

## PerR Controls Oxidative Stress Resistance and Iron Storage Proteins and Is Required for Virulence in *Staphylococcus aureus*

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**The *Staphylococcus aureus* genome encodes three ferric uptake regulator (Fur) homologues: Fur, PerR, and Zur. To determine the exact role of PerR, we inactivated the gene by allelic replacement using a kanamycin cassette, creating strain MJH001 (*perR*). PerR was found to control transcription of the genes encoding the oxidative stress resistance proteins catalase (KatA), alkyl hydroperoxide reductase (AhpCF), bacterioferritin comigratory protein (Bcp), and thioredoxin reductase (TrxB). Furthermore, PerR regulates transcription of the genes encoding the iron storage proteins ferritin (Ftn) and the ferritin-like Dps homologue, MrgA. Transcription of *perR* was autoregulated, and PerR repressed transcription of the iron homeostasis regulator Fur, which is a positive regulator of catalase expression. PerR functions as a manganese-dependent, transcriptional repressor of the identified regulon. Elevated iron concentrations produced induction of the PerR regulon. PerR may act as a peroxide sensor, since addition of external hydrogen peroxide to 8325-4 (wild type) resulted in increased transcription of most of the PerR regulon, except for *fur* and *perR* itself. The PerR-regulated *katA* gene encodes the sole catalase of *S. aureus*, which is an important starvation survival determinant but is surprisingly not required for pathogenicity in a murine skin abscess model of infection. In contrast, PerR is not necessary for starvation survival but is required for full virulence ( $P < 0.005$ ) in this model of infection. PerR of *S. aureus* may act as a redox sentinel protein during infection, analogous to the *in vitro* activities of OxyR and PerR of *Escherichia coli* and *Bacillus subtilis*, respectively. However, it differs in its response to the metal balance within the cell and has the added capability of regulating iron uptake and storage.**

The relationship between invading pathogenic bacteria and their host is dynamic, with bacteria having to rapidly adapt to the hostile and changing environment which they have entered. Metal ion acquisition is essential for pathogen proliferation, and limitation is a nonspecific host response to infection (10), which reduces the ability of bacteria to replicate and increases their susceptibility to clearance by the immune system. Iron is an essential nutrient *in vivo* and together with manganese is an important cofactor for bacterial antioxidant defense enzymes, e.g., catalase, peroxidase, and superoxide dismutase (SOD) (1, 48). Consequently, bacteria have evolved specialized proteins that monitor metal ion levels and respond accordingly by regulating gene expression (31, 51). These metalloregulatory proteins cluster in four distinct families represented by Fur (ferric uptake regulator), DtxR (diphtheria toxin repressor), MerR, and ArsR (53). The well-characterized DtxR from *Corynebacterium diphtheriae* (61) and Fur (26) have similar roles with respect to iron homeostasis and toxin synthesis; however, these two proteins share little amino acid homology, and their consensus DNA binding sequences are different.

Four metal ion-dependent repressors have been identified in *Bacillus subtilis*: three Fur-like proteins, Fur, PerR, and Zur (13, 28), and the recently identified DtxR-like protein, MntR (53). Fur controls iron homeostasis via a regulon of iron trans-

porters, iron siderophore transporters, and siderophore biosynthesis proteins. Zinc homeostasis is maintained by Zur-mediated repression of two operons encoding zinc uptake transporters. MntR is a bifunctional, manganese-responsive regulator of two manganese transporters, MntABCD and MntH, which are both selectively repressed by high levels of Mn(II) while MntR functions as a positive regulator of the *mntABCD* operon under low-Mn(II) growth conditions (53). MntH belongs to the Nramp family of proteins and has been proposed previously to have a role in the pathogenesis of *Salmonella enterica* serovar Typhimurium (42).

Genetic evidence revealed that *B. subtilis* PerR is a manganese- and iron-responsive transcriptional repressor of the genes encoding a catalase, alkyl hydroperoxide reductase (AhpCF); a Dps homologue (MrgA); and heme biosynthesis enzymes (13, 17, 18). Furthermore, PerR was hydrogen peroxide responsive, a property that makes it functionally analogous to OxyR, the well-characterized peroxide stress regulator in *Escherichia coli*. It was proposed elsewhere that peroxide stress activation of *B. subtilis* PerR was linked to the bound metal ion, a feature that would separate it mechanistically from OxyR, which is activated by H<sub>2</sub>O<sub>2</sub>-catalyzed disulfide bond formation and is not metalliferous (13, 69). A homologue of PerR in the gram-negative pathogen *Campylobacter jejuni* was similarly shown to repress catalase and AhpC in response to iron; however, regulation by manganese was not investigated (62). The *Streptococcus pyogenes* PerR was shown to be required for an inducible peroxide stress response (43). Intriguingly, this inducible response did not involve AhpC or MrgA, and it was

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suggested that a novel mechanism of peroxide stress management may exist in *S. pyogenes* (43).

*Staphylococcus aureus* successfully colonizes humans and commonly forms part of the flora of the anterior nares and the skin. Its versatility makes it an important pathogen that can infect a diversity of tissues causing a wide spectrum of diseases (24). Many staphylococcal virulence determinants are known and include adhesins, hemolysins, proteases, and superantigenic toxins that are mainly controlled by the interdependent, global regulators Agr and SarA (16, 20, 44). Intracellular survival in neutrophils (30), endothelial cells (63), epithelial cells (8), and osteoblasts (38) has been described previously, suggesting that both intracellular survival and extracellular multiplication play important roles in the pathogenesis of *S. aureus* infections. The determinants that promote in vivo survival intracellularly are poorly defined.

*S. aureus* has homologues of the four main metal ion-dependent repressors, Fur, PerR, Zur, and MntR. *S. aureus* Fur has been shown to be an iron-dependent repressor in vitro (68). Recently, we have shown that Fur represses the transcription of a number of iron uptake transporters and positively regulates, either directly or indirectly, catalase expression (37). Inactivation of *fur* was shown to impact on virulence, although this may simply be due to the fact that under some conditions growth rate was affected by the loss of Fur (37). *S. aureus* Zur mediates zinc homeostasis but is not important for virulence, and the functions of PerR and MntR have yet to be determined. The *Staphylococcus epidermidis* MntR homologue SirR was identified in vitro as an iron-regulated DtxR-like protein (34).

The importance of in vivo expression of the oxidative stress enzymes catalase and SOD has been suggested through the analysis of clinical isolates with reduced levels of expression of these enzymes (40, 46). In contrast, in vivo expression of one of the two SODs, SodA, was shown not to be important for pathogenicity despite contributing to survival in vitro (21).

In this report, we demonstrate that PerR controls a regulon of oxidative stress resistance and iron storage proteins in *S. aureus*. Furthermore, we demonstrate that PerR-dependent control of this regulon is important for pathogenicity.

