

## Functional Similarities between the *Listeria monocytogenes* Virulence Regulator PrfA and Cyclic AMP Receptor Protein: the PrfA\* (Gly145Ser) Mutation Increases Binding Affinity for Target DNA

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Most *Listeria monocytogenes* virulence genes are positively regulated by the PrfA protein, a transcription factor sharing sequence similarities with cyclic AMP (cAMP) receptor protein (CRP). Its coding gene, *prfA*, is regulated by PrfA itself via an autoregulatory loop mediated by the upstream PrfA-dependent *plcA* promoter. We have recently characterized *prfA*\* mutants from *L. monocytogenes* which, as a result of a single amino acid substitution in PrfA, Gly145Ser, constitutively overexpress *prfA* and the genes of the PrfA virulence regulon. Here, we show that about 10 times more PrfA protein is produced in a *prfA*\* strain than in the wild type. Thus, the phenotype of *prfA*\* mutants is presumably due to the synthesis of a PrfA protein with higher promoter-activating activity (PrfA\*), which keeps its intracellular levels constantly elevated by positive feedback. We investigated the interaction of PrfA and PrfA\* (Gly145Ser) with target DNA. Gel retardation assays performed with a DNA fragment carrying the PrfA binding site of the *plcA* promoter demonstrated that the PrfA\* mutant form is much more efficient than wild-type PrfA at forming specific DNA-protein complexes. In footprinting experiments, the two purified PrfA forms interacted with the same nucleotides at the target site, although the minimum amount required for protection was 6 to 7 times lower with PrfA\*. These results show that the primary functional consequence of the Gly145Ser mutation is an increase in the affinity of PrfA for its target sequence. Interestingly, similar mutations at the equivalent position in CRP result in a transcriptionally active, CRP\* mutant form which binds with high affinity to target DNA in the absence of the activating cofactor, cAMP. Our observations suggest that the structural similarities between PrfA and CRP are also functionally relevant and support a model in which the PrfA protein, like CRP, shifts from transcriptionally inactive to active conformations by interaction with a cofactor.

Virulence genes in the gram-positive, facultative intracellular pathogen *Listeria monocytogenes* are regulated by the pleiotropic transcriptional activator PrfA, encoded by the *prfA* gene (6, 8, 21, 25, 27). An ambient temperature of 37°C is necessary for the transcriptional activation of *prfA* and PrfA-dependent genes (24). This is, however, not sufficient for the full activation of the PrfA regulon. Wild-type strains express PrfA-regulated genes to a very low level in rich media (e.g., brain-heart infusion medium [BHI]) at 37°C (30), but strongly activate their transcription if cultured in BHI treated with activated charcoal (28–30) or if transferred from BHI to minimal essential medium (5). This requirement for a suitable combination of environmental signals of a physical and chemical nature may be a fail-safe mechanism used by *L. monocytogenes* to prevent the expression of virulence genes in situations in which they are not required, i.e., when the bacteria are outside an appropriate host niche. Recent observations have suggested that there is also a mechanism of negative regulation in *L. monocytogenes* which abolishes the expression of virulence genes in the presence of readily fermentable carbon sources, such as glucose or

cellobiose (26, 28). The molecular basis and biological relevance of this repression mechanism are unknown.

The primary structure of PrfA has significant similarities to that of *Escherichia coli* cyclic AMP (cAMP) receptor protein (CRP) and other members of the CRP-FNR family of bacterial transcription factors (21, 23). PrfA has, for example, a helix-turn-helix (HTH) motif in the C-terminal region, at the same position as in CRP and related proteins. This HTH motif has been shown to interact specifically with target DNA sequences called “PrfA-boxes,” which are 14-bp-long palindromes centered at position –41 relative to the transcription start site in PrfA-dependent promoters (3, 9, 11, 33). Binding to these PrfA-boxes is affected by the number of nucleotide mismatches they carry, becoming weaker as the sequence diverges from the perfect palindrome (4, 12, 34). The symmetrical structure of PrfA-boxes suggests that like CRP, PrfA binds to target DNA as a dimer, and there is experimental evidence that PrfA forms a homodimer in solution (9).

