFADFAWAREFSGM/VFTAIITAVTTTLKLT  169
PYLESINQLRFSSFSEXQPMRVEHSLTIVKLSCEMLIEREBEINOQ  172
Figure 5.
Figure 6.
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The Mannitol Operon Repressor MtlR Belongs to a New Class of Transcription Regulators in Bacteria
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