Protein Signatures That Promote Operator Selectivity among Paralog MerR Monovalent Metal Ion Regulators*

Received for publication, January 14, 2013, and in revised form, May 31, 2013 Published, JBC Papers in Press, June 3, 2013, DOI 10.1074/jbc.M113.452797

María V. Humbert‡§1, Rodolfo M. Rasia‡¶2, Susana K. Checa‡§2, and Fernando C. Soncini‡§2,3

From the ‡Instituto de Biología Molecular y Celular de Rosario (IBR, CONICET-UNR), ¶Departamento de Microbiología, and †Departamento de Química Biológica, Facultad de Ciencias Bioquímicas y Farmacéuticas, Universidad Nacional de Rosario, Consejo Nacional de Investigaciones Científicas y Técnicas, Ocampo y Esmeralda, 2000-Rosario, Argentina

Background: Two nucleotide bases distinguish promoters controlled by paralog MerR monovalent metalloregulators, avoiding cross-activation.

Results: Specific residues within the DNA-binding region of the regulators were identified as responsible for the selectivity in the operator recognition.

Conclusion: Co-evolution of both the regulator and its target operator sequences prevents cross-activation of paralog regulatory circuits.

Significance: The basis for regulator/operator specificity among MerR monovalent metalloregulators is described.

Two paralog transcriptional regulators of the MerR family, CueR and GolS, are responsible for monovalent metal ion sensing and resistance in Salmonella enterica. Although similar in sequence and also in their target binding sites, these proteins differ in signal detection and in the set of target genes they control. Recently, we demonstrated that selective promoter recognition depends on the presence of specific bases located at positions 3′ and 3 within the operators they interact with. Here, we identify the amino acid residues within the N-terminal DNA-binding domain of these sensor proteins that are directly involved in operator discrimination. We demonstrate that a methionine residue at position 16 of GolS, absolutely conserved among GolS-like proteins but absent in all CueR-like xenologs, is the key to selectively recognize operators that harbor the distinctive GolS-operator signature, whereas the residue at position 19 finely tunes the regulator/operator interaction. Furthermore, swapping these residues switches the set of genes recognized by these transcription factors. These results indicate that co-evolution of a regulator and its cognate operators within the bacterial cell provides the conditions to avoid cross-recognition and guarantees the proper response to metal injury.

Transcriptional regulators of the MerR family modulate transcription in response to different environmental signals, including heavy metal ions, organic compounds, or oxidative stress (1). These proteins have a structurally conserved N-terminal DNA-binding domain with two helix-turn-helix (HTH)4 motifs separated by a two-stranded antiparallel β-sheet, in a α1-α2-β1-β2-α3-α4 topology. This region is connected to a variable C-terminal effector-binding domain by an extended α-helix, forming a coiled-coil dimerization region. Usually, these regulators recognize pseudopalindromic operator sequences in ρ8-targeted promoters with longer (19- or 20-bp) spacers between the −35 and −10 elements that prevent open complex formation by RNAP without an activator (2–4). According to the current model of MerR-mediated induction, after the signal is detected by the regulator, the information is transduced to the DNA-binding domain, triggering localized base pair breaking and base sliding in the operator. This results in a realignment of the promoter elements that now allows proper RNAP-DNA interaction and transcription initiation. The solved structure of three MerR homologues bound to their target operators, the drug-binding BmrR and Mta proteins and the oxidative stress sensor SoxR, envisages a conserved mechanism for DNA recognition (3–5). Protein-DNA interactions involve hydrogen bonds and van der Waals contacts between amino acid residues mostly belonging to the α2-helix and the β2 strand of the N-terminal DNA-binding domain and the DNA backbone of the target operator sequence (3–5). It has been proposed that these contacts serve both to stabilize the distorted DNA conformation and to provide regulator/operator selectivity.