Evidence that PrfA and CRP are functionally related has been provided by our recent characterization of *prfA*\* mutants from *L. monocytogenes* (28, 29, 31). Mutatis mutandis, these *prfA*\* strains are analogous to the *crp*\* mutants of *E. coli* in that they constitutively overexpress *prfA* and PrfA-dependent genes under culture conditions in which the PrfA regulon is normally downregulated (e.g., at 37°C in BHI), to levels

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reached by wild-type strains only if cultured in charcoal-treated BHI (28–30). These *prfA*\* mutants carry a Gly→Ser substitution in residue 145 of PrfA that seems to increase the transcriptional activity of the regulator, releasing it from a variety of repressor signals including low temperature and growth on glucose or cellobiose (28–30). This mutation is located in a PrfA region of 11 amino acids (residues 141 to 151) with a sequence very similar (70% similarity) to that of the D  $\alpha$ -helix of CRP (29). Several *crp*\* mutations in *E. coli* that allow CRP to function in the absence of cAMP, the cofactor required for its allosteric activation, also map in this region (13, 15a, 20). One such CRP\* mutation, Ala144Thr, which presumably mimics the conformational change caused by the cofactor (19, 20), maps in the aligned proteins to the position equivalent to that of the Gly→Ser PrfA mutation (29). These observations led us to hypothesize that PrfA functions via a cofactor-mediated allosteric transition mechanism similar to that of CRP, and that the Gly145Ser mutation is a cofactor-independent PrfA\* form that is “frozen” in an active conformation (29).

In this study, we investigated the interaction of wild-type PrfA and mutant PrfA\* (Gly145Ser) with target DNA. As for CRP\* altered forms (2, 32, 35), the Gly145Ser mutant protein bound with higher affinity to specific DNA than did the wild-type protein, further supporting the notion that PrfA is a structural and functional homolog of CRP.

#### MATERIALS AND METHODS

***L. monocytogenes* strains and culture conditions.** P14, an *L. monocytogenes* wild-type strain of serovar 4b, and its *prfA*\* mutant, P14-A, have been described in detail elsewhere (28–31). *L. monocytogenes* EGD, a wild-type strain of serovar 1/2a, and its *prfA* deletion mutant,  $\Delta$ *prfA*, have also been previously described (3, 5, 29). They were grown in BHI broth at 37°C with shaking.

**General DNA techniques.** Restriction enzymes were purchased from Pharmacia and used as recommended by the manufacturer. The Expand high-fidelity PCR system (Boehringer Mannheim) was used to amplify specific DNA fragments. PCR products were purified from gels with the Qiaquick (Qiagen) gel extraction kit. Plasmid DNA was extracted from *E. coli* with a plasmid purification kit from Qiagen. DNA sequencing was performed with an Applied Biosystems 377 apparatus.

***L. monocytogenes* cell protein extracts, SDS-PAGE, and anti-PrfA immunoblotting.** Soluble protein extracts from *L. monocytogenes* were prepared and stored as described by Böckmann et al. (3). Total protein concentration was determined with the Bio-Rad Protein-Microassay. Sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) was performed with 12% acrylamide slab gels as described by Laemmli (22). For immunoblotting, proteins were electrotransferred from the gels to nitrocellulose sheets (Schleicher & Schuell), and PrfA was detected by using a previously described anti-PrfA polyclonal hyperimmune serum (3), peroxidase-conjugated secondary antibodies, and 4-chloro-1-naphthol.

