


SpoVG Regulates Cell Wall Metabolism and Oxacillin Resistance in Methicillin-Resistant *Staphylococcus aureus* Strain N315

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Increasing cases of infections caused by methicillin-resistant *Staphylococcus aureus* (MRSA) strains in healthy individuals have raised concerns worldwide. MRSA strains are resistant to almost the entire family of β -lactam antibiotics due to the acquisition of an extra penicillin-binding protein, PBP2a. Studies have shown that *spoVG* is involved in oxacillin resistance, while the regulatory mechanism remains elusive. In this study, we have found that SpoVG plays a positive role in oxacillin resistance through promoting cell wall synthesis and inhibiting cell wall degradation in MRSA strain N315. Deletion of *spoVG* in strain N315 led to a significant decrease in oxacillin resistance and a dramatic increase in Triton X-100-induced autolytic activity simultaneously. Real-time quantitative reverse transcription-PCR revealed that the expression of 8 genes related to cell wall metabolism or oxacillin resistance was altered in the *spoVG* mutant. Electrophoretic mobility shift assay indicated that SpoVG can directly bind to the putative promoter regions of *lytN* (murein hydrolase), *femA*, and *lytSR* (the two-component system). These findings suggest a molecular mechanism in which SpoVG modulates oxacillin resistance by regulating cell wall metabolism in MRSA.

Staphylococcus aureus is a versatile and dangerous pathogen in humans, especially while the frequency of staphylococcal infections caused by methicillin-resistant *Staphylococcus aureus* (MRSA) increases steadily (1). *S. aureus* has four penicillin-binding proteins (PBPs) which are involved in the last stages of peptidoglycan synthesis (2). MRSA strains have acquired an extra PBP (PBP2a), encoded by the *mecA* gene on the staphylococcal cassette chromosome *mec* (SCC*mec*), which has a remarkably lower affinity for β -lactams than does PBP2 and is responsible for resisting these drugs (3). Besides the expression of PBP2a, which is responsible for a higher level of β -lactam resistance, *S. aureus* has another primary β -lactam resistance mechanism: the expression of β -lactamase enzymes, encoded by the *blaZ* gene, which hydrolyze β -lactams such as penicillin. In addition, several additional native genes, such as *vraSR*, *pbp4*, and *fem* (factor essential for methicillin resistance), have been identified as being essential to the full expression of oxacillin resistance (4–6). Inactivation of *femA* and *pbp4* genes in the presence of an intact *mecA* gene usually results in strains with a low resistance to oxacillin. The VraS/VraR two-component regulatory system regulates the genes which are associated with cell wall biosynthesis, such as *pbp2*, *sgtB*, and *murZ* (7).

Most MRSA and glycopeptide-intermediately resistant *Staphylococcus aureus* (GISA) isolates are resistant to Triton X-100-induced autolysis, while instances of reduced resistance have been observed with concomitant increases of the autolysis rate (8–11). The expression of autolytic enzymes, including Atl, LytM, LytN, and Sle1 (12–15), is controlled by pleiotropic regulators, such as MgrA, SarV, SarA, LytSR, and ArlSR (autolysis-related locus) (16–20). Downstream of the *lytSR* operon is the *lrgAB* locus, the expression of which is regulated by LytSR (17). The *cidABC* and *lrgAB* operons may have an important role in the control of staphylococcal murein hydrolase activity and encode proteins analogous to the bacteriophage-encoded holins and antiholins (21–23).

SpoVG was initially identified in *Bacillus subtilis* and is involved in an unknown mechanism in sporulation (24). In non-sporulating bacteria, the σ^B -controlled *spoVG* affects capsule synthesis, expression of virulence factors, and antibiotic resistance

(25–27). SpoVG is considered a site-specific DNA-binding protein (28). Except that *spoVG* is under the control of σ^B , a small RNA SprX negatively regulates *spoVG* expression by direct anti-sense pairings at the internal translation initiation signals of *spoVG*, without affecting YabJ translation (29). SpoVG is one of the target genes under the control of SprX, which is involved in glycopeptide resistance. However, the molecular mechanism of how *spoVG* participates in antibiotic resistance was largely unknown. In this study, we have investigated the role of *spoVG* in autolysis and resistance to oxacillin in MRSA strain N315.