## MATERIALS AND METHODS

**Media and growth conditions.** *S. aureus*, and *E. coli* strains and plasmids are listed in Table 1. *E. coli* was grown in Luria-Bertani (LB) medium at 37°C. *S. aureus* was grown at 37°C with shaking at 250 rpm in brain heart infusion (BHI) broth (Oxoid), in chemically defined medium (CDM) (64), or in chemically defined metal-limitation medium (CL) (37). CLR medium consists of CL (which contains 400 µM magnesium sulfate) without glucose and replete with the following metals added at an 0.2 µM final concentration: calcium chloride, copper sulfate, ferrous sulfate, manganese chloride, nickel sulfate, molybdenum sulfate, and zinc sulfate. Colonies from non-Chelex-treated CL agar plates were used to inoculate a CLR preculture. Experimental 25-ml cultures in acid-washed, 250-ml flasks were inoculated at a starting optical density at 600 nm ( $OD_{600}$ ) of 0.002 prior to growth at 37°C. When included, antibiotics were added at the indicated concentrations: ampicillin, 100 mg liter<sup>-1</sup>; kanamycin, 50 mg liter<sup>-1</sup>; neomycin, 50 mg liter<sup>-1</sup>; and erythromycin and lincomycin, 5 and 25 mg liter<sup>-1</sup>, respectively.

**Construction of strains.** Derivatives of plasmid pAZ106, an integrating plasmid conferring resistance to erythromycin and containing a promoterless *lacZ* gene (16), or plasmid pAUL-A, a temperature-sensitive integrating plasmid conferring resistance to erythromycin (15), were constructed using PCR with *Pwo* polymerase (Roche) and standard cloning techniques (56). A plasmid for disrupting *perR* was constructed by PCR amplification of a 2-kb *perR* fragment

using primers OL7, OL8, OL9, and OL10 (Table 1) to insert an internal *ClaI* site by the site-directed mutagenesis method of Higuchi (33). This fragment was cloned into pAUL-A, creating pMAL5, and then a 1.5-kb *ClaI* fragment containing a kanamycin cassette from pDG782 (32) was inserted to produce pMAL7.

Transcriptional reporter fusions to the *perR*, *fur*, *kataA*, *ahpC*, *mrgA*, *bcp*, *ftn*, and *txxB* genes were made by PCR amplification of suitable DNA fragments using primers detailed in Table 1 followed by cloning into pAZ106, creating the plasmids pMAL9, pMAL10, pMAL11, pMAL12, pMAL13, pMAL28, pMAL29, and pMAL30, respectively. Typically, between 0.6 and 1.4 kb of DNA encompassing the start and promoter regions was amplified using *Pwo* polymerase. The purified DNA fragments were digested with *Bam*HI and *Eco*RI and cloned into plasmid pAZ106 digested with the same enzymes. The *perR* gene, amplified using primers OL82 and OL83, was cloned into the *E. coli*-*S. aureus* shuttle vector pCU1 (7), producing plasmid pMAL34.

All of the above suicide plasmids were integrated into the chromosome of electrocompetent *S. aureus* RN4220 (58) through homology with the parental copies by a Campbell-type event. These were then transduced into 8325-4 using phage φ11 (50), and individual clones were verified by Southern blotting or genomic DNA sequencing to confirm the structural integrity of the DNA at the integration site.

Random Tn917 insertions into the chromosome of *S. aureus* 8325-4 were created as described in the work of Watson et al. (65). Screening of transposants for a starvation survival defect was performed on CDM agar as described in the work of Watson et al. (65) with limiting, 0.1% (wt/vol) concentrations of glucose. Comparative starvation survival experiments were performed in glucose-limiting CDM with shaking (250 rpm) at 37°C.

**Genomic DNA sequencing.** DNA was isolated from lysed cells of overnight cultures of *S. aureus* using lysostaphin (100 µg ml<sup>-1</sup>) and Qiagen genomic DNA columns. Three micrograms of genomic DNA was sequenced from both ends of the transposon insertion using 16 µl of Big-Dye (ABI) premix in a 40-µl volume with 13 pmol of primer Tn1 or Tn2 (Table 1) in a 90-cycle PCR sequencing program (95°C, 30 s; 50°C, 20 s; 60°C, 4 min). PCR sequencing products were purified using DyeEx columns (Qiagen).

**β-Galactosidase assays.** Levels of β-galactosidase activity were measured as described previously (37). Briefly, 0.1-ml samples were harvested, and cell pellets were stored at -20°C. Thawed pellets were resuspended in 0.5 ml of ABT buffer (60 mM K<sub>2</sub>HPO<sub>4</sub>, 40 mM KH<sub>2</sub>PO<sub>4</sub>, 100 mM NaCl). The assay was started with the addition of 50 µl of freshly prepared MUG (4-methylumbelliferyl-β-D-galactoside) (10 mg ml<sup>-1</sup>), and the assay mixture was incubated at 25°C for 60 min. The assay was stopped with the addition of 0.5 ml of 0.4 M Na<sub>2</sub>CO<sub>3</sub>. The stopped assay mixture was then serially diluted in a 50:50 (vol/vol) mixture of ABT and Na<sub>2</sub>CO<sub>3</sub> in 96-well microtiter plates (Nunc). Fluorescence was measured using a Victor plate reader (Wallac) with a 0.1-s count time and calibrated with standard concentrations of MU (4-methylumbelliferone). One unit of β-galactosidase activity was defined as the amount of enzyme that catalyzed the production of 1 pmol of MU min<sup>-1</sup> OD<sub>600</sub> U<sup>-1</sup>. Assays were performed on duplicate samples, and the values were averaged. The results presented here were representative of three independent experiments that showed less than 20% variability.

**Catalase assay activity gels, H<sub>2</sub>O<sub>2</sub> challenge, and induction.** Catalase activity was detected after electrophoresis on a 10% (wt/vol) native polyacrylamide gel, pH 7.5 (45), of washed cells lysed with lysostaphin (100 µg ml<sup>-1</sup>). The double-staining method of Wayne and Diaz (66) was used to visualize bands of activity. Catalase activity was assayed spectrophotometrically at 240 nm as described by Beers and Sizer (9) using 50 mM potassium phosphate buffer (pH 7.0) with 19.6 mM hydrogen peroxide. Protein concentration was measured by the method of Bradford (12) using bovine serum albumin (fraction V; Sigma) as the standard. Hydrogen peroxide resistance was assayed as described in the work of Watson et al. (64) with the following modifications. Cells grown to exponential phase in CLR were washed and diluted in phosphate-buffered saline (PBS) to an OD<sub>600</sub> of 0.2 and, following challenge with 7.5 mM H<sub>2</sub>O<sub>2</sub>, were diluted in PBS containing 10 mg of catalase ml<sup>-1</sup> and then serially diluted in PBS. Viability was assessed by overnight growth on BHI agar. Hydrogen peroxide induction of gene expression using *lacZ* fusions was performed on cultures grown in CLR or Chelex-treated BHI medium to OD<sub>600</sub>s of 0.2, 2, and 8 by adding H<sub>2</sub>O<sub>2</sub> to a final concentration of 100 or 500 µM with shaking (250 rpm) at 37°C. Samples were removed for assay before and 60 min after the addition of H<sub>2</sub>O<sub>2</sub> and assayed for β-galactosidase.