**Expression in *E. coli* and purification of wild-type and Gly145Ser mutant PrfA proteins.** The *prfA* and *prfA*\* alleles from P14 and P14-A, respectively, were amplified by PCR with the oligonucleotide pair N-PR1 (5'-ATGACTCGAGACGCTCAAGCAGAAGAA-3') and C-PR1 (5'-CTGTAGATCTTTAATTTAATTTCCCA-3'), which contain *Xho*I (N-PR1) and *Bgl*II (C-PR1) restriction sites (underlined). The resulting *prfA*-containing DNA fragments were cloned in *E. coli* with the pMOSBlue T-vector kit (Amersham), and then transferred to pFLAG-MAC expression vector (Sigma) by using the *Xho*I and *Bgl*II sites, giving rise to the plasmids pF-PrfA and pF-PrfA\*(G145S). A fusion was created in these plasmids, resulting in a sequence that encodes a recombinant PrfA protein with an N-terminal tag of 14 amino acids including an 8-mer peptide marker (FLAG epitope). The whole open reading frame was checked by sequencing both strands in each expression plasmid. Recombinant PrfAs were overproduced in *E. coli* DH5 $\alpha$ . Host bacteria were grown at 37°C in 500 ml of Luria-Bertani medium containing ampicillin (50  $\mu$ g/ml) until the optical density at 600 nm was 1.0, and expression was induced by adding 1 mM IPTG (isopropyl- $\beta$ -D-thiogalactopyranoside). After 3 h, the induced bacteria were pelleted and lysed by suspension in 20 ml of lysis buffer A (50 mM Tris-HCl [pH 8], 5 mM EDTA, 50  $\mu$ g of sodium azide per ml, 0.25 mg of lysozyme per ml) and addition of 2 ml of lysis buffer B (1.5 M NaCl, 100 mM CaCl<sub>2</sub>, 100 mM MgCl<sub>2</sub>, 0.02 mg of DNase I per ml, 50  $\mu$ g of phenylmethylsulfonyl fluoride). Recombinant PrfAs, which did not form inclusion bodies in *E. coli*, were purified from the bacterial soluble extract by column affinity chromatography with anti-FLAG M<sub>2</sub> monoclonal antibody resin (Sigma), according to the manufacturer's instructions. The eluted fractions were analyzed by SDS-PAGE with Coomassie blue staining. The fractions containing >98% pure PrfA (which migrated as a band of 28.5 kDa, 1.5

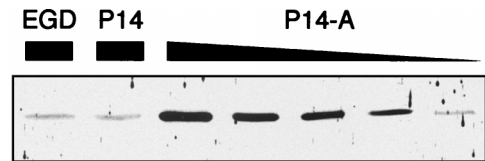


FIG. 1. Determination of PrfA protein in *L. monocytogenes* P14 (wild type), P14-A (*prfA*\* [Gly145Ser] mutant from P14), and EGD (control wild-type strain). Total cell extracts from these strains were subjected to SDS-PAGE in a 12% acrylamide gel (protein amounts loaded: P14 and EGD, 30  $\mu$ g; P14-A [from left to right], 30, 15, 10, 5, and 2.5  $\mu$ g) and analyzed by Western immunoblotting with an anti-PrfA hyperimmune serum. The PrfA protein is detected as a 27-kDa band. Note that equivalent amounts of PrfA protein are present in 30  $\mu$ g of the P14 and EGD extracts and 2.5  $\mu$ g of the P14-A extract.

kDa larger than predicted from the *prfA* sequence due to the presence of the extra 14 N-terminal amino acids) were collected, concentrated by Centricon devices (Amicon), and preserved at  $-20^{\circ}\text{C}$  with 20% glycerol.

**DNA mobility shift and footprinting assays.** A 136-bp double-stranded PCR fragment containing the *plcA*-*hly* promoter region was used as target DNA. It was amplified from strain P14 with primers YV3 (5'-TCCTATCTAGAAGTTA CTTTATGTC-3') and YV4 (5'-TATTGGATCCATTCGCTTCTAAAGATG-3'), which contain *Xba*I and *Bam*HI restriction sites (underlined). Previously described protocols were used for electrophoretic mobility shift assays with *L. monocytogenes* protein extracts or purified PrfA proteins (3). DNase I footprinting experiments were performed as previously described (9).

