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Role of Adaptor TrfA and ClpPC in Controlling Levels of SsrA-Tagged Proteins and Antitoxins in *Staphylococcus aureus*

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***Staphylococcus aureus* responds to changing extracellular environments in part by adjusting its proteome through alterations of transcriptional priorities and selective degradation of the preexisting pool of proteins. In *Bacillus subtilis*, the proteolytic adaptor protein MecA has been shown to play a role in assisting with the proteolytic degradation of proteins involved in competence and the oxidative stress response. However, the targets of TrfA, the MecA homolog in *S. aureus*, have not been well characterized. In this work, we investigated how TrfA assists chaperones and proteases to regulate the proteolysis of several classes of proteins in *S. aureus*. By fusing the last 3 amino acids of the SsrA degradation tag to Venus, a rapidly folding yellow fluorescent protein, we obtained both fluorescence-based and Western blot assay-based evidence that TrfA and ClpCP are the adaptor and protease, respectively, responsible for the degradation of the SsrA-tagged protein in *S. aureus*. Notably, the impact of TrfA on degradation was most prominent during late log phase and early stationary phase, due in part to a combination of transcriptional regulation and proteolytic degradation of TrfA by ClpCP. We also characterized the temporal transcriptional regulation governing TrfA activity, wherein Spx, a redox-sensitive transcriptional regulator degraded by ClpXP, activates *trfA* transcription while repressing its own promoter. Finally, the scope of TrfA-mediated proteolysis was expanded by identifying TrfA as the adaptor that works with ClpCP to degrade antitoxins in *S. aureus*. Together, these results indicate that the adaptor TrfA adds temporal nuance to protein degradation by ClpCP in *S. aureus*.**

The Gram-positive human pathogen *Staphylococcus aureus* causes invasive infections of the skin and bloodstream that can lead to life-threatening sepsis, endocarditis, and toxin-mediated syndrome (1). Unfortunately, transmission and infection are commonplace in hospitals and are increasing in frequency in community settings (2). This scenario is in part due to the ability of *S. aureus* to survive and spread from abiotic surfaces, such as gloves (3) and clothing (4), as well as from biotic niches, like the anterior nares of healthy adults (5).

The transcriptional regulation of numerous stress response mechanisms that facilitate its survival is well described (6, 7). However, posttranslational modifications of the proteome in *S. aureus* are poorly characterized in comparison, even though they play a significant role in the stress response and general house-keeping in many prokaryotes (8). These adjustments include recovery of denatured or aggregated proteins via GroEL- and ClpB-mediated refolding (9), tuning of the temporal and spatial activities of downstream regulatory networks (10), offsetting of starvation by increasing the available pools of amino acids (11), and hedging against unregulated cell death via toxin-antitoxin system activation (12, 13).

In *S. aureus*, proteolytic degradation is carried out by the intracellular proteases ClpP, FtsH, and ClpQ (also known as HslV). Of the three, ClpP appears to have the greatest impact in shaping the *S. aureus* proteome (14), and when disrupted, it causes significant defects in growth, survival, virulence gene expression, and cell wall maintenance (15, 16). FtsH has a modest role in regulating growth, stress resistance, starvation survival, and pathogenicity (17), but with a putative localization to the inner membrane, it is thought to mainly degrade membrane-bound targets. Very little is known about ClpQ, in part because a $\Delta clpQ$ mutant of *S. aureus* exhibits few phenotypes (18) and shows few alterations to stress responses.

The manner in which a protein is specifically degraded by one of these proteases is initiated by the identification of a degradation motif during partial ATP-dependent unwinding of the target (19). FtsH contains its own intrinsic unwinding domain to facilitate this process, but ClpP and ClpQ must associate with a member of the AAA+ ATPase family of chaperones (19) to accomplish this. This association with ClpP or ClpQ allows chaperones to unwind and deliver identified targets into the cognate protease's catalytic chamber.

S. aureus has three known chaperones that aid in ClpP- and ClpQ-mediated proteolysis, while a related fourth chaperone, ClpL, lacks the structures necessary to interact with ClpP or ClpQ (20). The chaperone ClpY (also known as HslU) assists ClpQ in recognition of target motifs, and strains lacking *clpY* have phenotypes similar to those of strains lacking *clpQ* (18), suggesting that few other partners exist for this chaperone/protease pair. The ClpC and ClpX chaperones associate with ClpP, with each selecting different groups of proteins for ClpP-mediated proteolysis. ClpX is primarily involved in virulence regulation (16), while ClpC tunes how the cell regulates its oxidative stress response (21), tricarboxylic acid cycle (21), toxin-antitoxin system activity (22), and survival in resource-limited medium (23).

Members of a second class of proteins called adaptors assist chaperones in controlling the proteolytic degradation of target

Received 14 August 2014 Accepted 9 September 2014

Published ahead of print 15 September 2014

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doi:10.1128/JB.02222-14

substrates (8). These proteins lack the AAA+ ATPase activity found in chaperones and instead facilitate motif identification either through direct interactions with a target (24), through selective motif occlusion (25), or by modification of a protein to alter its degradation rate (26). On the basis of homology to adaptors identified in *Bacillus subtilis*, there are at least three adaptors in *S. aureus*, namely, McsB (SA0482), YjbH (SA0860), and TrfA (SA0857). Studies have indicated the involvement of each in various stress responses (27–29), but many gaps remain in our understanding of the various roles played by these adaptors in *S. aureus*. Details of the protein targets of these *S. aureus* adaptors are limited, and to date, only two target proteins have been described to be regulated by *S. aureus* adaptors. McsB targets CtsR for ClpCP degradation (30), and the levels of the redox-sensing transcriptional regulator Spx have been shown to be controlled by the adaptor YjbH (31).

In this work, we investigated how the chaperones and adaptors of *S. aureus* assist in the proteolysis of several classes of proteins. More specifically, we found that the ClpCP proteolytic system and not the ClpXP system described in *Escherichia coli* (32) and *B. subtilis* (33) is essential for SsrA-tagged protein destruction in *S. aureus*. The adaptor TrfA was discovered to mediate this breakdown, primarily during late-logarithmic- and early-stationary-phase growth. This regulation is likely a result of the transcriptional regulation as well as the growth phase-dependent proteolytic degradation of TrfA by ClpCP. Additionally, a regulatory feedback loop was uncovered for a positive regulator of *trfA*, whereby Spx represses one of its promoters, which in turn affects *trfA* transcription. TrfA was also identified as the adaptor involved in the ClpCP-mediated degradation of antitoxins, revealing a role for the modulation of toxin activity in *S. aureus* (34). Together, these results reveal how the adaptor TrfA temporally tunes the ClpCP-mediated protein degradation of several classes of proteins involved in *S. aureus* physiology.

MATERIALS AND METHODS

Bacterial strains and culture conditions. Table 1 contains a list of the bacterial strains used in these studies. *E. coli* strains were routinely grown in LB, and *S. aureus* strains were routinely grown in Trypticase soy broth (TSB). Both strains were grown with shaking at 250 rpm at 37°C. Cell density was measured by determination of the absorbance (optical density [OD]) at 650 nm using an 18-mm borosilicate glass tube in a Spectronic 20D+ spectrophotometer. Ampicillin (Amp; 50 µg/ml), chloramphenicol (Cm; 10 µg/ml), or erythromycin (Erm; 2.5 µg/ml) was added to the appropriate medium when required by a particular strain. To disrupt transcription or protein synthesis, rifampin (200 µg/ml) or erythromycin (50 µg/ml) was added to the cultures, respectively. Promoter induction of pEPSA5 was performed using 0.1 to 0.5% xylose, when necessary. Thiol stress was induced by the addition of 5 mM diamide [1,1-azo-bis(*N,N*-dimethylformamide)]; Sigma-Aldrich].

DNA manipulations. *E. coli* plasmid purification was performed using Qiagen miniprep kits (Qiagen) per the manufacturer's instructions, while plasmid isolation from *S. aureus* was performed as described previously (35). Plasmid transformations in *S. aureus* were achieved via electroporation with a MicroPulser apparatus (Bio-Rad), using RN4220 as an intermediate between *E. coli* and relevant *S. aureus* strains.

Codon optimization of *venus*. To create a construct that yields a rapidly maturing fluorescent protein, the Venus gene was codon optimized for *S. aureus* using the codon frequencies calculated from the *S. aureus* COL genome, while avoiding the generation of additional restriction enzyme sites. Subsequently, 42-bp primers staggered to each neighboring sequence by 21 bp were combined in assembly reactions using HiFi PCR

Supermix (Invitrogen) to create ~300-bp fragments of *venus*, followed by the addition of flanking primers at the 5' and 3' ends of each 300-bp fragment using a new PCR. Following gel purification, equimolar amounts of each fragment were combined for a second assembly reaction with primers at the 5' and 3' ends of *venus*, and the PCR product was purified and sequenced. Both the primers and *S. aureus* codon-optimized *venus* sequences are available upon request.