MATERIALS AND METHODS

Bacterial strains, plasmids, and culture conditions. The bacterial strains and plasmids used in this study are listed in Table 1. *Escherichia coli* strains were cultivated with shaking (220 rpm) in lysogeny broth (LB) medium (Oxoid) or on lysogeny broth agar (LA) at 37°C. *S. aureus* strains were grown with shaking (220 rpm) in tryptic soy broth (TSB) medium (Difco) or on tryptic soy agar (TSA) at 37°C. When required, the media were supplemented with 100 μ g/ml of ampicillin or 50 μ g/ml of kanamycin for *E. coli* and 15 μ g/ml of chloromycetin for *S. aureus* strains.

Construction of the *spoVG* mutant strain. To create the *spoVG* mutant strain without extra genes introduced, the plasmid pBTs was used as previously described (30). The upstream and downstream regions of *spoVG*, which were amplified with primer pairs *spoVG*-mutant-up-F/*spoVG*-mutant-up-R and *spoVG*-mutant-down-F/*spoVG*-mutant-down-R (see Table S1 in the supplemental material), were

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TABLE 1 Strains and plasmids used in this study

Strain or plasmid	Relevant characteristic(s) ^a	Source or reference
<i>S. aureus</i> strains		
RN4220	8325-4, r ⁻ , initial recipient for modification of plasmids which are introduced into <i>S. aureus</i> from <i>E. coli</i>	NARSA ^b
N315	HA-MRSA, SCCmec type II	NARSA
N315Δ <i>spoVG</i>	N315 strain deletion of <i>spoVG</i>	This study
<i>E. coli</i> strains		
Trans1-T1	Clone host strain; F ⁻ Φ80 (<i>lacZ</i>) ΔM15Δ <i>lacX74 hsdR</i> (r _K ⁻ m _K ⁺) Δ <i>recA1398 endA1 tonA</i>	TransGen
BL21(DE3)	Express strain; F ⁻ <i>ompT hsdS_B</i> (r _B ⁻ m _B ⁻) <i>gal dcm</i> (DE3)	TransGen
Plasmids		
pBTs	Shuttle vector, temp sensitive, Amp ^r Chl ^r	
pBTsΔ <i>spoVG</i>	pBTs derivative, for <i>spoVG</i> deletion in strain N315, Amp ^r Chl ^r	This study
pET28a(+)	Expression vector with a hexahistidine tag, Kan ^r	Novagen
pET <i>SpoVG</i>	pET28a(+) derivative, with ORF of <i>spoVG</i> , Kan ^r	This study
pLI50	Shuttle vector, Amp ^r Chl ^r	Addgene
pLI <i>SpoVG</i>	pLI50 derivative, harboring ORF of <i>spoVG</i> and its promoter, Amp ^r Chl ^r	This study

^a r⁻, restriction system negative; Kan^r, kanamycin resistant; Amp^r, ampicillin resistant; Chl^r, chloramphenicol resistant.

^b NARSA, Network on Antimicrobial Resistance in *Staphylococcus aureus* (<https://www.beiresources.org/Collection/53/NARSA.aspx>).

ligated by SLiCE (31) to form an up-down fragment. Briefly, the two fragments were designed with a 20-bp overlap, the fragments (50 to 200 ng) were mixed at equimolar amounts, and then 1 μl of 1× SLiCE buffer (500 mM Tris-HCl, 100 mM MgCl₂, 10 mM dithiothreitol [DTT]; pH 7.5), 1 μl of SLiCE extract, 1 μl of 10 mM ATP, and double-distilled water (ddH₂O) were added to a final volume of 10 μl. The mixed solution was incubated at 37°C for 1 h and then amplified with primers *spoVG*-mutant-up-F/*spoVG*-mutant-down-R. The resultant fragment was digested with KpnI/EcoRI and then cloned into the KpnI/EcoRI-digested plasmid pBTs. The resulting plasmid, pBTsΔ*spoVG*, was first electroporated into *S. aureus* strain RN4220 for modification and subsequently transformed into strain N315. An allelic replacement mutant was selected using a previously described method (32) and was further confirmed by PCR and sequencing.

Plasmid construction. For the construction of complementary plasmid pLI*SpoVG*, DNA fragments covering the open reading frame (ORF) of *spoVG* and the promoter of the *yabJ*-*spoVG* operon were amplified from N315 genomic DNA using primer pairs *spoVG*-complement-F/*spoVG*-complement-R and *spoVG*-mutant-up-F/*SpoVG*-R (see Table S1 in the supplemental material), respectively. The two DNA fragments were ligated by SLiCE and the resulting fragment was then cloned into the KpnI-digested plasmid pLI50 (33).