#### RESULTS

**Amounts of PrfA in the wild-type and the *prfA*\* (Gly145Ser) mutant of *L. monocytogenes*.** The levels of expression of *prfA* are primarily controlled by the PrfA-dependent *plcA* promoter, from which a bicistronic transcript covering the *plcA*-*prfA* operon is generated. This *plcA*-*prfA* mRNA creates an autoregulatory loop that is essential for the normal function of the PrfA regulon, presumably because it ensures the synthesis of sufficient quantities of the PrfA protein (5, 7, 12, 24, 25, 29) (see Fig. 6). Even if the *prfA* gene remains intact, any interruption of this autoregulatory loop (e.g., by insertional mutagenesis in *plcA* or in the *plcA*-*prfA* intergenic region) leads to a PrfA<sup>-</sup> phenotype (7, 11, 25, 28, 29). *trans*-complementation experiments have suggested that the mutant form of PrfA synthesized from *prfA*\* (Gly145Ser), PrfA\*, is more effective than the wild-type protein at activating PrfA-dependent promoters (28, 29). This would result in PrfA\* constantly switching on the autoregulatory loop such that more PrfA protein was produced in the mutant *prfA*\* background than in the wild type. We tested this by analyzing cell extracts of the *L. monocytogenes* wild-type strain, P14, and its *prfA*\* mutant, P14-A, grown in BHI at 37°C, by Western blotting with anti-PrfA antibodies.

There was clearly more PrfA protein in P14-A than in P14 (Fig. 1), suggesting that PrfA\* did indeed activate its own synthesis by a positive feedback mechanism. The constitutive overexpression of PrfA-dependent virulence genes in *prfA*\* mutants is therefore presumably due to the sustained production of high levels of a transcriptionally active PrfA\* form. Densitometric analysis of the blots showed that the intracellular levels of PrfA\* were around 10 times higher in P14-A than in P14.

**DNA-protein complex formation by PrfA and PrfA\* (Gly145Ser) in cell extracts of *L. monocytogenes*.** We investigated whether the differences in virulence gene transcription in the wild type and *prfA*\* mutants (29, 30) were due to differences in the DNA-binding activity of the corresponding PrfA and PrfA\* proteins. Electrophoretic mobility shift assays were carried out with cell extracts from P14 and P14-A and a 136-bp PCR fragment containing the *plcA* promoter and its PrfA-box,

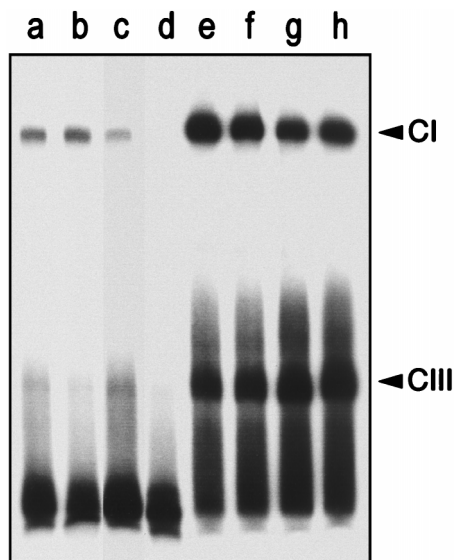


FIG. 2. Electrophoretic mobility shift assays with a 136-bp DNA fragment containing the PrfA-box of the *plcA-hly* promoter region and *L. monocytogenes* cell extracts. Lanes: a and b, P14 (30 and 60  $\mu$ g, respectively); c, EGD (30  $\mu$ g); d,  $\Delta$ *prfA* mutant from EGD (30  $\mu$ g); e to h, P14-A (30, 15, 10, and 5  $\mu$ g, respectively). Lanes b and h contain equal amounts of PrfA protein (Fig. 1). CI and CIII, respectively, low- and high-mobility specific PrfA-DNA complexes. (See text and references 3 and 9 for details.)

used as the target DNA. This PrfA-box is shared with the divergently transcribed *hly* gene and represents the "perfect" palindrome, to which PrfA presumably binds with maximal affinity (4, 34). Extracts from *L. monocytogenes* EGD, a wild-type strain in which the deduced amino acid sequence of PrfA is identical to that of P14 (29), and its *prfA* deletion mutant (EGD  $\Delta$ *prfA*) were used as controls.

Specific protein-DNA complexes (CI) (3, 9) were formed with the cell extracts from all of the PrfA-proficient strains used (Fig. 2). However, there was a higher level of complex formation with the PrfA\*-containing extract than with that containing wild-type PrfA, as determined from the intensities of the CI PrfA-dependent complexes formed (see lanes b and h, in which the amounts of PrfA protein are equivalent). The level of PrfA-dependent complex formation with extracts from EGD was identical to and as low as that with the extract from the P14 wild-type strain (lanes a and c). As expected, there was no binding activity observed with the EGD  $\Delta$ *prfA* extract (lane d). Therefore, the higher transcriptional activity of the PrfA\* (Gly145Ser) mutant form correlates with a higher affinity for target DNA.