Plasmid construction. To ensure the robust expression of our codon-optimized *venus* gene, we first cloned a strong phage repressor promoter of *saOUHSC_02234* from a lysogenized phage (36) into the EcoRI-XbaI restriction sites of shuttle plasmid pSK236. To place *venus* under the control of the phage repressor promoter, we cloned *venus* together with an upstream *sarA* ribosomal binding site into the SalI-PstI sites of pSK236. To add the *S. aureus* SsrA tag GKSNNNFVAA (25) to the C terminus of Venus (in which the construct was designated *venus*-SsrA), the following primers were used: 5'-AACTGCAGTTATGCTGCAACCTTGTAAGTT CATCCATTCC-3' and 5'-AACTGCAGTTAGGCAGCTACTGCGAAAT TATTGTTTGATTGCCAGTCTGTAAAGTTCATCCATTCC-3'. This sequence diverges from that of Flynn et al. (25) by its lack of an alanine at its N terminus, as this residue was not found in genomic sequences of clinical and laboratory isolates (data not shown). To add the VAA tag to the C terminus of Venus (in which the construct was designated *venus*-VAA), the following primer was used in place of the second primer listed above: 5'-ACGCGTCG ACGGAGGTTTAAACATGGTTTCTAAAGGTGAA-3'. To construct a separate recombinant plasmid for the induction of Venus expression, we cloned *venus* and the *venus*-VAA variant into the EcoRI and BamHI sites of pEPSA5 containing a xylose-inducible promoter (37).

For plasmid constructs capable of expressing *trfA* or *spx* under induction, we used a cloning strategy in pEPSA5 similar to the one used for *venus*. Briefly, *trfA* or *spx* was amplified by PCR with a 5' codon-optimized *myc* tag, and the amplified product was cloned into the EcoRI and BamHI sites of pEPSA5. In addition, *S. aureus* SH1000 strains capable of overexpressing *myc*-tagged antitoxin genes from *S. aureus* under the control of the xylose promoter of pEPSA5 were constructed as previously described (22).

To generate an *spx* promoter reporter construct, we cloned the distal P1 promoter or proximal P2 promoter region of *spx* (starting 245 bp or 102 bp before the *spx* start codon, respectively; amplification primers are available on request) into the EcoRI and XbaI sites of pALC1484, a shuttle plasmid containing a green fluorescent protein (GFP) reporter gene. Following sequencing verification, the plasmids were used to transform RN4220 and subsequent strains.

Construction of deletion strains. All sequenced *S. aureus* genomes contain the following three adaptor genes: *mcsB* (*sa0482*), *trfA* (*sa0857*), and *yjbH* (*sa0860*). As our relevant *clp* deletion strains are constructed in the *S. aureus* SH1000 background, we proceeded to construct deletion mutants of these adaptor genes in strain SH1000. Deletion mutants were constructed using the temperature-sensitive allelic replacement plasmid pMAD (38). In brief, chromosomal regions 1,000 bp up- and downstream of the gene of interest were amplified by PCR with primers proximal to the deletion site, joined by gene sewing, ligated into pMAD, and transformed into RN4220 and then into SH1000. The mutant was then generated by a temperature shift strategy to yield the mutant, as described previously (38). To generate a pMAD construct to complement a *trfA* deletion mutant, we inserted a DNA fragment with the intact gene flanked by 1,000 bp on each side into pMAD, followed by allelic replacement in the mutant. All constructs were verified by colony PCR and chromosomal sequencing.

Isolation of RNA and Northern blot hybridization. Overnight cultures of *S. aureus* were each diluted 1:1,000 and grown at 37°C with shaking at 250 rpm in 100 ml of TSB inside 500-ml sidearm flasks (flask/medium ratio, 5:1). Cell density was monitored by determination of the absorbance at 650 nm, and at the appropriate OD, aliquots of cells were pelleted and flash frozen at –80°C. RNA extraction was accomplished by lysing the cells with 1 ml of cold TRIzol reagent (Invitrogen) and 0.25 ml

TABLE 1 Strains and plasmids used in this study

Strain or plasmid	Genotype or characteristics	Source or reference
Strains		
<i>S. aureus</i>		
RN4220	Heavily mutagenized derivative of NCTC 8325-4	63
SH1000	NCTC 8325-4 with repaired <i>rsbU</i> gene	64
JM27	SH1000 <i>ftsH::tetL</i>	17
NCTC 8325-4 Δ <i>spx</i>	NCTC 8325-4 Δ <i>spx</i>	60
NCTC 8325-4 <i>spx</i> complement	NCTC 8325-4 <i>spx</i> mutant complemented with <i>spx</i>	60
ALC4976	SH1000 Δ <i>clpB</i>	22
ALC4977	SH1000 Δ <i>clpC</i>	22
ALC4978	SH1000 Δ <i>clpL</i>	22
ALC4979	SH1000 Δ <i>clpQ</i>	22
ALC4980	SH1000 Δ <i>clpX</i>	22
ALC4981	SH1000 Δ <i>clpY</i>	22
ALC5105	SH1000 Δ <i>clpP</i>	22
ALC6490	SH1000 <i>clpC</i> mutant complemented with <i>clpC</i>	22
ALC6491	SH1000 <i>clpP</i> mutant complemented with <i>clpP</i>	22
ALC7010	SH1000 Δ <i>mcsB</i>	This work
ALC7011	SH1000 Δ <i>trfA</i>	This work
ALC7137	SH1000 Δ <i>yjbH</i>	This work
ALC7272	SH1000 <i>trfA</i> mutant complemented with <i>trfA</i>	This work
<i>E. coli</i> XL1-Blue	General cloning strain	Agilent
Plasmids		
pMAD	<i>E. coli/S. aureus</i> shuttle plasmid with the <i>ori</i> pE194 ^{ts} ; <i>bgaB</i> Amp ^r Erm ^r	38
pEPSA5	Xylose-inducible shuttle vector; Amp ^r Cm ^r	37
pSK236	<i>S. aureus/E. coli</i> shuttle vector with pUC19 cloned into the HindIII site of pC194	65
pALC1484	pSK236 with promoterless <i>gfp-uvr</i> , a cycle 3 <i>gfp</i> allele	66
pALC6188	pEPSA5 with <i>mazE</i> with a 3' <i>myc</i> tag and <i>sarA</i> ribosome binding site	22
pALC6486	pEPSA5 with <i>axe1</i> with a 5' <i>myc</i> tag and <i>sarA</i> ribosome binding site	22
pALC6489	pEPSA5 with <i>axe2</i> with a 5' <i>myc</i> tag and <i>sarA</i> ribosome binding site	22
pALC6954	pSK236 with a phage repressor promoter and codon-optimized <i>venus</i>	This work
pALC6955	pSK236 with a phage repressor promoter and codon-optimized <i>venus-VAA</i>	This work
pALC7016	pEPSA5 with codon-optimized <i>venus-VAA</i> and <i>sarA</i> ribosome binding site	This work
pALC7113	pEPSA5 with <i>spx</i> with a 5' <i>myc</i> tag and <i>sarA</i> ribosome binding site	This work
pALC7151	pEPSA5 with <i>trfA</i> with a 5' <i>myc</i> tag and <i>sarA</i> ribosome binding site	This work
pALC7257	pMAD:: <i>trfA</i> (<i>sa0857</i>) complement	This work
pALC7422	pSK236 with a phage repressor promoter and <i>trfA</i> with a 5' <i>myc</i> tag	This work
pALC7867	pMAD:: Δ <i>trfA</i> (<i>sa0857</i>)	This work
pALC7868	pMAD:: Δ <i>mcsB</i> (<i>sa0482</i>)	This work
pALC7869	pMAD:: Δ <i>yjbH</i> (<i>sa0860</i>)	This work
pALC7803	pALC1484 with the <i>spx</i> P1 promoter in EcoRI and XbaI sites	This work
pALC7804	pALC1484 with the <i>spx</i> P2 promoter in EcoRI and XbaI sites	This work
pALC7901	pSK236 with a phage repressor promoter and codon-optimized <i>venus-SsrA</i>	This work

of 0.1-mm-diameter silica/glass beads in a Mini-BeadBeater 8 apparatus (Biospec Products) at its maximum setting for two 1-min pulses. RNA purification and subsequent Northern analysis were performed as described previously (39).