In Salmonella enterica serovar Typhimurium, two metal ion sensors of the MerR family, CueR and GolS, control the transcription of genes coding for factors responsible for monovalent metal ion resistance (6–8). These paralog transcription factors probably evolved from a common ancestor by gene duplication followed by divergence that rewired both signal recognition and the set of controlled genes (9, 10). The copper sensor CueR, present in most Gram-negative bacteria, induces the expression of its target genes in response to either Cu(I), Ag(I), or Au(I) (11, 12), whereas the horizontally acquired GolS regulator evolved to preferentially sense Au(I) ions (6, 7). As we
mentioned before, the set of genes controlled by each transcription factor also differs. CueR activates the expression of the P-type ATPase CopA, the multicopper oxidase CueO/CuiD, and the periplasmic copper-binding protein CueP (7, 8, 13), whereas GolS induces transcription of the Salmonella-specific genes encoding for another P-type ATPase, named GolT, the metal-binding protein GolB and the CBA-type GesABC efflux system (6, 14). We recently demonstrated that although these regulators recognize similar operator sequences at their target promoters, particularly at the −35 promoter region, the presence of only two distinctive nucleotide bases at the 3′- and 3′-positions relative to the center of the operator determines these positions (Fig. 1). Portions of the DNA-binding domain of each regulator by the series of GolS and CueR hybrid proteins replacing different selective recognition. To identify the amino acid residues DNA-binding domain of the transcription factors that direct must correlate with specific amino acid residues within the sequence are conserved in promoters that are proposed to be operators predicted to be controlled by the CueR group harbor C or G at these positions (15).

GolS-controlled promoters have an A at the 3′-position and a T at the 3′-position relative to the center of the operator, whereas operators recognized by CueR have either a C or a G at these positions (Fig. 1A). Switching these nucleotide bases in gol-like or cue-like operators is sufficient to swap the regulator dependence (15). In other words, the mutant golBcc promoter, which harbors a C at the operator positions 3′ and 3′ as PcopA, decreases the affinity for its native regulator GolS but has an increased binding to CueR compared with the wild-type golB promoter. A similar switch in regulator dependence is observed when the mutant copAAT and the wild-type copA promoters were compared. In fact, the A and T signature nucleotide bases at 3′- and 3′-positions from the center of the dyad operator sequence are conserved in promoters that are proposed to be transcriptionally controlled by GolS-like regulators, whereas operators predicted to be controlled by the CueR group harbor C or G at these positions (15).

The presence of selective nucleotide bases at the operators must correlate with specific amino acid residues within the DNA-binding domain of the transcription factors that direct selective recognition. To identify the amino acid residues involved in operator discrimination, here we constructed a series of GolS and CueR hybrid proteins replacing different portions of the DNA-binding domain of each regulator by the same region of the paralogous protein and tested their ability to direct expression from the golB or copA promoter. Together with site-directed mutagenesis and in silico modeling, these studies demonstrate that the residue at position 16 from the α2-helix of both CueR and GolS is essential for selective recognition. Our results provide additional evidence of the co-evolution of both the MerR regulators and the regulated genes to avoid cross-recognition and guarantee a proper response to metal injury.

**EXPERIMENTAL PROCEDURES**

**Bacterial Strains and Growth Conditions**—Bacterial strains (all derivatives of Escherichia coli W3110 strain or S. enterica serovar Typhimurium 104028, except when indicated) and plasmids used in this study are listed in Table 1. Oligonucleotides are listed in Table 2. Cells were grown at 37 °C in Luria-Bertani broth (LB) or on LB agar plates. Ampicillin, tetracycline, kanamycin, and chloramphenicol were used when necessary at 100, 15, 50, and 20 μg ml−1, respectively. Reagents, chemicals, and oligonucleotides were from Sigma, except for the Luria-Bertani culture medium, which was from Difco.

**Bacterial Genetic and Molecular Biology Techniques**—E. coli strains carrying deletions on the lacZ, cueR, or copA genes were generated in the W3110 strain by Lambda Red-mediated recombination (16) using the appropriate primer pairs listed in Table 2. The deletions were individually transferred to different W3110 derivatives by P1-mediated transduction (17) to generate the PB10305 strain. When necessary, the antibiotic resistance cassette inserted at the deletion point was removed using the temperature-sensitive plasmid pCP20 carrying the FLP recombinase (18).

Construction of hybrid cueR-Ns, cueR-HTH1S, cueR-(HTH1+L)S, and golS-Ns alleles was carried out by using mutagenesis by PCR overlap extension or SOE-PCR (19) (for the chimeric protein nomenclature used, see Figs. 1B and 2A). Briefly, we performed two independent PCRs using the complementary primers carrying the desired hybrid junction and oligonucleotides homologous to the 5′ and 3′ wild-type gene of interest harboring BamHI or HindIII restriction sites (Table 2). Then the products of both PCRs were purified and combined in a third PCR with the appropriate forward or reverse primers to generate the final product. To construct the cueR-(L+HTH2)S and cueR-HTH2S alleles, we amplified a first PCR product using GolS-Loop-Fw/GolS-HTH2-Rv or GolS-HTH2-Fw/ GolS-HTH2-Rv primer pairs (Table 2). Each product was employed as a primer in two independent PCRs along with CueR-ORF-Fw or CueR-ORF-Rv to generate the overlapping fragments used in the final PCR, as described above. The cueR-α2S, golS-α2R, or the alleles carrying point mutations were amplified in a single PCR using the forward oligonucleotides harboring the modification and the BamHI restriction site and the appropriate reverse primer. The final PCR fragments were purified and individually cloned into BamHI/HindIII-digested pUH21–2laqF or the pSU36 vector to generate the expression plasmids listed in Table 1.