**Interaction of purified PrfA and PrfA\* (Gly145Ser) with target DNA.** We characterized the differential interaction of PrfA and PrfA\* (Gly145Ser) with the target DNA in more detail by gel retardation assays with the purified proteins. The PrfA proteins from strains P14 and P14-A were produced in *E. coli* with a FLAG epitope fused to the N terminus, which allowed them to be purified by affinity chromatography with an anti-FLAG-peptide monoclonal antibody (see Materials and Methods). The recombinant purified proteins were called F-PrfA and F-PrfA\*, respectively. Addition of an N-terminal tag to PrfA has been shown to have no major effect on the DNA-binding function of the protein (3, 9). We used a recently described protocol with which direct, specific binding of purified PrfA can be observed in the absence of additional factors

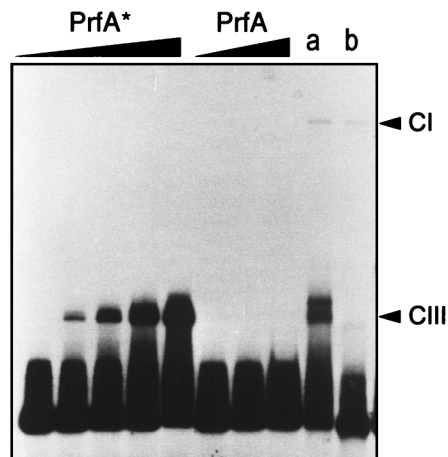


FIG. 3. Binding of the purified PrfA proteins to the *plcA-hly* promoter fragment. Various amounts of PrfA preparation were used (from left to right: PrfA\* [Gly145Ser], 0.5, 1.5, 3, 6, and 18 ng; PrfA, 120, 240, and 480 ng). Lanes a and b: low-mobility CI-specific protein-DNA complex formation by purified PrfA\* and PrfA proteins (50 ng each), respectively, in the presence of a PrfA-free *L. monocytogenes* cell extract (30  $\mu$ g) from EGD  $\Delta$ *prfA*. CIII, high-mobility specific PrfA-DNA complexes. (See text and references 3 and 9 for details.)

from the listerial cytoplasm (9). In this case, a high-mobility DNA-PrfA complex (CIII) is formed (9).

As little as 1.5 ng of F-PrfA\* was sufficient to produce a visible CIII complex with the 136-bp DNA fragment containing the *plcA-hly* PrfA-box (Fig. 3). This interaction was specific, as shown by the ability of the unlabeled specific probe and the inability of nonspecific DNA to compete out CIII complex formation (Fig. 4). In contrast, no mobility shift was detectable with F-PrfA, even at high protein concentrations (Fig. 3). Addition of PrfA-free *L. monocytogenes* extract (from EGD  $\Delta$ *prfA*) led to CI complex formation by F-PrfA\* and, also, by F-PrfA (Fig. 3). Therefore, although it cannot directly interact with DNA to form a visible CIII complex, purified wild-type PrfA is able to bind to its target site in the presence of additional factors from the listerial extract. These results show that the PrfA\* form is clearly more efficient than the wild-type protein at establishing direct interaction with the PrfA-specific target site.

We investigated whether the higher DNA-binding activity of PrfA\* was associated with a different pattern of interaction at

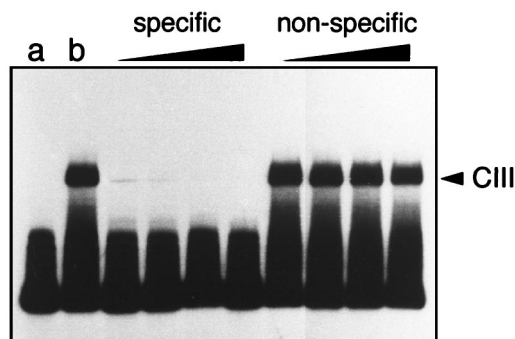


FIG. 4. Specificity of the interaction of PrfA\* (8 ng) with target DNA resulting in CIII complex formation. Competition assays with (from left to right) 50-, 100-, 200-, and 400-fold molar excess of specific (unlabeled 136-bp *plcA-hly* promoter fragment) and nonspecific DNA (from herring sperm). Lanes: a, control with the labeled 136-bp *plcA-hly* promoter fragment alone; b, control with the labeled probe plus 8 ng of purified PrfA\*.