Cell lysate preparation and Western blot analyses. Protein lysates were prepared as described before (22) from cells grown in a fashion similar to that for RNA extraction described above. At an OD of 650 nm (OD₆₅₀) of 1.1, growth was stalled with antibiotics (either 200 μ g/ml rifampin or 50 μ g/ml erythromycin, depending on the experiment), and a series of cells (10 ml each at a 15- or 30-min interval) were pelleted at 1°C, flash frozen in liquid nitrogen, and stored at -80°C. For lysis, the pellets were thawed on wet ice, washed twice with ice-cold sample buffer (50 mM Tris-HCl, pH 7.6, 5 mM EDTA), and resuspended in 200 μ l of sample buffer to which 100 μ l of 0.1-mm-diameter silica/glass beads was added. The cells were then disrupted in a Mini-BeadBeater 8 apparatus at its maximum setting for two 1-min pulses with a 1-min rest, followed by centrifugation at 20,000 \times g for 15 min at 1°C. Halt protease inhibitors

(1 \times ; Thermo Fisher) were then added to the lysates, and their protein concentrations were determined by Bradford assay using an FL600 microplate reader (BioTek Instruments).

For Western blot assays, SDS-PAGE was first performed with 30 μ g of lysate proteins, and then the proteins were transferred to polyvinylidene difluoride (PVDF) membranes using an iBlot Western blot transfer system (Invitrogen). Following blocking in 5% milk in Tris-buffered saline with Tween (TBS-T; 150 mM NaCl and 10 mM Tris, pH 8.0, with 0.1% [vol/vol] Tween 20) for at least 30 min, the membranes were then incubated with either mouse anti-GFP (Abcam) or mouse anti-Myc (Cell Signaling) antibodies in TBS-T for 2 h. The membranes were then washed in TBS-T and incubated with horseradish peroxidase-conjugated goat anti-mouse secondary antibody (Jackson ImmunoResearch). After additional washing, enhanced chemiluminescence (ECL) reagent (GE Healthcare) was added and the membrane was exposed to film. Scanned images from nonsaturated exposed film were then analyzed densitometrically using ImageJ software, and the half-lives of the

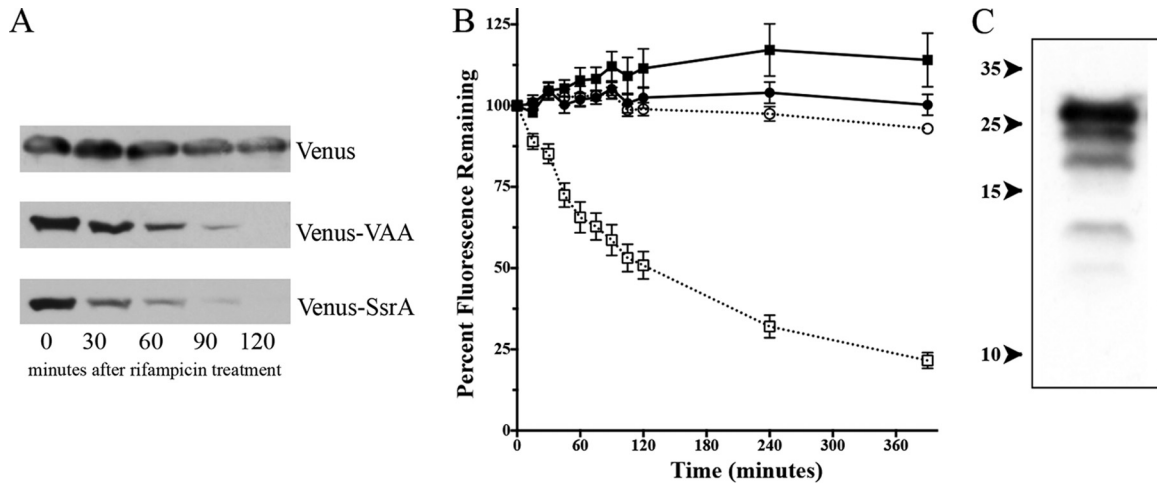


FIG 1 Venus and Venus-VAA protein levels in *S. aureus* SH1000 wild-type cells following rifampin treatment. (A) Western blot analysis of lysates of wild-type SH1000 with either pALC6954 (producing Venus), pALC6955 (producing Venus-VAA), or pALC7901 (producing Venus-SsrA) grown to an OD_{650} of 1.1 after rifampin treatment (200 $\mu\text{g}/\text{ml}$). The blot was probed with an anti-GFP primary antibody and then a secondary horseradish peroxidase-conjugated goat anti-mouse antibody. The experiments were repeated three times, and a representative blot of each set of lysates is displayed. (B) Venus fluorescence emitted from strains ALC6954 (producing Venus; circles) and ALC6955 (producing Venus-VAA; squares) either stalled by 50 $\mu\text{g}/\text{ml}$ erythromycin after reaching an OD_{650} of 1.1 (open symbols) or not treated with drug (closed symbols). The fluorescence at 540 nm was observed following excitation at 515 nm and normalized to the cell density at the OD_{650} . Values were then compared to the fluorescence at time zero. The result for each time point is the mean and standard deviation of three biological replicates. (C) Degradative laddering of Venus-VAA. The lysate from wild-type SH1000 expressing Venus-VAA from pALC6955 was resolved and immunoblotted with an anti-GFP primary antibody and a secondary horseradish peroxidase-conjugated goat anti-mouse antibody. Exposure to film was extended to reveal lower-molecular-mass bands. Numbers on the left are molecular masses (in kilodaltons).

proteins of interest were calculated using Prism (version 5) software (GraphPad).

Fluorescence analysis. To analyze the fluorescence of *S. aureus* cells expressing Venus or its variants, cells were monitored using one of two machines. For single time points, results for 200 μl of cells at various time points were read in a 96-well plate (Costar plate 3632) in an FL600 plate reader (BioTek) with 515-nm excitation and 540-nm emission filters (Omega Optical). In experiments with multiple time points, 150 μl of cells was shaken in a 96-well plate (Costar plate 3632) with a heated top plate to prevent condensation at 250 rpm and 37°C in a Tecan Infinite M1000 Pro apparatus (Tecan). Fluorescence was recorded using 515-nm excitation and 528-nm emission filters and 650-nm absorbance filters. In both cases, data are reported as the level of fluorescence per absorbance unit to normalize the cell density.

RESULTS

Establishing an *in vivo* assay for SsrA-tagged proteins in *S. aureus* using Venus fluorescence. During translation, ribosomes may stall for a variety of reasons, including stress, starvation, or premature degradation of the template mRNA. To reset the translational machinery, a hybrid RNA called transfer-messenger RNA (tmRNA) enters the open A site of the ribosome, shifting translation from the degraded or truncated mRNA to a small open reading frame with a stop codon within the tmRNA (40). The resulting 11-amino-acid tag, called the SsrA tag, is appended to the C terminus of the nascent but truncated protein and is then recognized as a signal for rapid degradation by ClpAP and ClpXP in *E. coli* (32) or ClpXP in *B. subtilis* (33). As the half-lives of SsrA-tagged proteins are short, reportedly <10 min (41), the tracking of such proteins is difficult *in vivo*. To investigate this system in *S. aureus* and to obtain an improved sensitivity compared to that obtained with the slowly folding non-codon-optimized cycle 3 *gfp* (42), we utilized a fast-folding yellow fluorescent protein (YFP)-tagged Venus variant (SEYFP-F46L) (43) which exhibits an accelerated ox-

idation step as a result of the F46L mutation, leading to rapid and enhanced fluorescence (43). In *S. aureus*, the SsrA tag has been predicted to be AGKSNNNFAVAA (25), but it has never been tested. Given that the N-terminal alanine of the predicted *S. aureus* sequence is not conserved in sequenced genomes (data not shown), we chose instead to append the sequence GKSNNNFAVAA to the C terminus of Venus. Furthermore, identifying the minimal unit of SsrA necessary for degradation was also of interest. Based on prior work in other organisms (44) suggesting that the final 3 amino acids are sufficient as a degradation signal, we created an additional construct appending the amino acids VAA to the C terminus of Venus.