The plasmids and the linear DNA fragments were introduced to E. coli strains by electroporation using a Bio-Rad device following the manufacturer’s recommendations. All constructs were verified by DNA sequencing.

**Metal Induction Assays**—The levels of expression of the lacZ reporter gene under the control of the native (PcopA or PgolB) or mutated (PcopAAT or PgolBcc) promoters in the presence of 10 μM AuHCl₃ or 100 μM CuSO₄ or without metal added were assessed essentially as previously described (15) using overnight cultures on LB. β-Galactosidase assays were carried out essentially as described (17).

**Protein Purification**—GolS, CueR, or the hybrid proteins GolS-α2S and CueR-α2S were overexpressed and purified from E. coli XL1-Blue strain essentially as described previously (14). 0.1 or 0.5 mM IPTG (for CueR or GolS variants, respectively) was added to the cultures at an A₆₀₀ of 0.6 to promote protein expression. All procedures were carried out at 4 °C. The protein profile of purified samples was determined by SDS-PAGE, and the concentration was calculated, recording absorbance at 280 nm with an ε of 4320 m⁻¹ cm⁻¹ (CueR), 6585 m⁻¹ cm⁻¹ (CueR-α2), 11,835 m⁻¹ cm⁻¹ (GolS), or 10,345 m⁻¹ cm⁻¹ (GolS-α2S), or by Bradford assay, using bovine serum albumin as a standard.

**Protein-DNA Interaction Analysis**—Electrophoretic gel mobility shift assays were performed using purified wild-type or
TABLE 1

Bacterial strains and plasmids used in this study

| Strain           | Relevant genotype | Reference or source |
|------------------|-------------------|---------------------|
| XL1-Blue         | recA1 endA1 gyrA96 thi-1 | Stratagene          |
|                  | hsdR17 supE44 relA1 lac    |                     |
|                  | [F proAR lacI52 Δ(M15)  |                     |
|                  | Ta105 (T-)]             |                     |
| W3110            | F' ΔIN[rpsL-5,tem-5] rph-1 | Ref. 27            |
| PB8885           | W3110 lacZ:cat | This study          |
| PB6731           | W3110 cueR::cat | This study          |
| PB10285          | W3110 lacZ:copA::kan | This study          |
| PB10285          | W3110 lacZ:copA | Ref. 13             |
| PB10305          | W3110 lacZ::copA cueR:cat | Ref. 13 |
| PB10683          | 14028ΔgolAgesABC ΔcueP | Laboratory         |
|                  | ΔcopA cueR::cat | stock               |

| Plasmid          | Relevant genotype | Reference or source |
|------------------|-------------------|---------------------|
| pKD3             | oriR 60, P4 bet’-gam’ Amp’ | Ref. 16 |
| pKD4             | ori 8 FRT Amp’ FRT | Ref. 16             |
| pCP20            | λ cl857 tsPr rpsL R 48 Amp’ Cm’ | Ref. 16 |
| pUH21–2 lacP      | orlβ34  Amp’ lacP | Ref. 28             |
| pb1205           | pUH::golS | Ref. 6              |
| pb1204           | pUH::golSER22 | This study          |
| pb1209           | pUH::golS-R22 | This study          |
| pSU36            | ori535’ Kmr’ | Addgene             |
| pb1239           | pSU36::golR | This study          |
| pb1395           | pSU36::golN53 | This study          |
| pb1391           | pSU36::golR-HTH1 + L5 | This study          |
| pb1392           | pSU36::golR-HTH2L5 | This study          |
| pb1393           | pSU36::golR-(L + HTH2) L5 | This study          |
| pb1394           | pSU36::golRHTH2 | This study          |
| pb1396           | pSU36::golR2222 | This study          |
| pb1423           | pSU36::golS2222 | This study          |
| pb1399           | pSU36::golR4121 | This study          |
| pb1424           | pSU36::golRHT121 | This study          |
| pb1400           | pSU36::golR2221 | This study          |
| pb1425           | pSU36::golR2222 | This study          |
| pb1401           | pSU36::golRASM2222 | This study          |
| pb1390           | pSU36::golS | Ref. 6              |
| pb1397           | pSU36::golS-N53 | This study          |
| pb1414           | pSU36::golS-N14 | This study          |
| pb1402           | pSU36::golS-M14 | This study          |
| pb1403           | pSU36::golS2222 | This study          |
| pb1416           | pSU36::golS2222 | This study          |
| pb1404           | pSU36::golSASM2222 | This study          |
| phMC1871         | pBR322, Tet’, lacP | Amersham, Biosciences |
| ph1225           | pPMC1871::PcopA | Ref. 15             |
| ph1222           | pPMC1871::PgolB | Ref. 15             |
| ph1223           | pPMC1871::PcopA2 | Ref. 15             |
| ph1220           | pPMC1871::PgolB2 | Ref. 15             |