The protein expression plasmid pET*SpoVG* was constructed by amplifying the ORF of *spoVG* with primers *SpoVG*-exp-F and *SpoVG*-exp-R (see Table S1) from strain N315 chromosomal DNA. The PCR product was digested with NcoI and XhoI and then cloned into the expression vector pET28a(+) (Novagen). The recombinant plasmid was sequenced to confirm that *spoVG* was free of mutations and in frame with the hexahistidine tag.

Oxacillin susceptibility assay. Determination of oxacillin MICs was performed by broth microdilution, as recommended by the Clinical and Laboratory Standards Institute (34). Population analysis profiles were established by plating appropriate dilutions of direct colony suspension on Mueller-Hinton agar with 2% NaCl containing increasing concentrations of oxacillin. The numbers of CFU were determined after 24 h of incubation at 37°C.

The growth experiments were performed as described elsewhere (20). In brief, overnight cultures of *S. aureus* were diluted into fresh TSB (Difco) with or without oxacillin to yield a starting optical density at 600 nm (OD₆₀₀) of 0.05 and inoculated into 50-ml flasks in a final volume of 10 ml. Oxacillin was added at sub-MIC. The mixtures were then incubated

with constant shaking (220 rpm) at 37°C, with the OD₆₀₀ measured hourly for total 9 h using a spectrophotometer (DU 800; Beckman Coulter). These assays were repeated at least three times, with similar results.

Triton X-100-induced autolysis assay. Triton X-100-stimulated autolysis was measured as described previously (35). Overnight-grown bacterial cells were diluted to an OD₆₀₀ of 0.05 in TSB and allowed to grow to the early exponential (OD₆₀₀ = 0.8) phase at 37°C with shaking (220 rpm). Cells were harvested, washed twice with 0.05 M Tris-HCl buffer (pH 7.5), resuspended in the original volume of Tris-HCl (0.05 M; pH 7.5) containing 0.05% (vol/vol) Triton X-100, incubated at 37°C with shaking, and checked for lysis by measuring the progressive decrease in absorbance (OD₆₀₀) at half-hour intervals using a microplate reader (Elx800; Bio-Tek). The experiment was repeated at least three times, with similar results.

Total RNA isolation and real-time quantitative reverse transcription-PCR (qRT-PCR). Overnight cultures of *S. aureus* were diluted 1:100 in TSB and cultivated to early exponential (OD₆₀₀ = 0.5), mid-logarithmic (OD₆₀₀ = 2.5), and early stationary (OD₆₀₀ = 6) phases. Cells were harvested and processed with 1 ml of RNAiso plus (TaKaRa) in combination with 0.1-mm-diameter silica beads in a FastPrep-24 automated system (MP Biomedicals). Residual DNA was removed with RNase-free DNase I (TaKaRa).

For the reverse transcription, the cDNAs were synthesized using a PrimeScript 1st Strand cDNA synthesis kit (TaKaRa) and qRT-PCR was performed with SYBR Premix *Ex Taq* (TaKaRa) using the StepOne real-time PCR system (Applied Biosystems). The quantity of cDNA measured by real-time PCR was normalized to the abundance of *hu* cDNA (36). All qRT-PCR assays were repeated at least three times.

Purification of SpoVG-His₆. Expression and purification of the recombinant SpoVG-His₆ protein were conducted as described previously, with modification (28). *E. coli* BL21(DE3) was transformed with the protein expression plasmid pET*SpoVG*, and the transformant was cultivated in LB at 37°C to an OD₆₀₀ of 0.5 and induced with 0.5 mM isopropyl-β-D-1-thiogalactopyranoside at 37°C for an additional 3 h. The cells were harvested and lysed by sonication in lysis buffer (50 mM Tris-HCl [pH 8.0], 300 mM NaCl). The hexahistidine-tagged SpoVG protein was purified with a nickel-nitrilotriacetic acid-agarose solution (Qiagen) by following the manufacturer's recommendation. The bound protein was eluted with an elution buffer (200 mM imidazole, 50 mM Tris-HCl [pH 8.0], 300 mM NaCl). The imidazole in the eluent was removed using a