**Cellular protein preparation.** The soluble cellular protein fraction was prepared by pelleting cells and resuspending them in 5 µl per 0.1 OD<sub>600</sub> U of 1 M Tris-HCl, pH 6.8, containing 20 µg of lysostaphin (Sigma). After 10 min of incubation at 37°C, an equal volume of Laemmli sample buffer was added before boiling for 10 min. The samples were centrifuged for 5 min before loading on a

TABLE 1. Strains, plasmids, and primers used in this study

Strain, plasmid, or primer	Genotype, description, or sequence <sup>a</sup>	Reference or source
<b>Strains</b>		
<i>E. coli</i>		
DH5 $\alpha$	$\phi$ 80 $\Delta(lacZ)M15 \Delta(argF-lac)U169 endA1 recA1 hsdR17 (r_k^- m_k^+) deoR thi-1 supE44 gyrA96 relA1$	56
<i>S. aureus</i>		
8325-4	Wild-type strain cured of prophages	Lab stock
RN4220	Restriction-deficient transformation recipient	Lab stock
MJH001	<i>perR::kan</i>	This study
MJH002	<i>ahpC::pAZ106 ahpC<sup>+</sup></i>	This study
MJH003	<i>bcp::pAZ106 bcp<sup>+</sup></i>	This study
MJH004	<i>fn::pAZ106 fn<sup>+</sup></i>	This study
MJH005	<i>fur::pAZ106 fur<sup>+</sup></i>	This study
MJH006	<i>katA::pAZ106 katA<sup>+</sup></i>	This study
MJH007	<i>mrgA::pAZ106 mrgA<sup>+</sup></i>	This study
MJH008	<i>perR::pAZ106 perR<sup>+</sup></i>	This study
MJH009	<i>trxB::pAZ106 trxB<sup>+</sup></i>	This study
ST16	<i>katA::Tn917-LTV1</i>	This study
SPW1	<i>sodA::Tn917-LTV1</i>	65
MJH102	<i>perR::kan ahpC::pAZ106 ahpC<sup>+</sup></i>	This study
MJH103	<i>perR::kan bcp::pAZ106 bcp<sup>+</sup></i>	This study
MJH104	<i>perR::kan fn::pAZ106 fn<sup>+</sup></i>	This study
MJH105	<i>perR::kan fur::pAZ106 fur<sup>+</sup></i>	This study
MJH106	<i>perR::kan katA::pAZ106 katA<sup>+</sup></i>	This study
MJH107	<i>perR::kan mrgA::pAZ106 mrgA<sup>+</sup></i>	This study
MJH108	<i>perR::kan perR::pAZ106</i>	This study
MJH109	<i>perR::kan trxB::pAZ106 trxB<sup>+</sup></i>	This study
MJH408	<i>perR::kan pMAL34 (perR<sup>+</sup>)</i>	This study
MJH418	8325-4 pMAL34 ( <i>perR<sup>+</sup></i> )	This study
<b>Plasmids</b>		
pAZ106	Promoterless <i>lacZ erm</i> insertion vector	16
pGem3Zf(+)	General cloning vector	Promega
pAUL-A	Temperature-sensitive <i>erm</i> integrational shuttle vector	15
pCU1	<i>E. coli-S. aureus cat</i> shuttle vector	7
pMAL5	pGem3Zf(+) containing a 2-kb <i>perR</i> fragment with engineered <i>ClaI</i> site	This study
pMAL7	pMAL5 containing a <i>kan</i> cassette in <i>ClaI</i> site.	This study
pMAL8	3.5-kb <i>BamHI-HindIII perR::kan</i> fragment from pMAL7 in pAUL-A	This study
pMAL9	1.08-kb OL7–OL11 <i>perR</i> promoter-containing fragment in pAZ106	This study
pMAL10	0.6-kb OL13–OL14 <i>fur</i> promoter-containing fragment in pAZ106	This study
pMAL11	1.2-kb OL15–OL16 <i>katA</i> promoter-containing fragment in pAZ106	This study
pMAL12	1.2-kb OL17–OL18 <i>ahpCF</i> promoter-containing fragment in pAZ106	This study
pMAL13	1.1-kb OL19–OL20 <i>mrgA</i> promoter-containing fragment in pAZ106	This study
pMAL28	0.6-kb OL53–OL54 <i>bcp</i> promoter-containing fragment in pAZ106	This study
pMAL29	1.4-kb OL60–OL61 <i>fn</i> promoter-containing fragment in pAZ106	This study
pMAL30	1.4-kb OL62–OL63 <i>trxB</i> promoter-containing fragment in pAZ106	This study
pMAL34	0.9-kb OL82–OL83 <i>perR</i> gene-containing fragment in pCU1	This study
<b>Primers</b>		
OL7	AATTGGATCCCATGGTTTGCAACGGGTG	This study
OL8	CCGGAAGCTTATCCTGAGCCAGGATCAAACCTCTCCAT	This study
OL9	GAATCAATTGCATCGATTGCGACAAGCAGGCG	This study
OL10	CGCCTCCTTGTGCGCAATCGATGCAATTGATTC	This study
OL11	CCAGAATTTCGAATCGACTTGATGAGTCTCCATATG	This study
OL12	AATTGGATCCTGTAATGGTTTGCCACTTTGCAG	This study
OL13	CCAGAATTCAGTAGCTTCGCGTTGTGGCGTTAGC	This study
OL14	AATTGGATCCATTACCAAACGGGTGAAACGTC	This study
OL15	CCAGAATTCACGCAGCTTGTTCAACATCC	This study
OL16	AATTGGATCCGACCACAATGCCAATACAACC	This study
OL17	CCAGAATTCTGAATGTACCACGTTGAGCTAAC	This study
OL18	AATTGGATCCATGGTAAGCGTGGCTTGGCTGC	This study
OL19	CCAGAATTCATCATCGCCAGCATTACC	This study
OL20	AATTGGATCCATGGCAATTTCTGTGTCGCGAGG	This study
OL53	GGTTGGATCCAATTTCTAATTCAGTTCGGTGTACC	This study
OL54	CCAGAATTCATAAATTGTCTCTAAAGTCACAAGC	This study
OL60	AGAAGGATCCGCGTTATAAGCGTTAAAGTCAC	This study
OL61	AGCAGAATTCTGCATGTGCACCTCTGTCCG	This study
OL62	AGAAGGATCCAGAACTGATTACGATTGGTAG	This study
OL63	AGCAGAATTCCTGCCAAATACTTTCCGTCATC	This study
OL82	AGAGGATCCACAGCGCATATAACTGGTAATG	This study
OL83	CCAGAATTCCTTATACTCACTTTATGGATAG	This study
PEX2	AGCTCCATGGTCTGACGCTCAGTGGAAACAACTC	This study
PEX3	CAGTAGCTTCGCGTTGTGGCGTTAGC	This study
PEX4	GGTGTAATTCCTTACGCGCTGCTGTGTCGC	This study
PEX5	TCTCGGTCTGATACTGGATGCC	This study
PEX6	TTACAACATCTTGTGATTACTC	This study
Tn1	GGAACGCGGTCTACTTACAAGCAGC	This study
Tn2	CTCACAATAGAGAGATGTCACCGTC	This study