Following introduction of the recombinant plasmids into *S. aureus* strain SH1000, initial work revealed significant fluorescence, while the cells exhibited no retardation of growth in the presence of any of these plasmids (data not shown). To verify that the SsrA tag and the abbreviated VAA tag on Venus were produced and functional, cells with the *venus* construct (pALC6954), the *venus-VAA* construct (pALC6955), or the *venus-SsrA* construct (pALC7901) were grown in TSB as described in Materials and Methods until the OD_{650} reached 1.1, whereupon 200 $\mu\text{g}/\text{ml}$ of rifampin was added to the culture and samples were withdrawn at predetermined time points for 2 h. Western blot analysis of whole-cell lysates probed with a mouse anti-GFP antibody capable of detecting Venus showed that the Venus protein levels rapidly decreased in both the strain containing Venus-VAA (pALC6955) and the strain containing Venus-SsrA (pALC7901), but no equivalent decrease was observed in cells producing untagged Venus (pALC6954) (Fig. 1A). This finding implied that Venus is stable in *S. aureus* and that addition of SsrA, notably, just the last 3 amino acids of such a tag (VAA), was sufficient to destabilize it. Densitometric analysis indicated that the expressed Venus-VAA construct

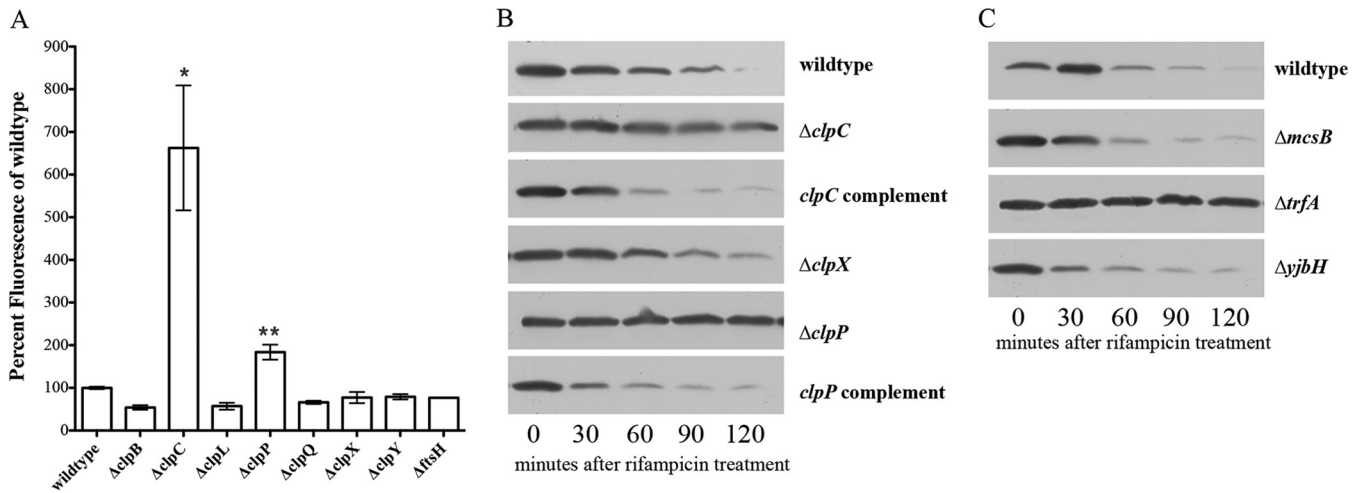


FIG 2 Venus levels in *clp* and adaptor mutant strains of SH1000. (A) Venus fluorescence analysis of overnight cultures of *S. aureus* strains containing pALC6955 (producing Venus-VAA). Venus fluorescence values (excitation and emission, 515 nm and 540 nm, respectively) were normalized to the OD₆₅₀ values and then reported as a percentage of the wild-type levels. Each value is the mean and standard deviation of three biological replicates. *, statistically significant differences ($P < 0.005$) compared to the wild type measured by Student's unpaired *t* test. (B) Western blot of Venus-VAA protein levels in *S. aureus* SH1000 wild-type and Δclp cells containing pALC7016 with a xylose-inducible promoter (producing Venus-VAA) after transcriptional arrest by rifampin (200 μ g/ml). Strains were grown to an OD₆₅₀ of 1.1 and then treated with rifampin (200 μ g/ml), and samples were taken every 30 min. Whole-cell lysates of these samples were then separated by SDS-PAGE on a 20% gel and transferred to PVDF for immunoblotting with an anti-GFP primary antibody and subsequently with a horseradish peroxidase-conjugated goat anti-mouse antibody. The experiments were repeated three times, and representative blots are displayed. (C) Western blot of Venus-VAA protein levels in *S. aureus* SH1000 wild-type and adaptor mutant strains containing pALC7016 (producing Venus-VAA). Whole-cell lysates of 0.5% xylose-induced samples obtained at different time points after rifampin treatment were resolved, blotted, and probed as described above for panel B. The experiments were repeated three times, and representative blots are displayed.

had a half-life of 47 min, while the Venus-SsrA construct had a half-life of 27 min, suggesting that while VAA is sufficient for degradation, other residues of the SsrA tag may play a role in proteolytic degradation.

To pursue our observation that the VAA residues of SsrA were sufficient for proteolytic degradation, we examined whether there was a functional correlation of protein degradation to fluorescence loss. To accomplish this, we used erythromycin (50 μ g/ml) at an OD₆₅₀ of 1.1 to stop protein synthesis (rifampin interferes with the fluorescence spectrum), followed by monitoring of cell density and fluorescence over 6 h. As seen in Fig. 1B, Venus and Venus-VAA levels remained largely unchanged over time in the absence of erythromycin. However, when erythromycin was added, the fluorescence levels of Venus-VAA (ALC6955) began to decrease with a half-life of 75 min, whereas the levels in ALC6954 (Venus) remained unchanged. This result suggested that the fluorescence associated with the degradation of Venus with the SsrA tag was reduced in the cell, albeit more slowly than the reduction of the protein (half-life, 47 min), as determined by Western blotting (Fig. 1A). The reason for this discrepancy is unknown but may be due to either Venus-VAA molecules retaining the capacity for emitting fluorescence even when partially degraded or the formation of *in vivo* foci that affect overall fluorescence (45). In support of the former possibility, we observed a laddering of lower-molecular-weight bands in Western blots of the Venus-VAA lanes following addition of rifampin (Fig. 1C), suggesting that proteolysis occurred in a stepwise fashion.

***clpC* instead of *clpXP* is involved in SsrA degradation in *S. aureus*.** With the availability of a reporter construct capable of displaying the relative amount of an SsrA- or VAA-tagged protein in a cell, we examined the role of ATP-dependent proteases and their cognate chaperones in the breakdown of SsrA-tagged protein

in various *clp* mutants in our collection using the Venus-VAA construct. As shown in Fig. 2A, Venus-VAA levels were higher in overnight cultures of $\Delta clpC$ and $\Delta clpP$ strains than in those of wild-type and other protease and chaperone mutant strains. Similar results were also observed in *clpC* and *clpP* mutants when grown for 6 h (data not shown). Interestingly, the amount of Venus-VAA that accumulated in the $\Delta clpP$ mutant was not as high as that which accumulated in the $\Delta clpC$ mutant. Given that $\Delta clpP$ strains exhibit lower growth rates, smaller colony sizes, and altered metabolic intracellular environments compared to those of the wild type (16), it is conceivable that these altered phenotypes may have accounted for the lower fluorescence in the *clpP* mutant than the *clpC* mutant. We next examined whether this difference in fluorescence between the $\Delta clpC$ and $\Delta clpP$ mutants was actually reflected in degradation of the Venus-VAA protein. Accordingly, we cloned the *venus-VAA* gene into pEPSA5, which allows Venus-VAA expression upon xylose induction, and then introduced this plasmid, pALC7016, into SH1000, $\Delta clpC$, and $\Delta clpP$ strains. Western blots of cell lysates taken from these xylose-induced cells following rifampin treatment were probed with anti-GFP antibody, revealing increased half-lives of Venus-VAA in the *clpC* and *clpP* mutants during 2 h of transcriptional stalling, while degradation was restored in the complemented *clpC* and *clpP* mutants (Fig. 2B). In contrast, a similar effect was not seen in the *clpX* mutant. Taken together, these data indicate that ClpC is the chaperone involved in SsrA-mediated degradation in *S. aureus*, contrary to the finding in *E. coli* (32) and *B. subtilis*, where ClpX is the primary chaperone that facilitates SsrA-mediation degradation (33).