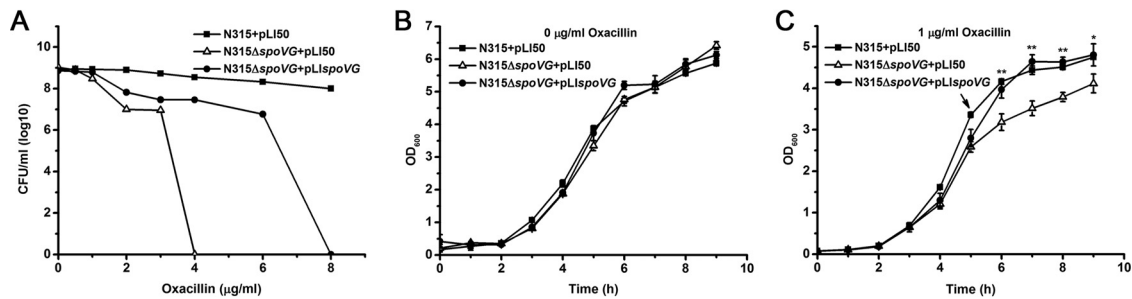


FIG 1 Oxacillin susceptibility assay. (A) Deletion of *spoVG* reduced oxacillin resistance in strain N315. The oxacillin resistance levels were evaluated by population analysis profiles as recommended by the Clinical and Laboratory Standards Institute. The colonies were counted after incubation at 37°C for 24 h. (B and C) Growth of the wild-type (N315+pLI50), the *spoVG* mutant (N315Δ*spoVG*+pLI50), and the *spoVG* complementary (N315Δ*spoVG*+pLI*spoVG*) strains in TSB broth at 37°C containing 0 or 1 μg/ml of oxacillin at the 5th hour. Statistically significant differences, calculated by the unpaired two-tailed Student *t* test, are indicated as follows: NS, not significant ($P > 0.05$); *, $P < 0.05$; **, $P < 0.01$; and ***, $P < 0.001$.

Centrifuge Biomax-5 column (Millipore), and the protein was stored with storage buffer (50 mM Tris-HCl, 25 mM KCl, 100 nM DTT, 10% [vol/vol] glycerol, 0.01% [vol/vol] Tween 20) at -80°C until use. Protein purity and concentration were assessed via SDS-PAGE and the bicinchoninic acid assay with bovine serum albumin as the standard. The final concentration of purified SpoVG was 30 μM.

EMSA. To determine whether the purified SpoVG binds to *lytN*, *lytSR*, and *femA* promoters, the biotin-labeled DNA fragments containing the putative promoter regions of *lytN* (254 bp), *lytSR* (253 bp), or *femA* (273 bp) were obtained by PCR with primers listed in Table S1 in the supplemental material. For electrophoretic mobility shift assay (EMSA), the biotin-labeled promoter regions of *lytN* (1 fmol), *lytS* (1 fmol), or *femA* (2 fmol) were incubated at 25°C for 30 min with various amounts of SpoVG in 10 μl of incubation buffer [50 mM Tris-HCl (pH 7.5), 1 mM DTT, 150 nM EDTA, 50 ng/ml of poly(dI-dC)]. After incubation, the mixtures were electrophoresed in a 5% native polyacrylamide gel in 1× Tris-borate-EDTA (TBE) buffer and then transferred to a nylon membrane in 0.5× TBE buffer. The band shifts were detected and analyzed according to the manufacturer's instructions. The images were obtained using ImageQuant LAS 4000 (GE). The unlabeled fragments of each promoter were added to the labeled fragments at a ratio of approximately 100:1 as specific competitors. The unlabeled DNA fragment of the *pta* ORF (100-fold or 200-fold) was added as a nonspecific competitor.

RESULTS

Influence of *spoVG* on oxacillin resistance in MRSA strain N315.

Either deletion of the *yabJ-spoVG* operon or inhibition of *spoVG* expression reduces antibiotic resistance in *Staphylococcus aureus* (25, 29). To investigate the regulatory mechanism in which SpoVG is involved in antibiotic resistance, we deleted *spoVG* in MRSA strain N315. Oxacillin resistance in the *spoVG* mutant was significantly reduced and the complementary strain exhibited a partial effect as shown by population analysis profiles (Fig. 1A). We also tested cell growth in TSB broth without antibiotic or exposed to 1 μg/ml of oxacillin (representing approximately one-quarter MIC for the mutant strain) at the 5th hour. No significant difference was observed when cells were grown in TSB broth without oxacillin (Fig. 1B), whereas the *spoVG* mutant exhibited a clear growth defect compared to the wild type in the medium containing oxacillin (Fig. 1C). In addition, there is no significant difference in vancomycin and teicoplanin resistance between the *spoVG* mutant and wild-type strains as shown by population analysis profiles (see Fig. S1 in the supplemental material). Ceftizoxime resistance in the *spoVG* mutant was significantly reduced, and the complementary strain also had a partial complementary action (see Fig. S1).