mutant regulators as described previously (15). The primers used to amplify the PcopA or PgolB promoter region are listed in Table 2.

Fluorescence anisotropy assays were carried out essentially as described (20, 21). Fluorescin-labeled double-stranded DNA fragments harboring the copA or golB operator sequences were generated by incubating pairs of single-stranded oligonucleotides (Table 2) at 85 °C for 10 min and then allowing the mix to cool at room temperature. The binding of native and mutant transcriptional regulators to PcopA andPgolB promoters was measured using a Varian Cary Eclipse fluorescence spectrophotometer (Agilent Technologies). All fluorescence anisotropy titrations were done in 10 mM Tris buffer with 10 mM NaCl, 2 mM MgCl2, and 5 mM DTT, pH 7.3. The fluorescence was excited at 495 nm. Anisotropy (r) was calculated as r = (I - G × I1)/(I1 + 2 × G × I1), where I1 and I are the fluorescence intensity parallel and perpendicular to the excitation polarization, respectively, and G is the correction factor for the instrument’s different responses to light of parallel and vertical polarizations. Binding isotherms were fitted to the equation, r = rP + (rP − rN) × (kA × x)/(1 + kA × x), where rP and rN are the anisotropy values for free and protein-bound DNA, respectively, x is the total protein concentration, and kA is the association constant of the protein–DNA complex, by nonlinear regression using the Sigma Plot software. Dissociation constants (kD) were estimated from the equation, r = rP + a × x/(b + x), where a = (rP − rN) × (kA × x)/(1 + kA × x) and b = kD.

In Silico Modeling—A structural model for CueR, CueR-α2g, and GolS was generated by homology modeling using Rosetta version 3.1. The structure of BmrR transcription factor bound to its promoter (Protein Data Bank code 3Q5R) was used as a template. 100 initial structures were generated using a cyclic coordinate descent algorithm combined with fragment assembly for modeling the CueR sequence with gaps. The 10 lowest energy structures were subsequently refined, and the lowest score structure was used as a model. The model structures were then superimposed on the structure of the complex used as a template in order to locate the position of the residues of interest (Ala-16 and Phe-19) with respect to the promoter DNA and compare their relative conformation in the different proteins.

RESULTS

Selective Regulator/Operator Recognition Is Directed by Amino Acids within the α2-Helix of the Sensor Protein—Salmonella CueR and GolS can distinguish their target operators from those of the paralog regulator by selectively recognizing two nucleotide bases located at the 3'- and 3-positions from the center of the operator (Fig. 1A) (15). To identify the amino acid residues within the DNA-binding domain of the transcription factor that direct operator discrimination, we designed chimeric proteins, in which the N-terminal DNA-binding domain of CueR (from Met-1 to Asn-68) was replaced by the equivalent region of GolS and vice versa (Fig. 1B). The encoding mutant alleles were cloned in plasmids and introduced either into an S. enterica serovar Typhimurium Δgol ΔcueP ΔcueR-copA or into an E. coli W3110 ΔcueR-copA ΔlacZ strain harboring lacZ reporter fusions to copA or golB promoters. (The use of strains deleted in copA helped to minimize differences in intracellular copper levels derived from partial activation of the copper transporter by the regulator variants.) The cells were grown in LB in the absence or presence of either 10 μM AuHCl or 100 μM CuSO4, concentrations required to attain the maximal induction of the reporter genes (7, 14). The chimeric CueR-N5 and GolS-N5 regulators were functional to activate transcription of the reporter genes in response to the metals, but their induction pattern resembled that attained by the paralog regulator (Fig. 1C) (data not shown). In other words, CueR-N5 activated the expression of the reporter gene more efficiently under the PgolB promoter than under the native PcopA promoter, resembling wild-type GolS, and GolSN5 switched its operator recognition preference, acting as a better inducer of CueR-regulated promoters. Because essentially identical results were obtained using either Salmonella or E. coli, we choose to continue the analysis in the latter species.