<sup>a</sup> In primer sequences, the restriction sites are underlined.



10% (wt/vol) polyacrylamide gel. Samples were blotted and N-terminally sequenced as described previously (16).

**RNA isolation and transcriptional start site mapping.** Total RNA was isolated from exponential cultures of *S. aureus* ( $OD_{600} = 1.0$ ) using the rapid liquid nitrogen chill method of Arnau et al. (6). Frozen cell pellets, briefly stored at  $-70^{\circ}\text{C}$ , were thawed, and RNA was rapidly extracted by cell disruption using the Fast-Prep blue kit (Bio 101) and a Fast-Prep system reciprocal shaker (Bio 101). Primer extension reactions were performed for *perR*, *katA*, *mrgA*, *ahpC*, and *fur* as described in the work of Horsburgh and Moir (36) using 100  $\mu\text{g}$  of RNA and 10 pmol of the corresponding primer (PEX2, PEX3, PEX4, PEX5, and PEX6, respectively). The PCR products used for cloning into pAZ106 were purified using a Qiagen PCR kit and then sequenced with a Sequenase 2.0 PCR kit (Amersham) using the same oligonucleotide used for the primer extension reaction.

**Virulence testing of strains in a murine skin abscess model.** Virulence of *S. aureus* strains was tested in an established murine abscess model of infection (16). Briefly, cells were grown to stationary phase in BHI (time, 15 h), then harvested by centrifugation, and washed twice in PBS. The cell numbers were adjusted to  $5 \times 10^8$  CFU  $\text{ml}^{-1}$ , and then 200  $\mu\text{l}$  of cell suspension was injected subcutaneously in female 6- to 8-week-old BALB/c mice. The precise inoculum was confirmed by serial dilution and counting on BHI agar. After 7 days, the mice were euthanized with  $\text{CO}_2$ , and skin lesions were aseptically removed and stored frozen in liquid nitrogen. The lesions were weighed, chopped, and homogenized in a mini-blender in 2.5 ml of ice-cold PBS. After 1 h of incubation on ice, the lesions were homogenized again before serial dilution of the suspension, and the total number of bacteria was determined by growth on BHI agar. Statistical significance was evaluated on the percent recovery of strains using Student's *t* test with a 5% confidence limit.

## RESULTS

**Identification of *perR* and *katA* in *S. aureus*.** The strain ST16 was isolated during a screen for *S. aureus* Tn917 transposants unable to survive glucose starvation after prolonged incubation on CDM agar plates. Genomic DNA sequencing in both directions from the transposon insertion of ST16 revealed inactivation of the *katA* gene encoding catalase. The 1,518-bp gene is monocistronic and encodes a protein of 505 amino acids with a molecular mass of 58.3 kDa (57). Upstream of the catalase gene is an imperfect palindrome of 18 bp, suggestive of transcriptional regulation by a member of the Fur family. A homologue of the *B. subtilis perR* gene, which controls peroxide stress resistance, was identified in the incomplete *S. aureus* 8325 genomic database (<http://www.genome.ou.edu>). The 441-bp *perR* gene is monocistronic and encodes a protein with a predicted molecular mass of 17.2 kDa. To characterize the role of PerR in *S. aureus*, we introduced a kanamycin resistance cassette into the *perR* gene using allelic replacement, thereby disrupting the chromosomal copy and creating strain MJH001.

**Peroxide resistance of MJH001 (*perR*) and ST16 (*katA*).** To investigate the importance of PerR and catalase in stress resistance of *S. aureus*, we examined the effect of adding hydrogen peroxide. MJH001 (*perR*) and ST16 (*katA*) mutants contrasted strikingly with 8325-4 (wild type). ST16 (*katA*) was hypersensitive to hydrogen peroxide, while MJH001 (*perR*) was more resistant than 8325-4 (wild type) (Fig. 1A). The *katA* gene encodes the sole catalase (Fig. 1B) of *S. aureus*, and the observed sensitivity was a consequence of a complete loss of all catalase activity as shown by enzyme assay (Fig. 1C). Database searching also failed to reveal any further catalase homologues in *S. aureus*. MJH001 (*perR*) was found to have increased levels of catalase activity (Fig. 1), as was reported elsewhere for *B. subtilis* (13) and *C. jejuni* (62) *perR* mutants. The addition of 20  $\mu\text{M}$  manganese to the growth medium reduced the amount of catalase present in 8325-4 (wild type) and decreased resistance

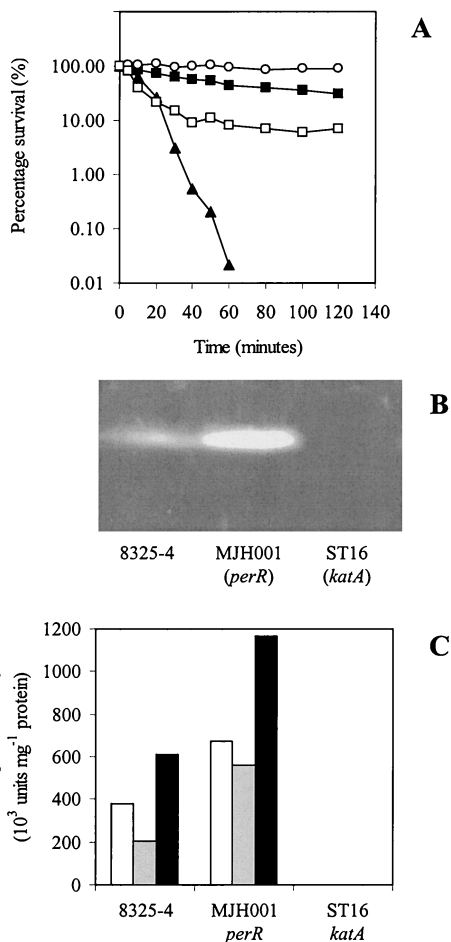


FIG. 1. (A) Effect of  $\text{H}_2\text{O}_2$  (7.5 mM) on washed, exponential-phase CLR-grown cells of 8325-4 (wild type) (■), MJH001 (*perR*) (○), and ST16 (*katA*) (▲) and 8325-4 (wild type) grown in CLR with 20  $\mu\text{M}$  manganese chloride (□). (B) Catalase activity gel. (C) Total catalase activity of washed, lysed, stationary-phase cells after 24 h of growth in CLR medium (open bars), CLR with 20  $\mu\text{M}$  manganese chloride (gray bars), or CLR with 20  $\mu\text{M}$  iron sulfate (black bars).