Effect of the *spoVG* mutation on Triton X-100-induced autolysis. Autolysis is commonly associated with the killing mechanism of penicillin and β-lactams. We examined Triton X-100-induced autolytic activity in the wild-type and *spoVG* mutant strains. The *spoVG* mutant exhibited increased autolysis rates compared with those of the wild type. The phenotype was restored by introducing the complementary plasmid pLI*spoVG* (Fig. 2), indicating that cell wall turnover was modulated by SpoVG.

Transcriptional analysis by qRT-PCR. The data presented above clearly indicate that SpoVG participates in the control of oxacillin resistance as well as in autolysis. Many genes involved in cell wall synthesis or cell wall degradation can influence methicillin resistance or autolysis. To determine whether the expression of these genes was altered in the *spoVG* mutant, we performed qRT-PCR to examine the mRNA levels of 17 potential target genes, among which 8 genes were associated with cell wall synthesis or oxacillin resistance (*femA*, *mecA*, *blaZ*, *pbp1*, *pbp2*, *pbp3*, *pbp4*, and *vraS*) and 9 genes were implicated in cell wall degradation (*atl*, *lytN*, *lrgA*, *arlS*, *sle1*, *lytS*, *cidA*, *lytM*, and *sarA*). The transcriptional levels of *mecA*, *femA*, and *blaZ* were decreased in the *spoVG* mutant during the early stationary phase of growth but not markedly altered at early exponential and mid-logarithmic phases (Fig. 3A). In addition to these genes, other target genes

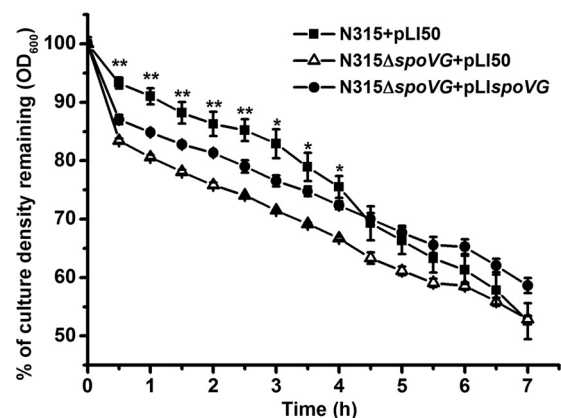


FIG 2 Triton X-100-induced autolysis. Autolysis of the wild-type (N315+pLI50), the *spoVG* mutant (N315Δ*spoVG*+pLI50), and the *spoVG* complementary (N315Δ*spoVG*+pLI*spoVG*) strains at 37°C in Tris-HCl buffer containing 0.05% Triton X-100 was determined by measuring changes in optical densities upon exposure to the detergent. *, $P < 0.05$; **, $P < 0.01$.