20512 JOURNAL OF BIOLOGICAL CHEMISTRY VOLUME 288 • NUMBER 28 • JULY 12, 2013
### Table 2

Oligonucleotides used in this study

| Primer name | Sequence (5'-3') | S'-Restriction site | Purpose |
|-------------|-----------------|---------------------|---------|
| lacZ-P1 | TTTGAGACCGCGAACTCCATTTCTCGAAGACTGTTTGCTTGTGA | | Deletion of the lacZ gene |
| lacZ-P2 | GCGAAATACGGGCGAACTCCATTTCTCGAAGACTGTTTGCTTGTGA | | Deletion of the lacZ gene |
| copA-P1 | AAAAGGTACATGCCAACTCCATTTCTCGAAGACTGTTTGCTTGTGA | | Deletion of the copA gene |
| copA-P2 | CCCATTTCTCGAAGACTGTTTGCTTGTGA | | Deletion of the copA gene |
| cueR-P1 | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAATCT | | Amplification of wild-type or hybrid cueR |
| cueR-P2 | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-NS | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-HTH1S | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-HTH2S | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-HTH3S | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-HTH4S | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-HTH5S | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-HTH6S | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-HTH7S | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |
| cueR-HTH8S | GAGGATCCATATGAATATTAGCGATGTGGCGAAAAAAACCGGTTTATCCCGTGTGGCGAAAAAACC | | Amplification of wild-type or hybrid cueR |

Protein Signatures for DNA Selectivity among MerR Regulators

**Generation of**

| Restriction site | Purpose |
|------------------|---------|
| BamHI | Amplification of wild-type or hybrid cueR |
| HindIII | Amplification of wild-type or hybrid cueR |
| XmaI EMSA | Fluorescence anisotropy |

**Fluorescence anisotropy**

- Fluorescence anisotropy
- Fluorescence anisotropy
- Fluorescence anisotropy
- Fluorescence anisotropy
- Fluorescence anisotropy
- Fluorescence anisotropy

**XmaI EMSA**

- Fluorescence anisotropy
- Fluorescence anisotropy
- Fluorescence anisotropy
- Fluorescence anisotropy
- Fluorescence anisotropy
- Fluorescence anisotropy

**Xm1**

- EMSA
- EMSA
- EMSA
- EMSA
- EMSA
- EMSA
To delimit the region of CueR responsible for operator discrimination, we replaced different portions from the N-terminal DNA-binding domain of CueR by the equivalent regions of GolS and analyzed the ability of each chimeric regulator to activate the expression of the reporter genes under either a GolS- or a CueR-controlled promoter (Fig. 2). Like CueR-NS, CueR-HTH1S and CueR-α2S, carrying the first HTH motif (from Met-1 to Lys-23) or the α2-helix (from Thr-13 to Lys-23) of GolS, respectively, drove a reporter gene expression like GolS (Fig. 2B). These hybrid proteins showed an improved activation of P<sub>golB</sub>-driven expression and a decreased induction of transcription from the native P<sub>copA</sub> promoter compared with wild-type CueR, suggesting that the operator selectivity resides within the α2-region. Furthermore, the equivalent α2-helix replacement in GolS resulted in a transcriptional regulator that exhibited a CueR-like expression pattern with an increased metal-activated expression of the reporter gene from P<sub>copA</sub> and almost no induction of transcription from the P<sub>golB</sub> promoter (Fig. 2B). The other CueR chimeric constructions with the replacement of either the α1-α2 loop or of the HTH2 domain from GolS (i.e. CueR-HTH2<sub>α</sub>, CueR-(HTH1+L)<sub>α</sub>, or CueR-(L+HTH2)<sub>α</sub>) resulted in less active or inactive regulators.
the mobility of the PcopA promoter compared with CueR. On the other hand, CueR-α2a substantially gained an apparent affinity for the PgoIB promoter. GolS-α2α shows much lower affinity for the PgoIB promoter compared with the parental GolS regulator, whereas it gained affinity for the PcopA region, resembling wild-type CueR (Fig. 3A).