to hydrogen peroxide (Fig. 1A) but did not reduce the amount of catalase or resistance in MJH001 (*perR*). Addition of 20  $\mu\text{M}$  iron produced the reverse effect, increasing catalase expression in both 8325-4 (wild type) and MJH001 (*perR*). This contrasts with both *B. subtilis* and *C. jejuni*, where PerR functions as both a manganese- and an iron-responsive repressor (13, 62). The increased level of *katA* expression when MJH001 (*perR*) was grown with 20  $\mu\text{M}$  iron sulfate added is due to a second regulator of catalase that was identified as Fur (37).

**The role of *perR* and *katA* in the growth of *S. aureus*.** The importance of PerR and catalase for growth in vitro and their effect on the sensitivity of the strains to different metal ions were evaluated by growth in the metal-depleted medium CL. MJH001 (*perR*) had an increased doubling time compared to that of 8325-4 (wild type) during exponential growth in CL medium (262 and 144 min, respectively) (Fig. 2A). This difference in growth rate was eliminated (doubling time, 133 and 150 min, respectively) by adding increased (20  $\mu\text{M}$ ) iron sulfate (Fig. 2B) to the cultures. ST16 (*katA*) showed a wild-type

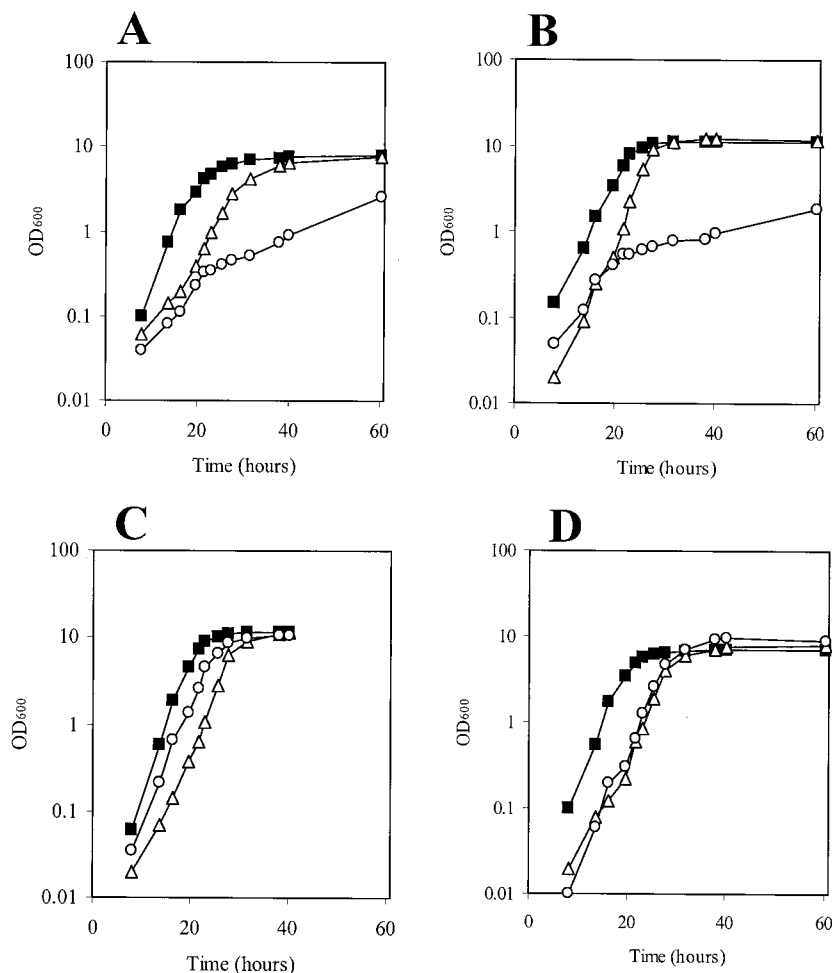


FIG. 2. Growth of 8325-4 (wild-type) (■), MJH001 (*perR*) (△), and ST16 (*katA*) (○) strains in CL with no added metal ions except magnesium (A), CL with 20  $\mu$ M iron sulfate (B), CLR (C), and CL with 20  $\mu$ M manganese chloride (D). All cultures were inoculated at an  $OD_{600}$  of 0.002.

growth rate in CLR and yet had marked growth defects in CL, where all metal ions (except magnesium) were absent from the culture (Fig. 2A), and in CL with 20  $\mu$ M iron sulfate (Fig. 2B). Growth in the metal-replete medium CLR produced a wild-type growth rate for ST16 (*katA*) (Fig. 2C). Further experiments showed that manganese alone was capable of restoring a wild-type growth rate to ST16 (*katA*) (Fig. 2D). Complementation of the mutation in MJH001 (*perR*) with the *perR* gene alone restored catalase activity and manganese-dependent repression to levels similar to those of the wild type (8325-4). This was demonstrated by assaying strains MJH408 (*perR::kan* [pMAL34 (*perR*<sup>+</sup>)]) and MJH418 (8325-4 [pMAL34 (*perR*<sup>+</sup>)]) during growth in CLR or CLR containing 20  $\mu$ M manganese chloride (data not shown).

**Identification of PerR-regulated genes in *S. aureus*.** Putative PerR binding sites were identified in the *S. aureus* incomplete genomic databases based on homology with the putative site upstream of *katA* (Fig. 3). A number of candidate sites were found upstream of the coding sequences of genes whose protein products were likely to have a role in oxidative stress or metal ion storage (Fig. 3). The *fur*, *fn*, *perR*, *trxB*, and *bcp* genes have not been identified as members of a PerR regulon

previously. To determine whether these candidate PerR binding sites were located in the promoter region of the genes, we determined the transcriptional start point for the *perR*, *fur*, *ahpC*, *katA*, and *mrgA* genes (Fig. 4). In each case, the PerR binding site was located in the promoter region close to the  $-35$  and  $-10$  elements. The PerR box in the *mrgA* gene was situated between the  $-35$  and  $-10$  elements in a position consistent with the high level of manganese-dependent transcriptional repression observed in 8325-4 (wild type) compared to that in MJH001 (*perR*) (Fig. 5).

