

Mapping the Transcription Start Points of the *Staphylococcus aureus* *eap*, *emp*, and *vwb* Promoters Reveals a Conserved Octanucleotide Sequence That Is Essential for Expression of These Genes^{∇†}

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Mapping the transcription start points of the *eap*, *emp*, and *vwb* promoters revealed a conserved octanucleotide sequence (COS). Deleting this sequence abolished the expression of *eap*, *emp*, and *vwb*. However, electrophoretic mobility shift assays gave no evidence that this sequence was a binding site for SarA or SaeR, known regulators of *eap* and *emp*.

The ability of *Staphylococcus aureus* to cause such diverse infections as endocarditis, pneumonia, skin infections, and biofilms is linked to its great repertoire of virulence factors, including adhesins, immunomodulatory molecules, and toxins (31). The *S. aureus* cell surface adhesins belong to one of two groups, the MSCRAMMs (microbial surface components recognizing adhesive matrix molecules), which include protein A, clumping factors A and B, and the fibronectin-binding proteins; and the SERAMs (secretable expanded repertoire adhesive molecules), which include the extracellular adherence protein (Eap), the extracellular matrix binding protein (Emp), and the extracellular fibrinogen-binding protein (Efb) (reviewed in references 5 and 7). The MSCRAMMs contain an LPXTG motif, which is involved in anchoring them to the staphylococcal cell surface (10, 32, 45), while the SERAMs lack this motif and may bind to the staphylococcal cell surface either covalently or via specific cell surface receptors (e.g., see references 11, 25, and 35). Together the MSCRAMMs and SERAMs facilitate the attachment of *S. aureus* to eukaryotic cells, platelets, extracellular matrix proteins, and inert surfaces (reviewed in reference 43) and may aid the survival and persistence of the staphylococci in the host due to their ability to interfere with the host's immune response (1, 4, 15, 22, 28, 30, 41, 44, 47).

Eap, Emp, and von Willebrand factor-binding protein (vWbp) are members of the SERAM family (5). While Eap is functionally well characterized (18), less is known about Emp and vWbp. Emp was described as an extracellular matrix binding protein, but additional functional roles have not yet been described (23). vWbp was identified during a screen for factors binding von Willebrand factor and was subsequently shown to be a coagulase (2, 3). Although the members of the SERAM family do not share significant sequence homology, they are

recognized as sharing similar functional properties, such as being important in adhesion and modulation of the host immune response to staphylococcal infections (1–3, 5, 12, 16, 17, 23, 29, 30, 42, 44, 47). What is not yet known, however, is whether the regulation of the SERAMs at the molecular level is governed by a common mechanism or factor.

We became interested in studying the regulation of *emp* and *vwb* as these two genes, together with *clfA*, are located adjacent to each other (*S. aureus* strain COL open reading frames SACOL0856-SACOL0858), with 223 bp separating *clfA* from *vwb* and 353 bp separating *vwb* from *emp*. As ClfA is also an important *S. aureus* virulence factor, an interesting scenario would be the cotranscription of *clfA*, *vwb*, and *emp*. We mapped the transcription start points of *emp* and *vwb* in *S. aureus* strain Newman using primer extension analysis as described in reference 19 and found that both genes had their own promoter (Fig. 1A). These findings suggest that the genes are not cotranscribed and fit with the observations by us and others that the expression profiles of the three genes are different (3, 19, 34, 46). *eap* was also found to be transcribed from a single promoter (Fig. 1A). Putative promoter elements were identified by analysis of the region upstream of the transcription start point. All three promoters have a conserved –10 box (Fig. 1B), but homology to the consensus –35 box, TTGACA (20, 33, 36, 37), is less conserved, particularly in the *vwb* promoter. However, we have found a conserved octanucleotide sequence (COS), AGTTAATT, that is just 5' to a putative –35 box in each promoter (Fig. 1B). Moreover, searching the *S. aureus* COL genome for this COS revealed a COS in the same position (i.e., immediately upstream of a putative –35 box) in the promoters of several important virulence factors (Table 1). A common feature of these virulence factors is that they are involved in modulating the immune response to *S. aureus* infections or antibiotic resistance (5, 9, 24, 48). Taken together, these data suggested that the COS could be important in the regulation of these genes.