The binding affinity for each regulator/operator pair was estimated by fluorescence anisotropy titration. In these assays, we used fluorescein-labeled 41-base pair double-stranded DNA containing the operator sequences from copA or golB. Each labeled dsDNA probe was titrated with increasing amounts of CueR, GolS, or the mutant variants CueR-α2a or GolS-α2a. Representative titration curves for each pair are shown in Fig. 3B. The curves for GolS/PgoIB and GolS-α2a/PcopA did not reach saturation because these proteins aggregate at high concentrations. This behavior is evidenced by a sharp increase in the anisotropy values induced by the contribution of light scattering from higher molecular weight particles to the measured anisotropy at protein concentrations higher than those reported. The saturation levels of anisotropy observed in both CueR variants (that reached saturation) are 0.102 and 0.133. The values obtained from fitting the available data on the complexes with GolS variants are 0.133 and 0.147, which are close to the values obtained for CueR. Assuming that similar kinds of complexes are formed by both proteins, we can conclude that the extrapolated saturation values are in good agreement with the physical system. With all of this information, the dissociation equilibrium constants (kD) were calculated for the interaction of the CueR/PcopA and GolS-α2α/PcopA as 141 ± 15 and 198 ± 49 nM, respectively. Similarly, the estimated kD values for GolS/PgoIB and CueR-α2a/PgoIB interactions were 12 ± 3 and 52 ± 12 nM, respectively. The equilibrium constants for the GolS/PcopA, CueR-α2a/PcopA, CueR/PgoIB, and GolS-α2a/PgoIB pairs could not be estimated because the low affinity of these interactions precluded the acquisition of binding data at protein saturating concentrations and thereby hindered fitting the experimental points (Fig. 3B). Nevertheless, previously estimated binding affinities for the GolS/PcopA and CueR/PgoIB heterologous interactions (by resonant mirror biosensor technology) were at least 1 and 2 orders of magnitude lower than those of the native CueR/PcopA and GolS/PgoIB interactions, respectively (15). These data indicate that, compared with the parental proteins, both the CueR-α2a and the GolS-α2a mutants exhibit stronger binding affinities for the otherwise heterologous promoters in detriment of the native target sequences, resembling the paralog regulators. Overall, these results indicate that the regulator/operator specificity resides in the α2-helix of these transcription factors.

The Amino Acid Residues at Positions 16 and 19 of the Regulator Determine Target Recognition—We analyzed the sequence differences within CueR or GolS α2-helix in order to identify the amino acid residues that direct operator recognition specificity (Fig. 4A). Considering that different members of the MerR family employ a similar DNA distortion mechanism for transcriptional activation (reviewed in Refs. 1 and 22–24), we also included in the analysis the sequence for the predicted α2-region from MtaN, BmrR, and SoxR, non-metal binding MerR proteins for which detailed structural information of the
regulator bound to their target DNA sequences is available (3–5), as well as the predicted region from the metal-responsive regulators MerR and ZntR. We focused on those residues that, according to the crystallographic studies performed on MtaN, BmrR, and SoxR, could establish hydrogen bonds or van der Waals contacts with DNA (residues at positions 15, 16, 18, 19, and 20 relative to CueR (Fig. 4A)). We observed that the Tyr-20 of CueR is conserved in these proteins, and the Ile-17 and Arg-18 residues are present in all metal-binding sensors (and not in the homologs responding to other signals), whereas the Lys residue at position 15 of both CueR and GolS is not conserved in other metal sensors. The residues at positions 16 and 19 differ between GolS and CueR. Thus, we carefully examined the identity of these residues in the different CueR-like and GolS-like homologs that we previously characterized for their operator selectivity (15). 

**FIGURE 4.** Amino acid residues at positions 16 and 19 of the α2-helix are essential for selective operator recognition. A, consensus motif for the α2-helix region of different metal-binding and non-metal-binding MerR proteins. The residues at position 16 and 19 are shaded (▽), whereas those residues conserved in the majority of the sequences are indicated in boldface type. DNA-contacting residues identified in the crystal structures of MtaN-DNA, BmrR-DNA, and SoxR-DNA complexes are highlighted by an asterisk. B, β-galactosidase activities (Miller units (M.U.)) from the PcopA or PgoIB reporter fusions as in Fig. 1 expressed on the W3110 lacZ copA cueR::cat cells carrying the expression plasmids for the indicated CueR or GolS hybrid or mutant proteins. Bacteria were grown overnight in LB (−) or in LB supplemented with 100 μM CuSO4 (Cu). The data correspond to mean values of at least three independent experiments done in duplicate. Error bars, S.D.

**FIGURE 5.** The identity of the residues at position 16 and 19 is conserved among CueR-like and GolS-like proteins. Phylogenetic tree obtained by comparison of the full-length CueR-like and GolS-like regulators. The tree was constructed by Bayesian inference as described previously (15). The α2-helix sequence for each CueR or GolS homologue was extracted and listed on the right. Residues 16 and 19 are highlighted.