**The effect of PerR on transcription of candidate genes.** To determine whether PerR regulated the expression of the identified genes, *lacZ* fusions were made to each of the eight genes listed in Fig. 3 to monitor transcription from their promoters. All of the identified genes displayed PerR-dependent regulation (Fig. 5) as evinced by significantly increased transcriptional activity from the promoters in MJH001 (*perR*) compared to that for 8325-4 (wild type). Furthermore, elevated levels of manganese repressed transcription of the genes, with the exception of *trxB*, in 8325-4 (wild type) (Fig. 5), but not in MJH001 (*perR*) (data not shown). This is consistent with the known manganese-responsive transcriptional repression activ-

AATTATAATTATTATAAAT  
 GATTAGAATTATTATAAAT  
 GATTAGAATTATTATAAAT  
 AATAATAATTATTATAAAT  
 TTTTTTAATTATTAGTAGG  
 AATTATAATTATTATTATT  
 GCATATAATTATTATTATT  
 TCATAAAATTATTATAAATG

- at TA t AATTATTAT a At -

TTA\*AAT\*ATTAT

*katA* Catalase  
*ahpCF* Alkyl hydroperoxide reductase  
*mrgA* Ferritin-like Dps  
*perR* Peroxide regulon regulator (PerR)  
*fur* Ferric uptake regulator (Fur)  
*ftn* Ferritin  
*trxB* Thioredoxin reductase  
*bcp, pdh* Bacterioferritin comigratory protein,  
 3-phosphoglycerate dehydrogenase

*S. aureus* consensus sequence

*B. subtilis* consensus sequence (19)

FIG. 3. Alignment of the putative PerR boxes identified in the incomplete *S. aureus* genome databases. Boxes were identified as described previously (37). An *S. aureus* consensus sequence was compiled from all of the sequences identified and compared to the *B. subtilis* consensus (19).

ity previously reported for PerR from *B. subtilis* (18). The addition of calcium, copper, cobalt, nickel, molybdenum, or zinc to CLR had no significant effect on transcription of *katA* or *mrgA* (data not shown). Transcription of all of the PerR-regulated genes was found to be consistently greater in CLR with 20  $\mu$ M iron sulfate added than in CLR medium alone (Fig. 5). Titration of the iron-dependent induction showed concentrations above 2  $\mu$ M to be effective (data not shown). In the case of *katA* and *ftn*, iron-dependent induction was found to be as great as the level of transcription resulting from *perR* inactivation (Fig. 5). Transcription of *katA* is positively regulated, either directly or indirectly, by Fur in an iron-dependent manner (37).

SOD activity in *S. aureus* was not affected by *perR* inactivation (data not shown). Interestingly, when catalase levels were assayed in a *sodA* mutant (SPW1) of *S. aureus*, we found a higher specific activity (698 U mg<sup>-1</sup>) than that for 8325-4 (wild type) (379 U mg<sup>-1</sup>). Since this suggested that increased superoxide stress in the cell ultimately leads to activation of the PerR regulon, we tested the effect of paraquat on induction of *mrgA*. The addition of 10  $\mu$ M paraquat to an early- or mid-log culture of MJH007 (*mrgA-lacZ mrgA*<sup>+</sup>) produced a threefold increase in  $\beta$ -galactosidase activity when assayed 60 min after paraquat addition (data not shown).

#### Induction of PerR-regulated genes with hydrogen peroxide.

PerR was originally described as a peroxide-responsive repressor and was shown to derepress transcription of the *katA*, *ahpC*, *hemA*, and *mrgA* genes of *B. subtilis* after the addition of 100  $\mu$ M hydrogen peroxide (19). Peroxide induction of the *S. aureus* PerR regulon *lacZ* reporter fusions was tested in 8325-4 (wild type) by adding 100 or 500  $\mu$ M hydrogen peroxide to early- and mid-exponential and early-stationary-phase cultures grown in CLR medium or Chelex-treated BHI. A 100  $\mu$ M concentration of hydrogen peroxide was found to produce little induction compared to a 500  $\mu$ M concentration, possibly due to the very high levels of catalase activity present in *S. aureus*. Using a concentration of 500  $\mu$ M hydrogen, increased transcription was observed in early- and mid-log-phase cultures grown in CLR for *ahpC* (threefold), *bcp* (twofold), *ftn* (threefold), *katA* (twofold), *mrgA* (sixfold), and *trxB* (twofold) but

not for *fur* or *perR* (data not shown). Induction was not observed in early-stationary-phase growth.

**The effect of PerR and KatA on cellular protein profiles.** The soluble cellular protein fraction of the strains revealed that MJH001 (*perR*) overexpressed a number of proteins compared to 8325-4 (wild type) (Fig. 6). The addition of manganese repressed these proteins in 8325-4 (wild type), while iron induced their expression; neither metal affected expression of these proteins in MJH001 (*perR*) (data not shown). To confirm that the major proteins were members of the PerR regulon, they were N-terminally sequenced. The N-terminal sequence of the 17-kDa (SNQQDVVKEXNQXANXTVA), 20-kDa (LSKNXLEAL), and 23-kDa (SLINKEILPF) proteins matched the predicted translation of the *mrgA*, *ftn*, and *ahpC* genes of *S. aureus*, respectively. AhpC was one of four major proteins from *S. aureus* induced by osmotic upshock after growth in 2.5 M NaCl (5). Apart from AhpC, the other proteins induced during osmotic upshock do not match the sizes of those overexpressed in MJH001 (*perR*), suggesting that this osmotic induction was not PerR mediated. ST16 (*katA*) was found to overexpress the same proteins as MJH001 (*perR*), albeit at a lower level (data not shown). Bsai et al. (14) and Antelmann et al. (3) have shown that inactivation of *katA* and *ahpC* in *B. subtilis* results in derepression of the PerR regulon. From this, it was inferred that PerR senses the accumulation of both hydrogen peroxide and organic hydroperoxide.

Analysis of stationary-phase exoprotein profiles showed no apparent difference between MJH001 (*perR*) and 8325-4 (wild type). This was confirmed for specific proteins using antisera raised against alpha-hemolysin (Hla) and staphylococcal serine protease A (SspA). In addition, the zones of hemolysis on rabbit blood agar were identical for MJH001 (*perR*) and 8325-4 (wild type) (data not shown).