To investigate the importance of the COS, we deleted it in the *eap*, *emp*, and *vwb* promoters in a two-step PCR. For deleting the COS in the *emp* promoter, two primer pairs were used. Primers *emp*-cs_R (5'-GTTTACTTCAATTATACTGA AAATTC-3') and *emp*-cs_F (5'-GAATTTTCAGTATAATT

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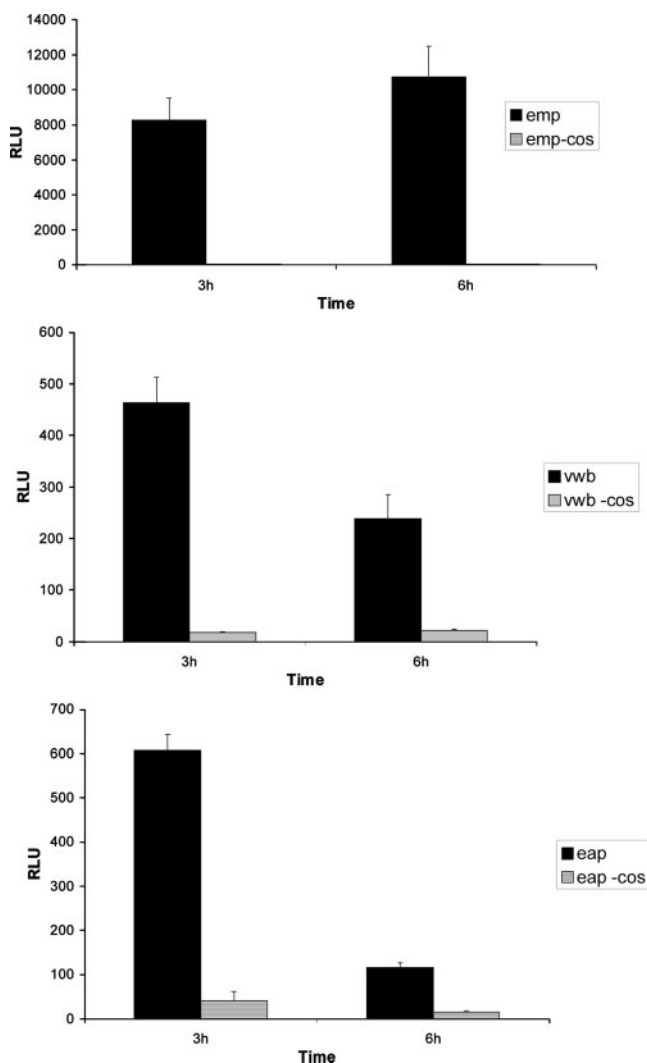


FIG. 2. Effect of deleting the COS in the *emp* (A), *vwb* (B), and *eap* (C) promoters. A β -galactosidase assay was used for measuring *emp* and *vwb* promoter activity, and the bioluminescence assay described in Harraghy et al. (19) was used for measuring *eap* expression. The data shown are the means \pm standard errors of the means of the results of at least two independent experiments. RLU, relative light units.

TTCGGACTC-3') and *vwb*PR1 (5'-CCTAAGCTTTAATTTT CCCTAATTAAC-3') amplified the entire promoter region, while primers *vwb*-cs_F (5'-CTACCTTTTTAAAATGTTGAT GAA-3') and *vwb*-cs_R (5'-ATTCATCAACATTTTAAAAA GG TAG-3') were the complementary internal primers lacking the COS. The COS in the *eap* promoter was deleted by using a QuikChange mutagenesis kit (Stratagene) using primers QCF1 (5'-GATAATTTATTATTAATATTCCAAAAATA GAGAAAGTCTGGC-3') and QCR1 (5'-GCCAGACTTTCT CTATTTTTTGGAATATTAATAATAAATTATC-3'). All clones were sequenced to confirm the deletion of the COS and that no additional mutations had been introduced during cloning. The mutated promoters were cloned in their respective reporter gene vectors and transduced into strain Newman as described in reference 19. The expression of *eap* was analyzed by using a bioluminescence assay, while *emp* and *vwb* were

analyzed by using a β -galactosidase assay. As shown in Fig. 2, deleting the COS in all three promoters severely repressed the expression of the reporter gene. To exclude the possibility that the deletion of the COS per se was responsible for the decrease in expression, we mutated the COS in the *emp* promoter, changing the sequence from AGTTAATT to TCATAATT (thereby changing the first three nucleotides of the COS while leaving the putative -35 box intact) by using a QuikChange mutagenesis kit (Stratagene) with primers QCF3_ *emp* (5'-GA CAACGTTTACTTTCATCATAATTATTATACTGAAAATT CTGG-3') and QCF3_ *emp*-r (5'-CCAGAATTTTCAGTATA ATAATTATGATGAAGTAAACGTTGTC-3'). As shown in Fig. 3, mutagenesis of the COS in the *emp* promoter resulted in a >50% decrease in *emp* expression but did not completely abrogate expression. This is likely due to the partial homology of the region to the COS. Taken together, our findings suggested that the COS could be the binding site for a regulator of *eap*, *emp*, and *vwb*.