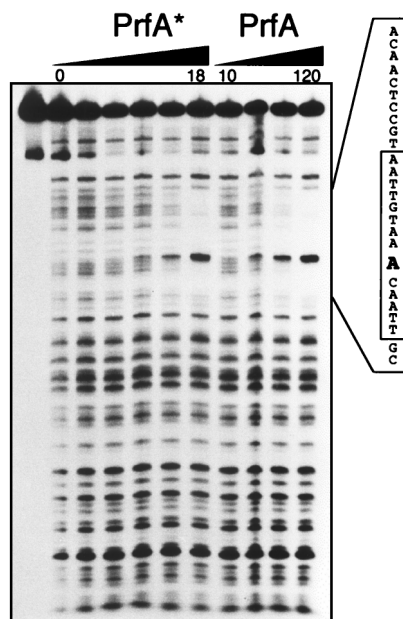


FIG. 5. DNA footprinting experiments with various amounts of the purified PrfA proteins (PrfA\*, 0, 0.5, 1.5, 3, 6, and 18 ng; PrfA, 10, 20, 40, and 120 ng) and the *plcA-hly* promoter fragment. The protected sequences, identical for the two PrfA proteins, are indicated on the right. The palindromic PrfA binding site is boxed, and numbers indicate the nucleotide position with respect to the transcription start site of the *hly* mRNA. The hypersensitive nucleotide (A) at position -40, close to the center of the palindrome, is in boldface. To the left is shown the uncleaved probe.

the PrfA site by footprinting analysis of the DNA sequence protected by F-PrfA and F-PrfA\* in the *plcA-hly* promoter region (Fig. 5). Various amounts of the purified PrfAs were used to further evaluate the differential binding affinity of the two proteins for the target site. In contrast to the results obtained with gel retardation assays, we did detect direct interaction of F-PrfA with target DNA. This probably results from the different experimental conditions for the two techniques, particularly the relatively high concentration of poly(dI-dC), used in the mobility shift experiments as nonspecific competitor to minimize nonspecific or low-affinity protein-DNA interactions (9). However, the amount of purified wild-type protein required for complete protection was significantly higher (6 to 7 times) than that for F-PrfA\*. The DNA region protected from DNase I digestion was exactly the same for both purified proteins (positions -58 to -33 relative to the transcriptional start site of *hly*, including the PrfA-box palindrome and 10 bp upstream and 2 bp downstream from it) (Fig. 5) and was concordant with that previously reported for PrfA (9).

## DISCUSSION

We have shown, by electrophoretic mobility shift and footprinting assays, that (i) wild-type PrfA interacts weakly with the specific target DNA, and (ii) a mutant PrfA form, PrfA\* (Gly145Ser), has a much higher DNA-binding activity than the wild-type protein. There was no difference between the DNA sequences footprinted by the mutant and wild-type PrfA proteins, demonstrating that the primary functional consequence of the Gly145Ser substitution is a very significant increase in binding affinity for the target DNA. Since *prfA* is positively regulated by its own product, PrfA (24, 25, 28, 29) (Fig. 6), the expected outcome of the mutation in vivo is an increase in the

levels of the PrfA protein. This has been also demonstrated herein. Our results are consistent with the physiological properties of wild-type and *prfA\** (Gly145Ser) backgrounds of *L. monocytogenes*, which in normal culture media express low and constitutively high levels of PrfA-dependent genes, respectively (28–30).