TrfA is necessary to break down SsrA in *S. aureus*. To determine if other additional components were involved in degrading SsrA-tagged proteins in *S. aureus*, we examined the role(s) played

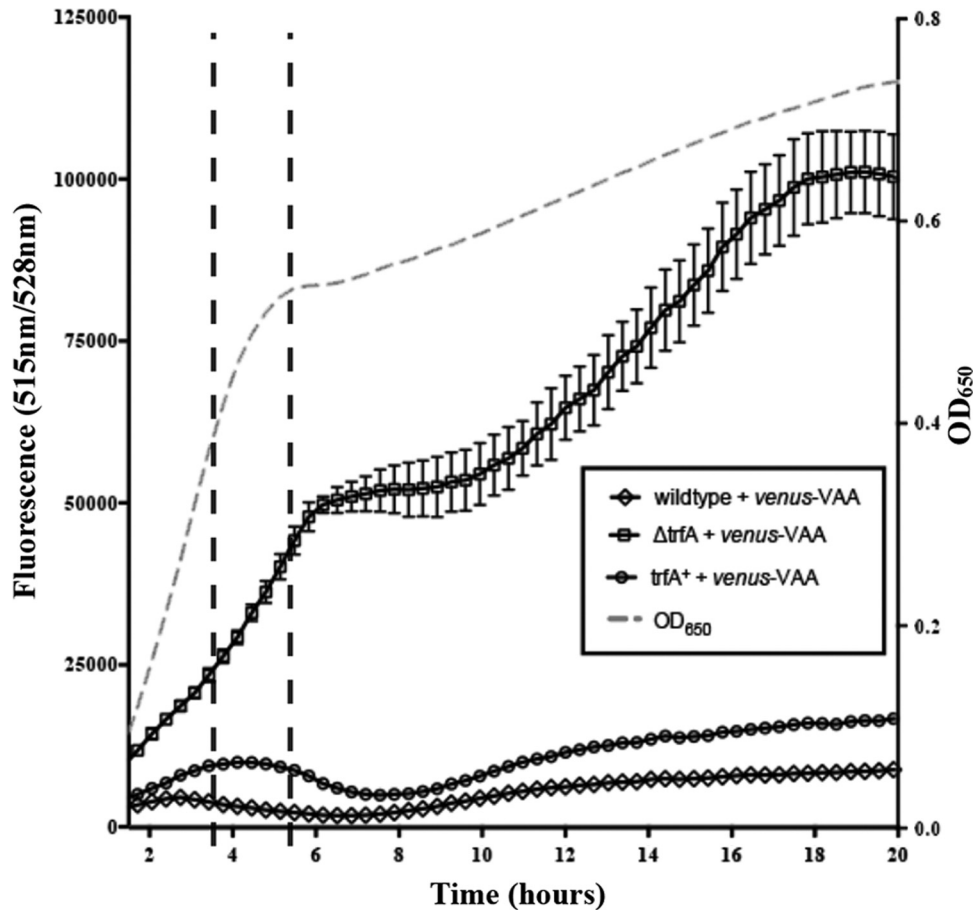


FIG 3 Venus-VAA levels in the wild type and the $\Delta trfA$ mutant over 20 h of growth. The raw Venus fluorescence of the *S. aureus* SH1000 wild-type, $\Delta trfA$, and $trfA$ -complemented ($trfA^+$) strains containing pALC6955 (producing Venus-VAA) over 20 h is shown. Cell densities at an OD_{650} were measured, while Venus fluorescence was detected at 540 nm (black lines), with each value representing the mean of three biological replicates.

by various adaptor proteins. The *S. aureus* genome contains genes for three known adaptors: *mcsB* (*sa0482*), *trfA* (*sa0857*), and *yjbH* (*sa0860*). We first transformed the pEPSA5-Venus-VAA construct pALC7016 into *mcsB*, *trfA*, and *yjbH* mutants. Western blot analysis of Venus-VAA breakdown in these adaptor mutants with pALC7016, grown to an OD_{650} of 1.1, after which rifampin was added, showed that only *trfA* was involved in degrading Venus-VAA (Fig. 2C). These data indicate that TrfA, together with ClpCP, controls SsrA-mediated degradation in *S. aureus* through the 3 terminal amino acids of the SsrA tag.

To examine how deletion of *trfA* impacted SsrA degradation at various phases of growth, we sampled fluorescence levels during 20 h of growth in TSB medium for wild-type, $\Delta trfA$, and $trfA$ -complemented strains expressing Venus-VAA (Fig. 3). Starting from overnight cultures, strains were back-diluted 1/100 and grown to an OD_{650} of 0.6, whereupon they were normalized to an OD_{650} of 0.1, and 150 μ l of this culture was aliquoted into a well of a 96-well plate. Cell growth was comparable among the three strains when incubated at 37°C with shaking for 20 h (data not shown). Analysis of the strains at these time points revealed that the $\Delta trfA$ strain (Fig. 3, squares) produced much higher levels of both raw and normalized fluorescence at all time points than the wild-type (Fig. 3, diamonds) and $trfA$ -complemented (Fig. 3, circles) strains. This suggests that Venus-VAA levels are generally

elevated in a $\Delta trfA$ strain relative to those in the wild-type and complemented strains.

The fluorescence intensity of Venus-VAA and its 47-min half-life make it possible to uncover conditional changes of fluorescence between strains. Specifically, the raw fluorescence in all strains (Fig. 3) increased at various rates in early logarithmic phase up through 3.5 h, but at late logarithmic phase (3.5 to 5.5 h), Venus-VAA levels began to decrease in the wild-type and $trfA$ -complemented strains but not a $\Delta trfA$ strain (Fig. 3). This decrease or flattening of both the raw and normalized fluorescence of strains containing *trfA* was maintained well into stationary phase, while the fluorescence of Venus-VAA continued to climb in the $\Delta trfA$ strain. This suggests that *trfA* has a lesser role in SsrA regulation in early and mid-logarithmic phase (when there was no decrease in fluorescence in the wild type) but gains in importance as cells begin to exit logarithmic phase and enter stationary phase.

Transcriptional regulation of *trfA*. *trfA* is transcribed from three promoters, generating transcripts of closely related sizes of 751, 818, and 898 bases, which can be seen as a single transcript in a 0.7% agarose gel (28). The changes to *trfA* transcription during various growth phases remain unclear, however. To investigate *trfA* transcript levels over time, we conducted Northern blot analyses using RNA taken from SH1000 cells grown to various time points. Our data demonstrate (Fig. 4A, unstimulated time points)