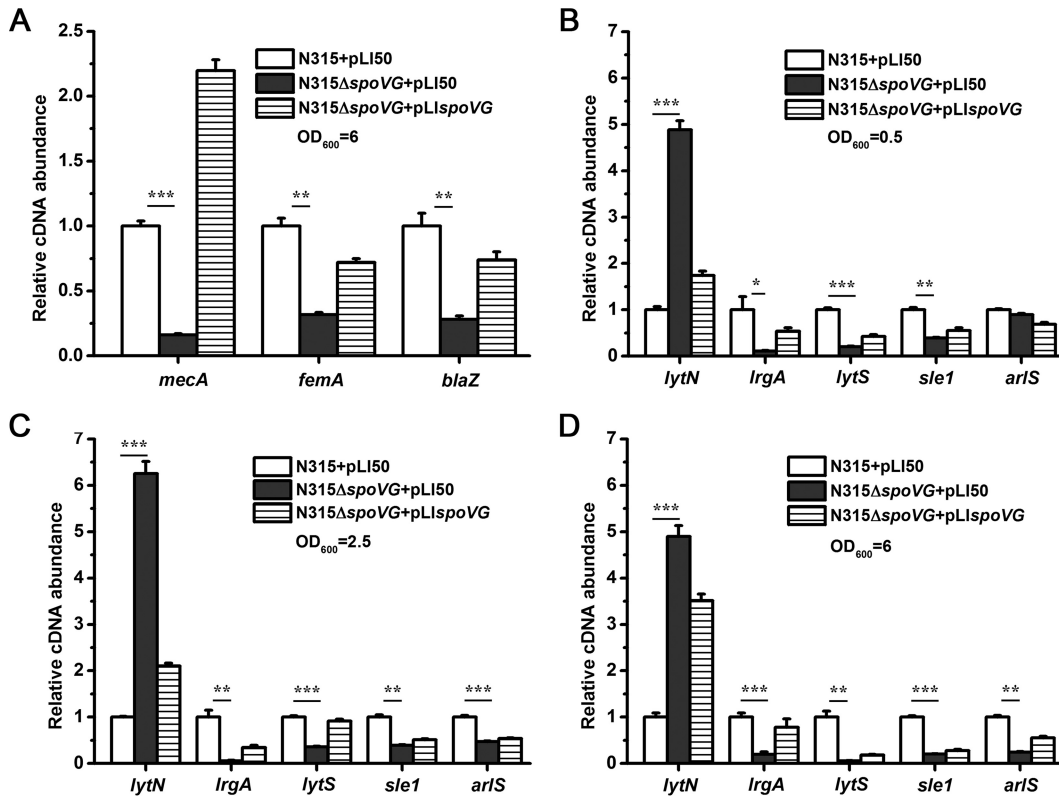


FIG 3 Transcriptional analysis of cell wall-related genes. The relative transcription levels of several cell wall biosynthesis- and hydrolysis-related genes in the wild-type (N315+pLI50), the *spoVG* mutant (N315Δ*spoVG*+pLI50), and the *spoVG* complementary (N315Δ*spoVG*+pLI*spoVG*) strains are shown. (A) Relative transcription levels of *femA*, *mecA*, and *blaZ* genes in early stationary phase of growth. (B to D) Relative mRNA levels of target genes (*lytN*, *lrgA*, and *sle1*) and regulatory loci (*lytSR* and *arlSR*) involved in autolysis in early exponential (OD₆₀₀ = 0.5), mid-logarithmic (OD₆₀₀ = 2.5), and early stationary (OD₆₀₀ = 6) phases. *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001.

involved in methicillin resistance (*pbp1*, *pbp2*, *pbp3*, *pbp4*, and *vraS*) were not significantly changed in the mutant strain (see Fig. S2 in the supplemental material). A significant increase in the transcript level of *lytN*, encoding a murein hydrolase, was observed in the *spoVG* mutant compared with those in the wild-type and complemented strains in the different growth stages (Fig. 3B to D). In contrast, the mRNA levels of *lrgA* (antiholin) and *sle1* (murein hydrolase) were significantly reduced in the *spoVG* mutant (Fig. 3B to D). The transcriptional levels of *lytS* and *arlS* were also found to be decreased in the mutant strain (Fig. 3B to and D). The transcript levels of *atl*, *lytM*, *cidA*, *mgrA*, and *sarA* were not significantly affected in the *spoVG* mutant (see Fig. S2). These data

indicate that SpoVG controls regulatory genes as well as target genes involved in cell wall metabolism.

Binding of SpoVG to the putative promoter regions of *lytN*, *femA*, and *lytSR*. To test whether the target genes are under the direct control of SpoVG, EMSA was performed with biotin-labeled putative promoter regions and the recombinant SpoVG (SpoVG-His₆). The results showed that SpoVG can retard the mobility of the *lytN* promoter in a dose-dependent manner (Fig. 4A). This shifted band disappeared in the presence of an approximately 100-fold excess of unlabeled *lytN* promoter DNA, but not in the presence of a 200-fold excess of an unlabeled coding sequence DNA of *pta*. EMSA was also performed using the promoter re-