In our previous study, we showed that *sarA* and RNAlII are involved in the regulation of *eap* and *emp* and that *sae* is essential for the expression of both genes (19). Six of the 11 genes in Table 1 are also regulated by *sae* (39). To investigate whether SaeR was binding to the *eap*, *emp*, and *vwb* promoters, SaeR was amplified from *S. aureus* Newman using primers *sae*R_F2 (5'-GGCATAACATATGACCCACTTACTGATC-3') and *sae*R_R3 (5'-CCCCAAGCTTATCGGCTCCTTTCAA ATTTATATCC-3'), cloned in the pET28a vector (Novagen), and overexpressed in *Escherichia coli*. The purified protein was subsequently assessed for binding to each promoter (see the supplemental material for the DNA sequences used) by using electrophoretic mobility shift assays (EMSA). However, no binding of SaeR to the promoters was found (data not shown). As it is possible that SaeR needs to be phosphorylated to bind to its target promoters (13), we decided to purify the DNA binding domain of SaeR and looked for binding of this to the promoters using EMSA. However, we did not observe any binding of the SaeR DNA binding domain to the three promoters. These findings, as well as the observations that *vwb* is not regulated by *sae* (39) and that some *sae*-regulated genes, e.g., *scn* and *chp* (40), do not have a COS in their promoter, suggest that the COS is not the binding site for SaeR.

The COS is similar to a proposed binding site for SarA (AGTTAAG) (38). As SarA is known to be involved in the regulation of *eap* and *emp*, and SarA binding to different promoters has been demonstrated (6, 38), we investigated whether the COS could be a binding site for SarA. Although SarA binds to the *eap*, *emp*, and *vwb* promoters (N. Harraghy and J. Kormanec, unpublished data), deleting the COS did not have any effect on SarA binding to the three promoters (data not shown), indicating that the COS is not essential for SarA binding.

In summary, we have identified a COS in the *eap*, *emp*, and *vwb* promoters, as well as in the promoters of several genes recognized as being involved in modulation of the immune response to staphylococcal infection. The nature of the relationship between the SERAMs (5) and leukocidins is intriguing as it was recently shown that, in some strains, the expression of the Pantone-Valentine leukocidin interferes with the regulation of the other major group of staphylococcal adhesins, the MSCRAMMs (27). Although it is unlikely that the leuko-

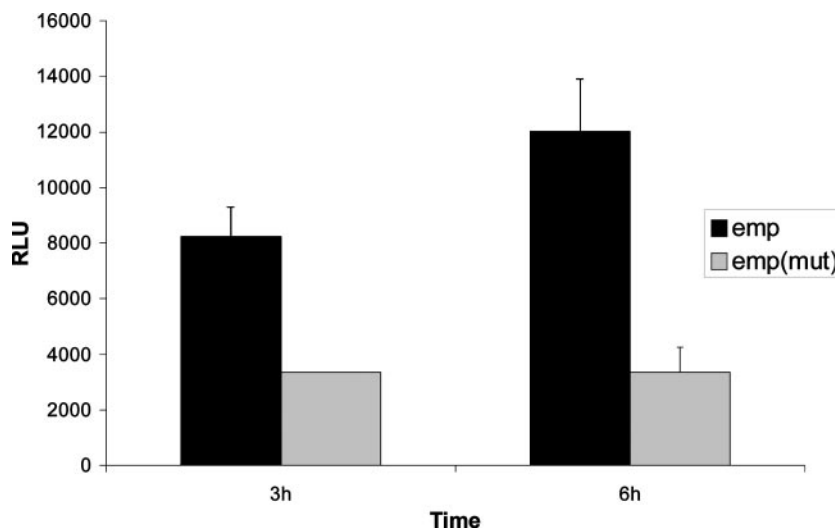


FIG. 3. Effect of mutating the COS in the *emp* promoter [*emp*(mut)]. The COS was changed from AGTTAATT to TCATAATT, and the effect on *emp* expression was assayed by using a β -galactosidase assay as described in reference 19. The data shown are the means \pm standard errors of the means of the results of two independent experiments. RLU, relative light units.

cidins described here affect the regulation of the SERAMs, it is possible that they share a common regulator.

Our findings suggest that the COS has an important functional role because deletion of the COS in the *eap*, *emp*, and *vwb* promoters, as well as mutation of the COS in the *emp* promoter, affected the expression of the reporter gene. Although deleting the COS only partially disrupted the proposed -35 box (in the case of the *eap* promoter, there is only one mismatch in comparison with the original promoter) the deletion dramatically affected promoter activity. Moreover, the mutation of the COS in the *emp* promoter, which preserved the -35 box and maintained homology to the COS (only three bases were different), also affected promoter activity, although not to the same extent as when the COS was deleted. Thus, the changes in the expression of the reporter genes appear to be the result of modifications to the COS and the possible loss of a transcription factor-binding site. Our findings suggest that the COS is the binding site for an as-yet-unidentified regulator of *eap*, *emp*, and *vwb* that may function together with SaeR. The existence of such a factor was postulated by Goerke et al. (14) and is supported by work in our laboratory, as well as the recent findings of Kuroda et al. (26). Emerging data from microarray studies and ongoing work in our laboratory will help reveal such candidates.

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