PrfA and CRP exhibit significant similarities at the level of primary structure (21, 23, 33). The observations here reported with PrfA are also very similar to those for the structure-function relationships of CRP, supporting the notion that PrfA acts via a regulatory mechanism similar to that of the *E. coli* transcription factor. The inactive form of CRP binds to specific DNA with very low affinity, so the interaction is not normally detectable in gel retardation assays. However, if complexed with the activating cofactor, cAMP, CRP undergoes a conformational change that is associated with a dramatic increase in affinity for the target DNA (2, 16, 17, 35). Binding of the cofactor is thought to alter intersubunit alignment and interdomain orientation in CRP, ultimately resulting in the protrusion of the F  $\alpha$ -helix which is part of the HTH DNA-binding motif, thereby facilitating productive specific protein-DNA interaction (1, 14, 16, 19, 20). CRP\* mutations in the D  $\alpha$ -helix, which spans residues 139 to 150, close to the hinge region connecting the N-terminal cAMP-binding domain and the C-terminal DNA-binding domain of the CRP subunit (13, 15a), are thought to evoke the conformational change caused by cAMP, resulting in transcriptional activation in the absence of significant amounts of the cofactor (14, 19, 20, 32). In this D  $\alpha$ -helix, any amino acid substitution introducing a larger side chain at position 144, which aligns with PrfA residue 145 (the site of our PrfA\* mutation) (29), results in a cAMP-independent CRP\* phenotype (19). According to the determined crystal structure of CRP, amino acid 144 faces residue 190 in the F  $\alpha$ -helix (20, 37). Therefore, the cAMP-independent phenotype is presumably due to the larger side chain pushing the DNA-binding sequence outward (19, 20). The PrfA\* mutation studied, mapping to a region that is remarkably homologous to the D  $\alpha$ -helix of CRP (29), is similar to the CRP\* mutation previously characterized at position 144, Ala to Thr (14, 15a, 20), and involves the replacement of a small amino acid (Gly) with a larger one (Ser). A second CRP\* mutation described at position 141 in the D  $\alpha$ -helix is also a Gly→Ser substitution, and we recently characterized another PrfA\* mutation that maps nearby within the same D  $\alpha$ -helix-homologous region, which also involves a replacement by a bulkier amino acid (36). Except for the fact that PrfA has an extra 25 amino acids at the extreme C terminus, the C-terminal domain of CRP is very similar (45% identity, 60% similarity from amino acid 128 to amino acid 201) to the corresponding region in PrfA (21, 23, 36). These observations suggest that the mutations resulting in the PrfA\* phenotype are associated with conformational changes in the DNA-binding domain similar to those that are thought to occur in CRP\* mutant proteins.

Our findings provide support for our model of PrfA-dependent regulation (29). In this model, similar to that proposed for CRP, PrfA undergoes an allosteric transition from an inactive to an active conformation upon interaction with an environmentally regulated low-molecular-weight cofactor. A key element of this model is the positive autoregulatory circuit of *prfA*, an aspect in which the listerial regulatory gene also resembles the transcriptional control mediated by *crp* in *E. coli* (15). (See Fig. 6 for a detailed description of the model.)

Two typical features of CRP are particularly well conserved in PrfA. One is the HTH motif in the C-terminal region, for which the functional similarity between the two proteins has been already documented (33). The other is a series of short