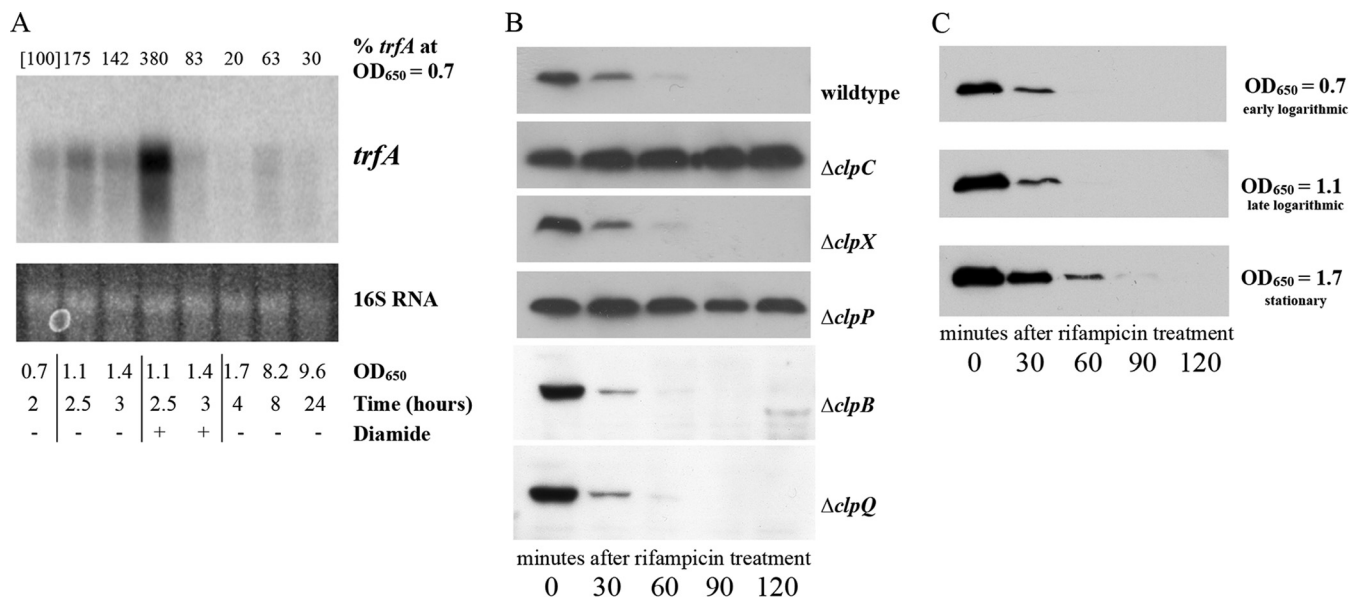


FIG 4 Transcriptional and posttranslational regulation of *trfA*. (A) Northern blot of *trfA* during growth and thiol stress. *S. aureus* SH1000 was grown to various ODs under conditions with and without 5 mM diamide for thiol stress. The RNAs obtained from these cells (20 μ g each) were resolved on a denaturing agarose gel, blotted to a Hybond XL membrane, and hybridized with a 300-bp ³²P-radiolabeled *trfA* DNA fragment. The 16S rRNA of the ethidium bromide-stained gel was used as the loading control for the blot. The percentages of the *trfA* transcript at a given time point relative to the amount of *trfA* transcript at an OD₆₅₀ of 0.7 (set equal to 100%, as indicated in brackets), determined by densitometric analysis, are reported at the top of the image. (B) Western blots of whole-cell lysates of various SH1000 strains transformed with pALC7151 (producing TrfA with an N-terminal *myc* tag). Cells were induced with 0.1% xylose at an OD₆₅₀ of 1.1 and then treated with rifampin (200 μ g/ml). Samples taken every 30 min were immunoblotted with a mouse anti-Myc primary antibody, followed by a horseradish peroxidase-conjugated goat anti-mouse antibody. The experiments were repeated three times, and representative blots are displayed. (C) Western blot analyses of TrfA in wild-type strain SH1000 at various time points following rifampin treatment. Wild-type strain SH1000 containing pEPSA5 expressing N-terminally Myc-tagged TrfA was induced with 0.1% xylose at early logarithmic phase (OD₆₅₀ = 0.7), late logarithmic phase (OD₆₅₀ = 1.1), or stationary phase (OD₆₅₀ = 1.7) and then treated with rifampin (200 μ g/ml). Samples were then immunoblotted with an anti-Myc primary antibody, followed by a horseradish peroxidase-conjugated goat anti-mouse antibody. The experiments were repeated three times, and representative blots are displayed.

that the *trfA* transcripts in SH1000 increased from early log phase and peaked at late log phase (OD₆₅₀, 1.1), followed by a significant decrease in stationary phase to a point where transcription was all but halted during deep-stationary-phase culture (at 24 h).

trfA has also been shown to be responsive to certain stresses (28). Reverse transcription-PCR (RT-PCR) analysis showed that one or more of the *trfA* transcripts were elevated following antibiotic or thiol stress. To evaluate the resiliency and duration of this *trfA* induction, we examined *trfA* levels following the addition of 5 mM diamide. *S. aureus* cells were grown to an OD₆₅₀ of 0.7 and then split for continued growth or challenge with 5 mM diamide. Samples were withdrawn 30 and 60 min later (corresponding to OD₆₅₀s of 1.1 and 1.4, respectively). As seen in Fig. 4A, comparing the results at time points with and without diamide, the level of transcription of *trfA* more than doubled at 30 min after diamide addition relative to that of untreated cells at an analogous OD. Remarkably, this induction of *trfA* transcripts nearly vanished at 60 min after diamide induction (i.e., at an OD₆₅₀ of 1.4), presumably due in part to the rapid inactivation of diamide within 20 to 30 min after addition (Ana Posada, personal communication).

TrfA is degraded by ClpCP. The adaptor MecA of *B. subtilis* shares 36% amino acid sequence identity to TrfA, primarily in the first 80 amino acids. MecA controls competence by targeting the transcriptional regulator ComK for degradation through ClpCP (46), and MecA is itself proteolytically regulated by this system (46). Although *S. aureus* does not contain the competence machinery of *B. subtilis*, it is of interest to ascertain if *S. aureus* utilizes

a similar method of posttranslational regulation. Accordingly, we induced the expression of TrfA with a codon-optimized N-terminal Myc tag using the xylose-inducible plasmid pEPSA5. The Myc tag was chosen because it has been demonstrated not to alter the pattern of proteolysis in other proteins (22). At an OD₆₅₀ of 0.6, *S. aureus* cells were induced with 0.2% xylose and grown to an OD₆₅₀ of 1.1. At this point, rifampin was added to stall transcription and samples were withdrawn at predetermined time points for 2 h. Whole-cell lysates of these aliquots were then examined by Western blotting with anti-Myc antibodies. As observed in Fig. 4B, TrfA levels decreased rapidly in the wild-type strain SH1000 with an ~22-min half-life, as determined by densitometric analysis. Similar rates were seen for the *clpB*, *clpQ*, and *clpX* deletion mutants. However, TrfA levels remained relatively steady in the Δ*clpC* and Δ*clpP* strains, indicating that the ClpCP protease system controls the degradation of TrfA, as it does in *B. subtilis*. These data, combined with the transcriptional transiency of *trfA* induction after thiol stress, suggest that TrfA levels can be rapidly induced under specific conditions and just as rapidly recede to preinduced levels following the cessation of a specific stress.

The combination of TrfA's proteolytic degradation and its transcriptional peak during late logarithmic phase gives a sense of how TrfA operates in the cell. However, to fully understand how TrfA changes during growth in medium, we were interested to determine whether the TrfA degradation rate was maintained at other growth phases. Similar to our previous time point experiments, we grew wild-type SH1000 expressing N-terminally Myc-

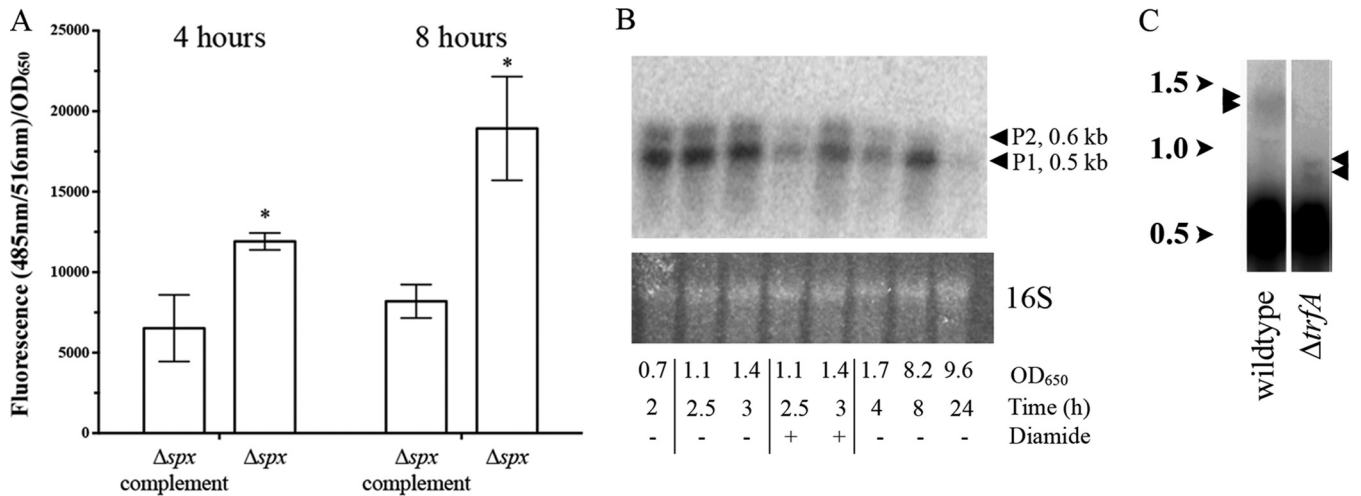


FIG 5 *spx* regulation of *trfA* and *spx* transcription. (A) GFP fluorescence driven by the *spx* P1 promoter on a shuttle plasmid containing a GFP reporter (pALC7803) in the Δ *spx* and *spx*-complemented strains at the 4- and 8-h time points (late log and early stationary phases, respectively). To minimize variations in fluorescence attributable to cell density, the data are presented as the average number of fluorescence units for triplicate samples per unit of absorbance at 650 nm. *, a statistically significant difference ($P < 0.05$) compared to the results for *spx*-complemented strain, as measured by Student's unpaired *t* test. (B) Northern blot of *spx* transcription from *S. aureus* SH1000 at various growth phases with and without 5 mM diamide. RNA (20 μ g each) was resolved on a denaturing agarose gel, blotted as described above, and probed with a 300-bp ³²P-radiolabeled *spx* DNA probe. As a loading control, ethidium bromide-stained 16S rRNA was included, and the results are shown at the bottom. Arrowheads, sizes of the transcripts generated from the P1 and P2 *spx* promoters. (C) Northern blots of RNA extracted from the *S. aureus* SH1000 and Δ *trfA* strains at the late exponential phase of growth (OD₆₅₀ = 1.1) and probed with a 300-bp ³²P-radiolabeled *spx* probe. Faint bands are identified by arrowheads to the side of each lane. The numbers to the left indicate kilobases.

tagged TrfA from pEPSA5 to various ODs under conditions of xylose induction and, following the addition of rifampin, took samples at various time points over 2 h. Western blots of lysates of these cells were probed with anti-Myc antibody and revealed that while cells in early logarithmic phase degraded TrfA similarly to those in late logarithmic phase (half-life, ~22 min), cells in stationary phase took longer to degrade TrfA (half-life, ~37 min) (Fig. 4C). This suggests that even with the decreased transcription of *trfA* in stationary phase (Fig. 4A), TrfA may still be present in stationary-phase cells.