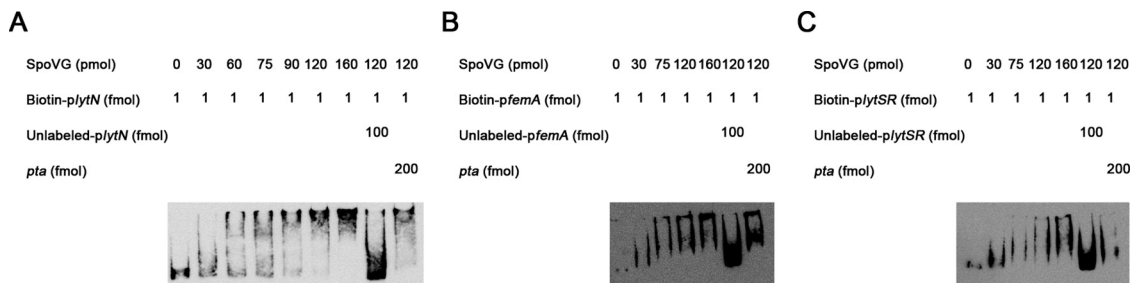


FIG 4 Electrophoretic mobility shift assay of the purified SpoVG with the biotin-labeled DNA fragments containing the putative promoter regions of *lytN* (254 bp), *femA* (273 bp), or *lytSR* (253 bp). Increasing concentrations of purified SpoVG and 1 or 2 fmol of the biotin-labeled probe were used in the reactions. The specific competitor concentration was 100-fold that of the labeled promoter, and the concentration of the nonspecific competitor (*pta*) was 100-fold or 200-fold.

gions of *femA* and *lytSR*, and similar band shift patterns were observed with the *femA* and *lytSR* promoters (Fig. 4B and C). These data suggest that SpoVG can regulate the expression of *lytN*, *femA*, and *lytSR* by directly binding to their promoter regions.

DISCUSSION

Previous studies have shown that SpoVG is the major factor of the *yabJ-spoVG* operon required in *S. aureus* for antibiotic resistance. In this study, we have also found that *spoVG* deletion can reduce oxacillin resistance in MRSA strain N315. SpoVG can directly regulate *fntB*, which is one of the *fem* or *aux* genes (26, 28). It has been reported that *femA*, also one of *fem* or *aux* genes, is involved in the synthesis of pentaglycin cross-links (5). Our studies indicate that SpoVG acts as a positive regulator for *femA*. A decrease of the mRNA level of *femA* in the *spoVG* mutant implies that the synthesis process of peptidoglycan cross-links may be damaged, thus reducing oxacillin resistance. With the normal function of PBP2, the decreased expression of *mecA* is likely to be associated with the lower resistance to oxacillin. In addition, the decrease in the transcript level of *blaZ* in the *spoVG* mutant suggests that the ability to hydrolyze β -lactams decreased. PBP2 is the only native PBP which is capable of both transpeptidation and transglycosylation of the peptidoglycan in *S. aureus*. PBP2a is capable of replacing the transpeptidase domain of PBP2, and the transglycosylase domain of PBP2 is necessary for resistance (37, 38). PBP4 has a low affinity for most β -lactam antibiotics and possesses transpeptidase and carboxypeptidase activities. The deletion of *pbp4* in MRSA strain MW2 affects the transcription of *pbp2* in cells challenged with oxacillin (6). Disruption of *pbp4* also caused increased sensitivity to ceftizoxime in the ZOX3 mutant, which has a decreased affinity to ceftizoxime in the ZOX3 mutant, which has a decreased affinity of its PBP2 variant for ceftizoxime (39). Therefore, two native *S. aureus* transpeptidases (PBP2 and PBP4) and an acquired transpeptidase (PBP2A) are likely to cooperate functionally in the biosynthesis of peptidoglycan and susceptibility to antimicrobial agents. How the activities of PBP1, which is involved in the enhancement of the activity of daptomycin against MRSA in the presence of β -lactam antibiotics, and the “nonessential” PBP3 of *S. aureus* interact in this cooperative system remains to be explored (40, 41). The *VraS/VraR* system regulates many cell wall biosynthesis-related genes, and the inactivation of it yielded significant decrease of resistance against teicoplanin, β -lactams, bacitracin, and fosfomycin (7). However, the transcript levels of *pbp1*, *pbp2*, *pbp3*, *pbp4*, and *vraS* were not affected by SpoVG in strain N315. In addition, our results indicate that there were no dramatic differences in vancomycin and teicoplanin resistances between the *spoVG* mutant and wild-type strains. This result is not in agreement with the previous reports (25, 29), perhaps due to the diverse genetic backgrounds of strains.