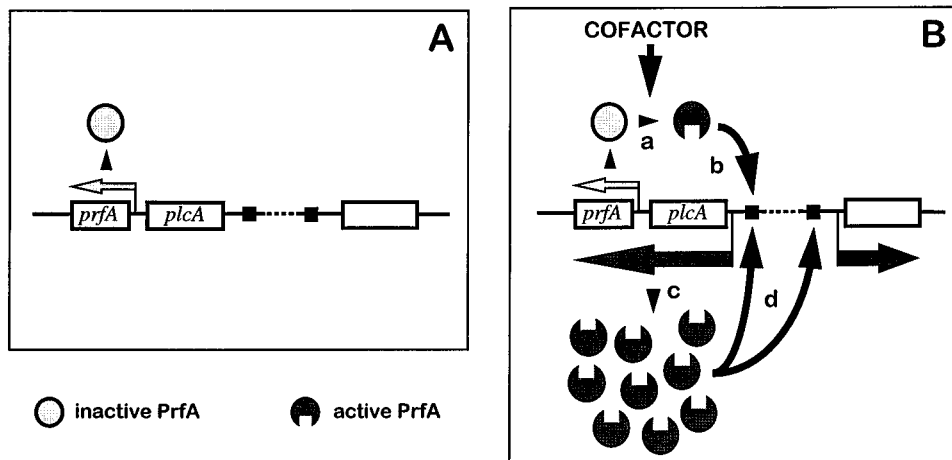


FIG. 6. Model for PrfA-mediated gene regulation (29). Central to this model is the assumption that PrfA has two functional conformations, inactive and active, and shifts from one to the other on interaction with a hypothetical activating cofactor, the intracellular concentrations of which depend on the environmental conditions. A key element is also that *prfA* can be expressed in two different ways: (i) constitutively and at low levels, from monocistronic transcripts driven by promoters in the *plcA-prfA* intergenic region (represented by a small light-gray arrow above *prfA* on panels A and B); and (ii) dependent on PrfA, from bicistronic transcripts originating from the *plcA* promoter (dark gray arrow below *plcA* and *prfA* on panel B), thereby creating an autoregulatory circuit. The regulation mechanism would be as follows. Under normal conditions, there is no cofactor, and, thus, the PrfA protein is synthesized at low, basal levels from the monocistronic transcripts (A). However, if *L. monocytogenes* senses a suitable combination of activating environmental signals (a temperature of 37°C and a particular composition of the extracellular medium), the intracellular concentration of the hypothetical cofactor increases (B). This cofactor interacts with the inactive PrfA protein synthesized from monocistronic transcripts (a), causing a conformational change that results in a significant increase in the binding affinity of PrfA for its target DNA (b) (PrfA sites are indicated by black squares). The transcriptionally active PrfA causes the synthesis of more PrfA (in active conformation) by positive feedback (c), which boosts the transcription of all the PrfA-dependent genes (d) (dark gray arrows; the empty rectangle represents any PrfA-dependent gene). The PrfA regulon remains switched on as long as there are sufficiently high levels of the cofactor in the bacterial cytoplasm, but the system is rapidly switched off if the activating environmental signals cease and the concentration of the cofactor drops. A second level of regulation is provided by the differential response of the PrfA-dependent promoters according to the structure of the PrfA target site, which affects the binding affinity of PrfA. (See references 4, 7, 11, 12, and 34 for details about this *cis*-acting control mechanism.) Evidence for a negative autoregulation mechanism involving a putative PrfA-binding site in the *plcA-prfA* intergenic region has been also presented (11, 12), which would add complexity to the transcriptional control mediated by PrfA. The proposed regulatory model is highly versatile and makes possible an immediate, fine-tuned adaptive response to rapidly changing environmental conditions, such as those encountered by the soil bacterium *L. monocytogenes* during its transition from free to parasitic life and within the various compartments and tissues of the infected host.

antiparallel  $\beta$ -strands delimited by glycine residues, which may form a  $\beta$ -roll structure involving most of the N-terminal half of the protein (21, 23). The prediction of such a structure in PrfA is quite intriguing, because in CRP it forms the pocket in which the activating cofactor, cAMP, is buried in the N-terminal domain of the protein (20). However, cAMP is undetectable, and it is not known to function as an effector molecule in gram-positive bacteria (18). In fact, most of the residues in CRP that are important for cAMP binding are not conserved in PrfA (21, 36), and addition of exogenous cAMP does not result in PrfA activation (36). It is, however, unknown whether cAMP is taken up by *Listeria*. Preliminary studies with the extrinsic fluorescence probe 8-anilino-1-naphthalenesulfonic acid (ANS) have shown that as for CRP (16), the addition of cAMP to purified PrfA results in a significant fluorescence quenching (36). This indicates that cAMP induces a conformational change, but not necessarily that it allosterically activates PrfA. The cyclic nucleotide cGMP, for example, does not functionally activate CRP, but it does bind to it with an affinity similar to that of cAMP, and there are cAMP analogs that bind to CRP and cause a conformational change similar to that elicited by cAMP but do not activate transcriptional function (10). It is therefore possible that the putative cofactor for PrfA is a cyclic nucleotide similar to cAMP. We are currently working toward the identification of this putative PrfA cofactor and the genetic characterization of the signal transduction machinery that connects the PrfA system with the extracellular environment.

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