***spx* activates *trfA* but represses its own promoter.** Previous studies of *spx* transcription by RT-PCR have shown that the gene product of *spx* likely activates one or more of these transcripts of *trfA* (28). Spx is a thiol-sensitive transcriptional regulator that is itself proteolytically regulated and is responsible for controlling the adaptation and survival of *S. aureus* in environments with oxidative stress.

To examine whether the transcriptional regulation of *trfA* by *spx* was subject to any autoregulation, we constructed transcriptional fusions of the *spx* P1 (distal) and P2 (proximal) promoters linked to GFP, as described in Materials and Methods, to yield pALC7803 and pALC7804, respectively. These plasmid constructs were used to transform the Δ *spx* mutant and its complemented strain of *S. aureus*. The results revealed that for the P2 promoter construct pALC7804, GFP levels were equivalent between the Δ *spx* and complemented strains (data not shown). However, GFP fluorescence levels were consistently higher in the Δ *spx* strain with the P1 promoter (Fig. 5A) at multiple time points than in its complemented strain. These data indicate that *spx* represses its own distal (P1) promoter but does not regulate its proximal (P2) promoter.

To better assess the growth conditions under which Spx represses its own promoter, we examined *spx* transcription at the

same times of growth used for measuring *trfA* transcription (Fig. 4A). Surprisingly, *spx* transcription was relatively constant through log and early stationary phases under unstimulated conditions (Fig. 5B). However, following the addition of 5 mM diamide, *spx* transcription was greatly reduced after 30 min (OD₆₅₀ = 1.1 with diamide) and recovered after 60 min as the diamide effect tapered (OD₆₅₀ = 1.4 with diamide). Interestingly, protein levels of Spx actually increase following diamide treatment in both *B. subtilis* and *S. aureus* (31, 47). This increase in Spx levels alongside a corresponding decrease in *spx* transcript levels following diamide treatment (Fig. 5B), as well as the increase in *spx* promoter fusion activity in a Δ *spx* strain in general (Fig. 5A), supports our notion that the Spx protein represses its own transcription in *S. aureus*.

We also examined the amount of transcriptional read-through that occurs from the *spx* locus into the adjacent gene, *trfA* (Fig. 5C). A *rho*-independent terminator was predicted between *spx* and the adjacent *trfA* gene ($-\Delta G = 14.5$ kcal/mol) (48), so to examine its effect on *trfA* transcription during cultured growth, we isolated RNA from wild-type strains grown to mid-log phase (OD₆₅₀ = 1.1). In addition to the two previously noted *spx* transcripts (28), several putative read-through bands at ~1.4 kb were also observed with extended exposure (Fig. 5C, arrowheads on the left). Relative to the smaller *spx* transcripts, these were estimated by densitometric analysis to be 3 to 4% of total *spx* transcripts. Importantly, these larger bands decreased in size in the Δ *trfA* strain, corresponding to the reduction of *trfA* (720 bp) in the mutant. Therefore, the terminator between *spx* and *trfA* is strong and able to terminate transcription in ~95% of cases.

TrfA does not regulate Spx protein levels in *S. aureus*. Spx has been shown to be proteolytically regulated by YjbH in *S. aureus* (31) and *B. subtilis* (49). However, YpbH, a MecA paralog with which TrfA shares 26% identity, also recognizes and degrades Spx

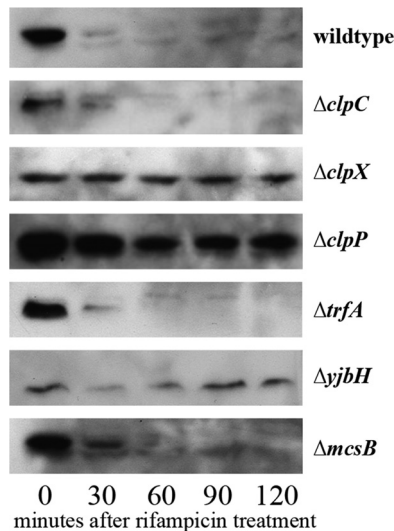


FIG 6 Role of *S. aureus* adaptors in Spx degradation. Western blots of whole-cell lysates of the SH1000 wild-type, $\Delta trfA$, $\Delta yjbH$, $\Delta clpC$, $\Delta clpX$, and $\Delta clpP$ strains with pALC7113 (producing Spx with an N-terminal myc tag). Cells were grown to an OD_{650} of 1.1 with 0.1% xylose and then treated with rifampin (200 μ g/ml). Cell lysates taken every 30 min were immunoblotted with a mouse anti-Myc primary antibody, followed by a horseradish peroxidase-conjugated goat anti-mouse antibody. The experiments were repeated three times, and representative blots are displayed.

in *B. subtilis* (50). To determine whether YjbH and TrfA share Spx as a substrate or whether another adaptor, such as McsB, was involved, we exogenously expressed Spx with an N-terminal myc tag from pALC7113 in various *clp* and adaptor deletion strains and tracked the degradation rates in transcriptionally stalled cells. As shown in Fig. 6, only in the *yjbH* mutant did the level of Spx persist through 120 min after rifampin treatment, whereas Spx was barely detectable at 30 min after rifampin treatment in the wild-type, $\Delta mcsB$, and $\Delta trfA$ strains. Furthermore, Spx levels remained high in the $\Delta clpX$ and $\Delta clpP$ strains but not the $\Delta clpC$ strains. Together, these data indicate that only *yjbH* acts as the facilitator of Spx breakdown in *S. aureus* through the ClpXP system.

TrfA is the adaptor regulating *S. aureus* antitoxin degradation. Toxin-antitoxin systems are small genetic loci that serve a variety of functions, from plasmid maintenance, chromosomal stabilization, and persistor formation to a bacterial form of programmed cell death (51). Several classes of toxin-antitoxin systems exist, based in part on whether the antitoxin is RNA (type I and type III) or protein (type II). In all cases, the antitoxin is short-lived due to its rapid degradation but is stabilized by its binding to its cognate toxin. The antitoxin also inactivates the toxin, protecting the bacteria until levels of antitoxin drop in response to a variety of environmental stresses (52). There are three known type II toxin-antitoxin systems in *S. aureus*, Axe1/Txe1, Axe2/Txe2 and MazEF, all of which are known to be proteolytically regulated by ClpCP (22). Given that TrfA works with ClpCP to degrade SsrA-tagged proteins in *S. aureus*, we wanted to determine whether TrfA also contributed to antitoxin degradation. For this purpose, the wild-type strain SH1000 and its isogenic *trfA* mutant were individually transformed with recombinant pEPSA5 plasmids carrying the antitoxin gene *mazE*, *axe1*, or *axe2*. For detection, we attached either a 5' or 3' myc tag (depending on the success of expression) to the antitoxin genes (Table 1) (22). At an

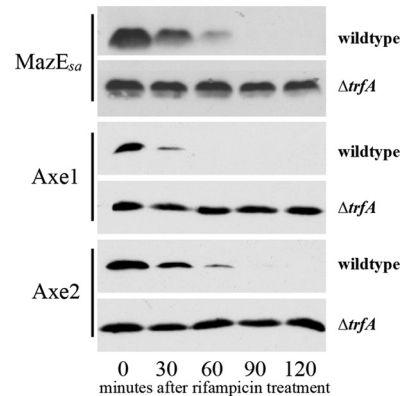


FIG 7 Degradation of antitoxins in *trfA* mutants. The SH1000 wild-type and $\Delta trfA$ strains with either pALC6188 (*mazE_{sa}*), pALC6486 (*axe1*), or pALC6489 (*axe2*) were grown to an OD_{650} of 1.1 with 0.1% xylose and then treated with rifampin (200 μ g/ml). Cell lysates taken every 30 min were immunoblotted with an anti-GFP primary antibody, followed by a horseradish peroxidase-conjugated goat anti-mouse antibody. The experiments were repeated three times, and representative blots are displayed.

OD_{650} of 0.6, cells were induced with 0.2% xylose and grown to an OD_{650} of 1.1, at which point the cells were transcriptionally stalled by addition of rifampin. Western blots of whole-cell lysates from these time points were probed with an anti-Myc antibody, disclosing that the levels of *S. aureus* MazE (*MazE_{sa}*), Axe1, and Axe2 all remained constant in the $\Delta trfA$ strain for 120 min after rifampin treatment, while those of the parental strain decreased rapidly after 30 to 60 min (Fig. 7). Degradation rates comparable to the rate for the wild type were also observed in the *mcsB* and *yjbH* mutants of *S. aureus* (data not shown). Together, these results demonstrate that the adaptor TrfA facilitates antitoxin degradation via ClpCP in *S. aureus* (22), hence linking a new set of proteins to adaptor-mediated proteolysis in bacteria.