**The role of PerR and catalase in starvation survival.** ST16 (*katA*) was identified in a carbon starvation screen due to its inability to survive extended incubation on glucose-limiting CDM. Catalase is induced postexponentially, and starved *S. aureus* cells were shown previously to be highly resistant to elevated hydrogen peroxide (64). The reduced viability of ST16 (*katA*) was demonstrated by prolonged aerobic incubation at

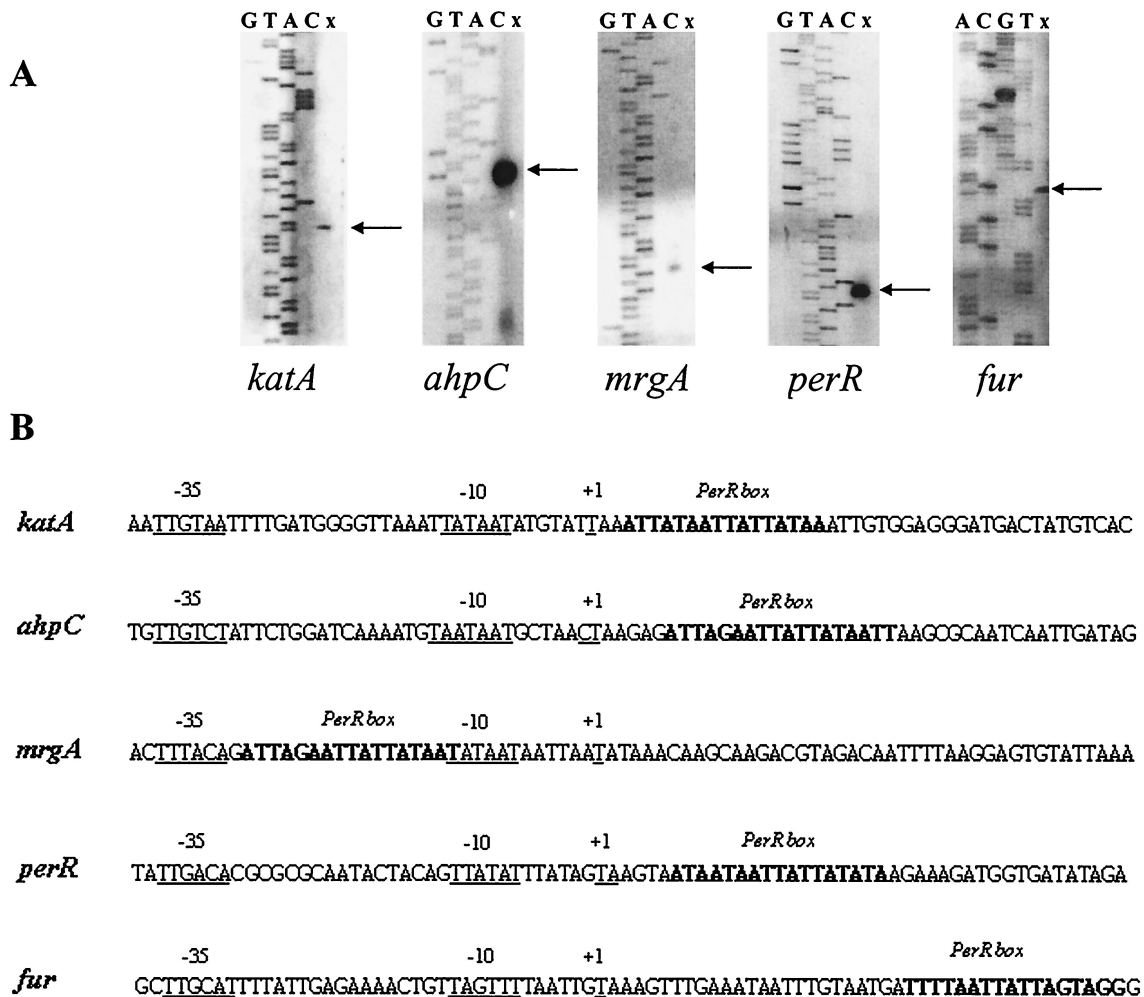


FIG. 4. (A) Mapping of the 5' end of the *perR*, *katA*, *mrgA*, *ahpC*, and *fur* mRNA by primer extension. RNA was isolated from exponential cultures of 8325-4 (wild type) grown in CLR ( $OD_{600}$  of 1.0), and 100  $\mu$ g of RNA was used for each reaction. The potential transcription start sites are marked with arrows. Lanes G, T, A, and C show the dideoxy sequencing ladder obtained with the same primer used for primer extension (x). (B) Sequence of the identified promoter regions. Potential  $-35$  and  $-10$  regions and transcriptional start sites are underlined. PerR boxes are in boldface.

37°C (Fig. 7). In contrast, MJH001 (*perR*) retained a wild-type level of viability in this assay.

**The importance of PerR and catalase in vivo.** Catalase was previously proposed to be an important virulence determinant in *S. aureus* (40, 46) since catalase and SOD levels correlated with virulence in clinical isolates. The availability of single defined mutations in *sodA* (21) and now *katA* has enabled us to test this fully. While *sodA* encodes one of two SODs, making the importance of SOD activity in virulence harder to test, *katA* encodes the sole catalase of *S. aureus*. Together with MJH001 (*perR*) and 8325-4 (wild type), we tested the pathogenicity of ST16 (*katA*) in a murine subcutaneous skin abscess model of infection (Fig. 8). ST16 (*katA*) was not attenuated in this model ( $P = 0.124$ ) in two independent experiments ( $n = 7$  and  $n = 9$ ). Conversely, MJH001 (*perR*) was significantly attenuated in this model of infection ( $P < 0.005$ ) and produced smaller lesions ( $0.237 \pm 0.143$  g) compared to both 8325-4 ( $0.528 \pm 0.173$  g) and ST16 ( $0.582 \pm 0.217$  g).

## DISCUSSION

*S. aureus* PerR controls a large regulon of oxidative stress components including catalase, alkyl hydroperoxide reductase, the thioredoxin-dependent thiol peroxidase (Bcp), and thioredoxin reductase. It also regulates the iron storage protein ferritin and the ferritin-like Dps homologue, MrgA. Furthermore, *S. aureus* PerR has an autoregulatory capacity and modulates expression of the iron homeostasis regulator Fur. In *B. subtilis*, PerR has so far been shown to have a more specific role as a controller of catalase, alkyl hydroperoxide reductase, MrgA, and heme biosynthesis enzymes.