![Protein Signatures for DNA Selectivity among MerR Regulators](image-url)
In view of these observations, we constructed single and double mutant versions of CueR and GolS at these positions, replacing GolS residues at position 16 and/or 19 by those present in CueR and *vice versa* and assayed their ability to activate transcription from *P* \(_{Copa}\) and *P* \(_{GolB}\) in the presence of copper. Although residues at positions 14 and 22 are not predicted to interact with the DNA, we also analyzed the role of these residues in the selective operator recognition because they differ between CueR and GolS (Fig. 4A).

The activation profile driven by either CueR\(_{A16M/F19Y}\) or GolS\(_{M16A/Y19F}\) mutants resembled the α2-variants of CueR and GolS, respectively. These mutants have an improved induction of transcription from the heterologous promoter and a diminished induction from their innate target promoter compared with the wild-type regulators (Fig. 4B). Furthermore, CueR\(_{A16M}\) and GolS\(_{M16A}\) displayed patterns of transcriptional induction similar to the double mutant proteins. Replacement of the residue at position 19 (CueR\(_{F19Y}\) and GolS\(_{Y19F}\)) had only minor effects on their operator recognition pattern (Fig. 4B). These results clearly indicate that the identity of the residue at position 16 is a main determinant of regulator/operator selectivity among *gol* and *cue* regulons. Moreover, our observations also pointed out that the residue at position 19 finely tunes the selectivity. As expected, mutant regulators with replacements at positions 14 and 22 displayed wild-type patterns of transcriptional induction (Fig. 4B), indicating that these residues are not involved in operator discrimination.

**The Residues at Position 16 and 19 of the Regulator Recognize the Selective Operator Bases 3′ and 3′—** There is currently no structural information about the target operator recognition by MerR metalloregulators. Thus, assuming that all MerR homologs interact with DNA in a similar manner, we used the available crystallographic structure of the drug-binding homolog BmrR bound to its target promoter (3) to simulate the interaction of CueR or GolS with DNA. We selected BmrR because, like CueR and GolS, it recognizes target promoters with a 19-bp spacer between the −35 and −10 elements and interacts with dyad-symmetric sequences separated by 1 bp (4, 15). In the models, we replaced the amino acid residues present in the N-terminal region of BmrR (from residue 1 to 88) by those present either in CueR, CueR-α2\(_{α}\), or GolS. As shown in Fig. 4C, the side chain of the residue at position 16 in both CueR and GolS approached the DNA backbone toward the nucleotide bases located at position 3 from the center of the operator (15). Substitution of the small methyl side chain of the alanine by a bulkier methionine will probably shift the position of the HTH motif with respect to the operator DNA. The side chain of the residue at position 19 was also oriented toward the interface between nucleotide bases at positions 3 and 4, supporting the experimental data about the role of the residue at position 16 in directing selectivity in the operator recognition.

To verify the role of amino acid residues at positions 16 and 19 in the distinction of the operator nucleotide bases at positions 3′ and 3′, we compared the promoter preferences of the mutant regulators CueR\(_{A16M/F19Y}\) and GolS\(_{M16A/Y19F}\) with the wild-type regulators, using innate *P* \(_{Copa}\) and *P* \(_{GolB}\) promoters or the mutated versions, *P* \(_{Copa}\) and *P* \(_{GolB}\), in which the operator nucleotide bases at positions 3′ and 3′ were switched by those present in the heterologous operators (15) (also see Fig. 6A). Here again, we observed that the induction of expression from the wild-type and mutated *gol* and *cue* promoters by CueR\(_{A16M/F19Y}\) is similar to that obtained using GolS, whereas the induction of the analyzed promoter by GolS\(_{M16A/Y19F}\) mimicked that of wild-type CueR (Fig. 6B). In other words, CueR\(_{A16M/F19Y}\) was more efficient in activating expression from *P* \(_{Copa}\) from wild-type *P* \(_{Copa}\). Conversely, it better recognized wild-type *P* \(_{GolB}\), which harbors A and T at position 3′ and 3′, respectively, than *P* \(_{GolB}\), which harbors C at these two positions, exhibiting a pattern of induction similar to GolS. Similarly, GolS\(_{M16A/Y19F}\) better recognized promoters with Cs at positions 3′ and 3′, such as the native *P* \(_{Copa}\) or the mutant.
**Protein Signatures for DNA Selectivity among MerR Regulators**

PgoiB\textsubscript{CC} promoters, than promoters with A and T at these positions (i.e. PgoiB and PcpoA\textsubscript{CC}). Overall, these results established the importance of the amino acid residue at position 16 and, to a lesser extent, at position 19 in the selection of the *Salmonella* cue and gol regulon target operators.