DISCUSSION

S. aureus cells adjust the levels of their intracellular proteins through alterations in either transcriptional priorities or degradation rates. Chaperones and adaptors assist with the latter by tuning the substrate specificity of various proteases, allowing proteins to be selectively removed. One way that regulated proteolysis is utilized by many bacteria is to recover from errors in translation that may result in stalled ribosomes. In this process, tmRNA, a tRNA/mRNA hybrid that contains a small open reading frame for an 11-amino-acid tag called SsrA, enters the A site of the stalled ribosome, and as translation restarts, it is appended to the C terminus of the nascent protein. When newly tagged protein is freed from the ribosome, SsrA is then recognized as a signal for degradation. This is performed by both ClpXP and ClpAP (32, 53) in *E. coli*, while in *B. subtilis* only ClpXP is used (33).

Interestingly, we discovered that SsrA-mediated proteolysis in *S. aureus* is performed not by ClpXP but by ClpCP (Fig. 2A and B). Previous work has shown that ClpX is involved in regulating toxin production and virulence in *S. aureus* (16, 54), while ClpC plays a modulatory role in responding to stress, metabolic changes, and starvation (23). Therefore, it is not surprising that the degradation of SsrA-tagged proteins in *S. aureus* falls under the regulation of ClpCP (Fig. 8). Furthermore, SsrA degradation by proteases other than ClpXP and ClpAP is not without precedent, as Lon is used in

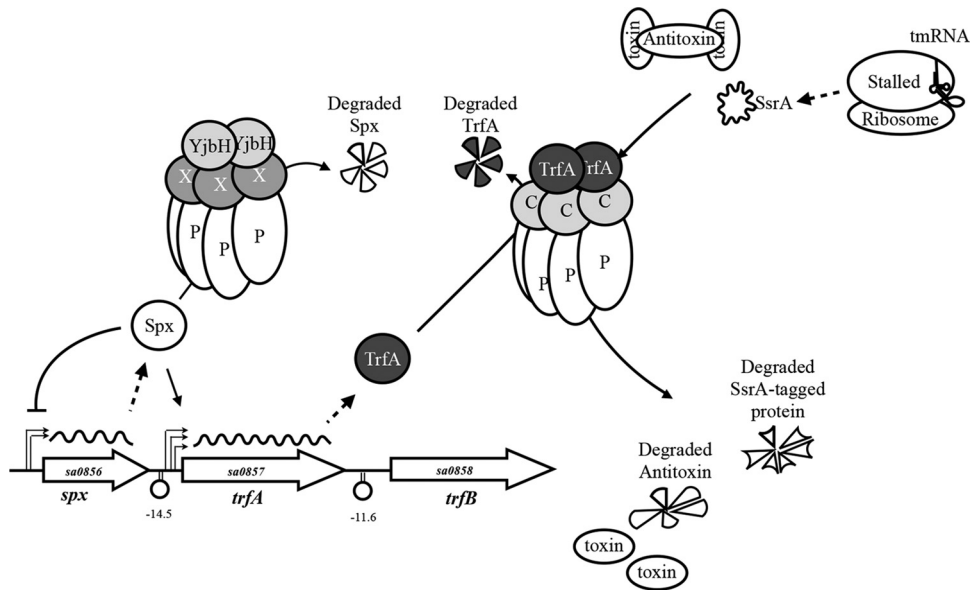


FIG 8 Proposed model for TrfA regulation and activity in *S. aureus*. The adaptor TrfA associates with ClpCP to selectively degrade SsrA-tagged proteins and class II antitoxins, such as MazE_{sp}. TrfA is itself degraded by ClpCP, which gives TrfA temporal nuance. The transcription of *trfA* is initiated from three upstream promoters, at least one of which is positively regulated by Spx. Spx activity is controlled by the degradation of the Spx protein via ClpXP and YjbH under normal conditions and with decreased proteolysis during oxidative, thiol, or antibiotic stress. Spx represses its own distal P1 promoter, creating a negative-feedback loop where less *spx* is transcribed during such stresses. X, ClpX; P, ClpP; C, ClpC.

such a capacity in the small genome of *Mycoplasma* (55), and Tsp (56), Lon (57), and FtsH (58) are all used in *E. coli* degradation of the SsrA tag, albeit at a lower frequency than ClpXP and ClpAP. While the reason that *S. aureus* swapped ClpXP for ClpCP in such a role is not known, ClpC in *S. aureus* shares some homology to ClpA in *E. coli* (36% identity, 56% similarity). It is possible that *S. aureus* (or a progenitor of it) utilized both ClpC/ClpA and ClpX for SsrA-mediated breakdown in a manner similar to that in which they are utilized in *E. coli*. However, as this bacterium evolved and ClpC became dominant in controlling the stress response and ClpX became dominant in controlling virulence, the ability of ClpX to degrade SsrA-tagged proteins may have become less of a necessity and ultimately lost entirely.

In *E. coli*, adaptors have long been known to complement chaperones in controlling SsrA-mediated degradation. For example, the adaptor SspB normally competes for the same SsrA motif as ClpA, blocking its recognition of SsrA-tagged proteins (25). SspB levels increase during heat shock (25), effectively redirecting ClpAP to degrade other targets, including misfolded proteins. No homologs of SspB or any other *E. coli* adaptors exist in *S. aureus*, so the mechanism by which TrfA mediates SsrA-mediated degradation in *S. aureus* may be novel (Fig. 8). Although the 3 C-terminal amino acids of the SsrA tag are sufficient for TrfA-mediated degradation of SsrA-tagged proteins in *S. aureus*, it is plausible that an additional factor(s) may mediate recognition of the entire native SsrA tag by ClpCP since Venus-SsrA appears to be degraded faster than Venus-VAA (Fig. 1A). Furthermore, as TrfA seems to have the greatest influence on SsrA degradation during late logarithmic and early stationary phases, the yet unidentified adaptor(s) may assist with degradation during other growth states.

The regulation of TrfA has also begun to be unraveled by our studies. Similar to MecA in *B. subtilis* (46), TrfA is degraded by the very protease that it assists in proteolysis. The 22-min half-life of

TrfA likely ties its activity to its transcription rate, suggesting that TrfA functions largely during logarithmic and early stationary growth phases, as well as to the oxidative stress response. In the last case, the transcription of *trfA* is upregulated in an Spx-dependent manner (28, 59), but it is quickly restored to baseline levels following abatement of thiol-related stresses (Fig. 4A). Such a pulse of transcription during these oxidative stresses could temporarily redirect ClpCP to process more TrfA-specific targets.

The thiol stress-sensitive transcriptional regulator Spx is degraded by ClpXP under normal growth conditions (60), but it is stabilized during oxidative stress in *S. aureus* (28, 31), allowing levels to rise. YjbH has been shown to contribute to this regulation in *S. aureus* (31), but YpbH, an adaptor in *B. subtilis* that has 26% identity to TrfA, has been shown to also control Spx degradation (50). We clarified this uncertainty in *S. aureus* by determining that only YjbH regulates Spx degradation (Fig. 6 and 8).

In *B. subtilis*, one of the three promoters of *spx* is upregulated following thiol stress (61), which further contributes to an increase in Spx levels. On the basis of our data, an alternative system appears to exist in *S. aureus*, where Spx (directly or indirectly) represses its own promoter. Accordingly, following the increase in Spx levels after exposure to 5 mM diamide, the transcription of *spx* is repressed. Considering the *spx*-dependent activation of *trfA*, this suggests that Spx can act as both an activator and a repressor in *S. aureus*. This brings the Spx function in line with that in *B. subtilis*, wherein it functions both to enhance RNA polymerase activity (47) and to act as an anti- α factor repressing transcription (62).

Besides SsrA-tagged proteins, our studies here also expand the set of proteins regulated by TrfA to include the labile antitoxins of the type II toxin-antitoxin systems (Fig. 7 and 8). This finding is of interest to us because we have shown previously that ClpCP is the major proteolytic system for the degradation of class II antitoxins

in *S. aureus* (22). While these protein-based antitoxins have been known to be degraded by a variety of ATP-dependent proteases and chaperones (52), to our knowledge, this is the first time that an adaptor has been linked to antitoxin degradation. As toxin-antitoxin systems are intimately linked to environmental and antibiotic-induced stresses, our results indicate that an adaptor such as TrfA may also participate in this stress response by modulating the degradation of antitoxins in *S. aureus*.

ACKNOWLEDGMENT

This work was supported by NIH grant RO1 AI91801 (to A.L.C.).

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