Since our data in this study were consistent with the previous report that reduced resistance was observed with concomitant increases of autolysis rate in *S. aureus* isolates, we further investigated the mRNA levels of cell wall degradation-related genes by qRT-PCR. Two regulatory systems, LytSR and ArlRS, have been shown to modulate autolytic activity negatively (17, 22). LytSR positively regulates *lrgA* and *lrgB*, a set of genes downstream of *lytSR*. Our studies indicate that SpoVG acts as a positive regulator for *lytSR* and, in turn, activates *lrgA* and leads to the repression of murein hydrolase activity. LytN, a murein hydrolase, is negatively regulated by SpoVG, which is consistent with the hypothesis that SpoVG is a negative regulator of autolysis in *S. aureus*.

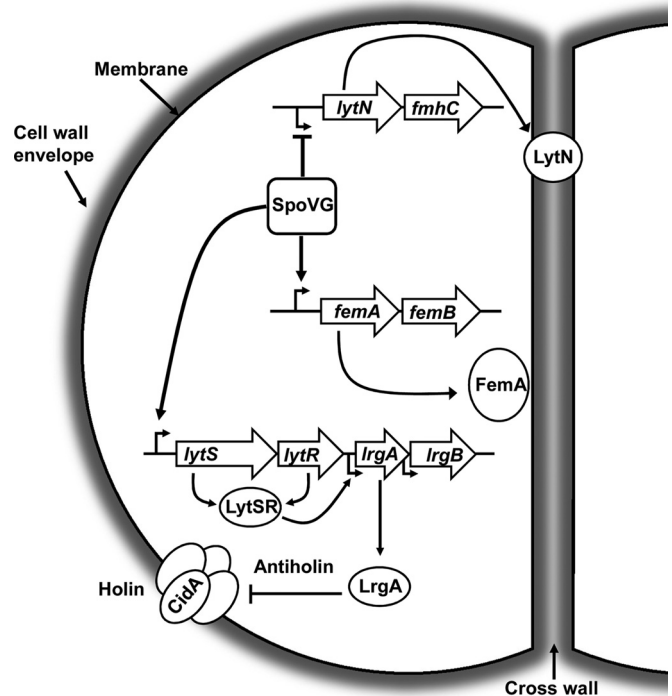


FIG 5 SpoVG is involved in the control of cell wall metabolism and oxacillin resistance in *Staphylococcus aureus* strain N315. SpoVG positively regulates the expression of *femA*, which is involved in the synthesis of pentaglycin cross-links. LytN, a murein hydrolase, is negatively regulated by SpoVG, which is consistent with the hypothesis that SpoVG is a negative regulator of autolysis in *S. aureus*. Besides, SpoVG acts as a positive regulator for *lytSR* and, in turn, activates *lrgA*, leading to the repression of murein hydrolase activity.

Our data suggest that SpoVG indirectly affects the expressions of *arlRS* and *sle1*. Sle1 is a peptidoglycan hydrolase which has been implicated in cell separation of *S. aureus*. But, unlike Atl (a major autolysin), Sle1 is not directly involved in autolysis of *S. aureus* (14). In summary, these results indicate that SpoVG as a transcriptional factor regulates the expression of *lytN*, *femA*, and *lytS* genes involved in cell wall metabolism (Fig. 5). Even though we identified several genes involved in oxacillin resistance and autolysis under the control of SpoVG, it cannot be excluded that other genes or small regulatory RNAs may be involved in the regulatory process.

Previous studies have shown that the 5' noncoding regions of *cap5*, *fntB*, *esxA*, and *lukED* all contain at least two 5'-TAATT_{T/A}-3' sequences to which SpoVG binds with high affinity and specificity (28). In our study, we have also found at least two 5'-TAATT_{T/A}-3' sequences existing in the 5' noncoding regions of *lytN*, *femA*, and *lytSR*. Whether this motif or other surrounding DNA sequences or structures contribute to the binding of SpoVG to target genes remains to be determined.

In conclusion, our findings reveal that SpoVG modulates β -lactam antibiotic resistance through promoting cell wall synthesis and inhibiting cell wall degradation in the presence of an intact *mecA* gene. While the molecular mechanism of *mecA* expression regulated by *mecR1-mecI* has been widely recognized, antibiotic resistance caused by auxiliary factors deserves further study.

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