**DISCUSSION**

Transcription factors must be capable of locating their specific target sequences along the chromosome, avoiding their unproductive or even harmful interaction at ectopic places. This appears particularly relevant when similar regulatory proteins with almost identical target recognition operators coexist in a single cell. *Salmonella* has two structurally related but functionally distinctive metalloregulators of the MerR family that orchestrate the cellular response to the presence of toxic amounts of monovalent metal ions in the environment (25). By rewiring both the input signal detection and the recognized operator sequences, the horizontally acquired gold sensor GolS is able to induce the expression of its target genes without interfering with the function of the copper homeostasis regulator CueR. In a previous report, we demonstrated that selectivity in the recognition of GolS or CueR target operators is achieved by subtle modifications of the operator sequences (regulon signature nucleotide bases are displayed at the 3 and 3’ operator positions (15)), which are accompanied, as is shown in this work, by subtle modifications at the DNA-binding motif of these transcriptional regulators.

We constructed a set of hybrid proteins between GolS and CueR to identify the region that directs operator recognition (Fig. 1 and 2). These studies allowed us to focus on the α2-helix (from residue 14 to 22), which can be defined as the minimal region necessary for operator discrimination among the gol and cue regulons. *In vitro* experiments confirmed these observations, showing that the solely replacement of this motif in each regulator lowered its affinity for its innate promoters and increased its affinity for the paralog operators (Fig. 3).

There is no structural information on any metal-sensing MerR regulator bound to its target operator, and based on the available biochemical and genetic data, it was postulated that the DNA distortion mechanism for transcriptional activation is conserved among all family members (reviewed in Refs. 1 and 22–24). Therefore, we assumed that CueR and GolS interact with their target sequences in a similar manner as the non-metal binding MerR homologs BmrR and MtaN, from which the crystallographic structure of protein-DNA complexes is available (3, 5). In other words, the axis of symmetry of the CueR/GolS dimer is facing the minor groove at the center of the palindromic sequences and have the 3–3’ signature (C/G)(C/G) have Ala, Ser, or Thr at position 16, and the DNA distortion mechanism for transcriptional activation is corroborated by the observation that double mutant regulators CueR\textsubscript{A16M/F19Y} and GolS\textsubscript{M16A/Y19F} were as effective as the hybrid CueR-α2S and GolS-α2\textsubscript{M} respectively, to activate the promoters controlled by the paralog regulator (Fig. 4B). In addition, CueR\textsubscript{A16M/F19Y} and GolS\textsubscript{M16A/Y19F} activated transcription of the reporter gene from the promoter with its 3’ and 3 bases replaced more efficiently than from the wild-type promoters (Fig. 6). We hypothesize that the failure of GolS to recognize operators bearing C/G substitutions at positions 3 and 3’ could be explained by the formation of a third hydrogen bond between the pairing CG bases, which renders a base pair less deformable than the AT base pair. This would result in either steric or electrostatic interference with the interaction of GolS, which has the more voluminous yet hydrophobic methionine residue at position 16.

The presence of the distinctive Met residue at the α2-helix of the DNA-binding domain of GolS is extended to all GolS-like proteins having the 3–3’ AT signature in the operators of genes predicted to be controlled by them (Fig. 5) (also see Ref. 15). By contrast, CueR homologs that recognize promoters with the signature (C/G)(C/G) have Ala, Ser, or Thr at position 16, and indeed, replacement of the Ala-16 residue of CueR by Thr did not affect its operator selectivity (Fig. 4B). As expected, the residue at position 19 is not conserved, but interestingly, close GolS homologs harbor a Tyr at this position, whereas all CueR homologs have a Phe. Indeed, the recently characterized *Cupriavidus metallidurans* CH34 gold sensor CupR, which controls genes with the characteristic 3–3’ AT operators (26), harbors the GolS-like Met-Phe signature at the N-terminal DNA-binding domain. Notoriously, several MerR homologs that recognize promoters with the signature (C/G)(C/G) have Ala or Thr at position 19, whereas CueR-like and GolS-like regulators rely exclusively on the interaction of the amino acid residue at position 16 and the signature nucleotide base at position 3 of the operator.

Overall, these studies suggest that, along evolution, rewiring of both the transcriptional regulators GolS and CueR and the regulatory elements in their target genes confers novel abilities to detect distinct environmental cues, avoiding at the same time cross-regulation that would jeopardize the adequate response to a specific stress.

Acknowledgment—We thank Eleonora García Véscovi for valuable experimental advice and critical reading of the manuscript.