MJH001 (*perR*) has a reduced growth rate in CLR medium that contains low levels of iron and manganese. Supplementing the medium with iron negated this defect, and we hypothesize that the observed high level of overexpression of ferritin and MrgA may sequester intracellular iron, effectively limiting free iron in the cell. MrgA in *B. subtilis* functions as a Dps-like



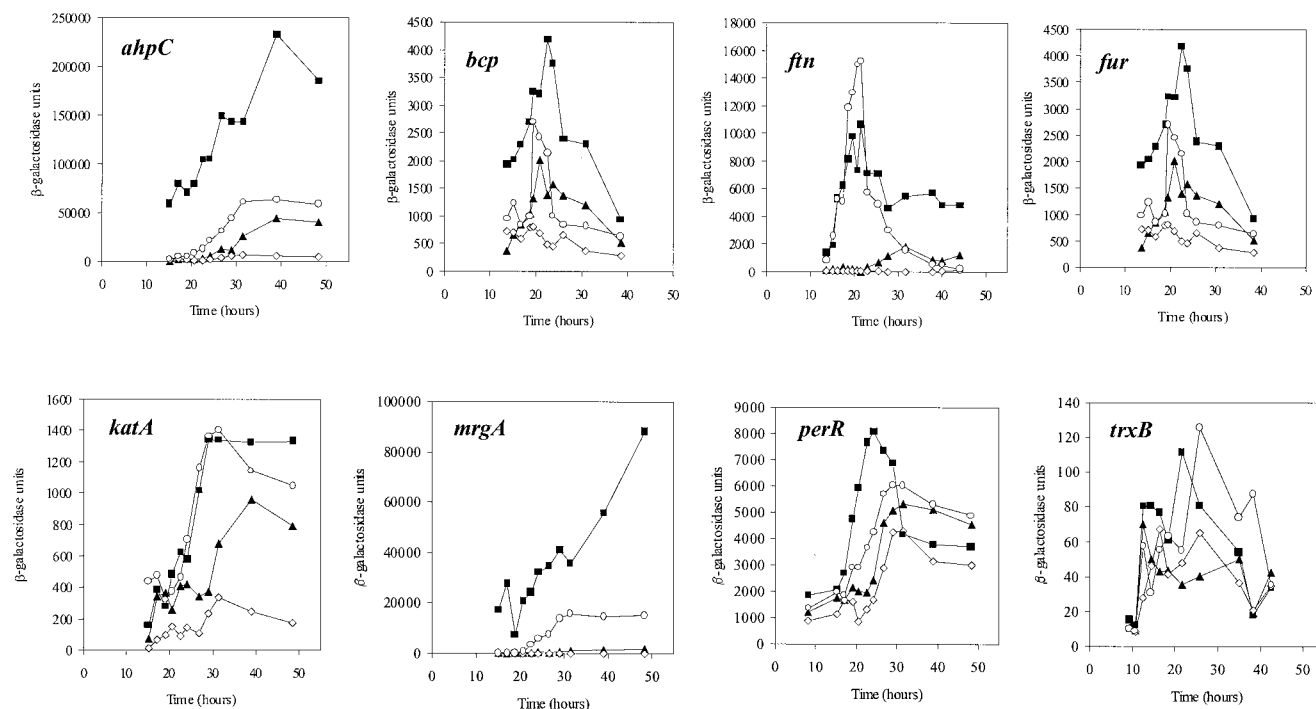


FIG. 5. Analysis of transcription from promoter-*lacZ* fusions during growth in CLR medium. 8325-4 (wild type) containing the fusion indicated was grown in CLR medium ( $\blacktriangle$ ), CLR with 20  $\mu\text{M}$  manganese ( $\diamond$ ), and CLR with 20  $\mu\text{M}$  iron sulfate ( $\circ$ ). Transcription of the *lacZ* fusions in MJH001 (*perR*) ( $\blacksquare$ ) is shown on each graph. Samples were removed at the times indicated and sampled for  $\beta$ -galactosidase activity. The strains grew with slightly different growth rates as indicated in Fig. 2, but for reasons of clarity only the changes in  $\beta$ -galactosidase activity are shown. The *lacZ* fusions shown are MJH002 (*ahpC-lacZ*), MJH003 (*bcp-lacZ*), MJH004 (*ftn-lacZ*), MJH005 (*fur-lacZ*), MJH006 (*katA-lacZ*), MJH007 (*mrgA-lacZ*), MJH008 (*perR-lacZ*), and MJH009 (*trxB-lacZ*), and these same fusions in the *perR* mutant background are MJH102, MJH103, MJH104, MJH105, MJH106, MJH107, MJH108, and MJH109, respectively.

protein. These proteins have the ability to bind DNA (17, 47) and sequester iron to protect the DNA from oxidative damage (29, 67), and *B. subtilis* MrgA is important for cellular defense against hydrogen peroxide (17).

The indirect control of iron homeostasis by PerR regulation of Fur and the direct control of iron storage proteins allow *S. aureus* to coordinate the intracellular availability of free iron with the level of antioxidant proteins present in the cell. The deleterious effect of hydroxyl radicals on growth resulting from hydrogen peroxide and Fe(II) participating in the Fenton reaction has been demonstrated clearly in *S. aureus* (54), and elevated iron levels have been shown to predispose *S. aureus* to killing by monocytes and macrophages but not polymorphonuclear granulocytes (35, 55). Regulation of *fur* in *E. coli* is mediated by a number of regulators including OxyR, the well-characterized, gram-negative bacterial functional analogue of PerR (60).

Expression of the PerR regulon is repressed by elevated manganese concentrations, and this is likely to be due to the antioxidant properties of manganese in vivo. Numerous studies have shown that several manganese complexes can catalyze the disproportionation of hydrogen peroxide (11, 59) while others can act as superoxide scavengers (27). *Lactobacillus plantarum* maintains high intracellular levels of manganese as a substitute for having enzymes that protect against reactive oxygen species (4). Que and Helmann (53) recently described the function of MntR of *B. subtilis* as a manganese homeostasis regulator

controlling expression of two manganese uptake transporter systems, MntABC and MntH (Nramp); it has a similar role in *S. aureus* (S. J. Wharton, M. J. Horsburgh, and S. J. Foster, unpublished data). SirR from *S. epidermidis* is a homologue of MntR, although its role in metal ion homeostasis is not known (34). TroR was recently described as a DtxR-like manganese-dependent regulatory protein that regulates a manganese transport system in *Treponema pallidum* (52). A direct regulatory link between the PerR and MntR regulons has not been reported, although Kehres et al. (42) have reported that MntH in *S. enterica* serovar Typhimurium is induced by peroxide and has a putative OxyR binding site. We have shown that there is PerR-dependent control of Fur, the regulator of iron homeostasis in *S. aureus* (37). Should such a link between PerR and the MntR regulon exist, it would demonstrate a complex interdependence between antioxidant defense and the levels of iron and manganese in the cell. In *S. aureus*, the importance of PerR as a central regulator of antioxidant defenses and of iron storage proteins was shown by the reduced virulence of a *perR* mutant in a skin abscess model of infection. Measured levels of manganese in the human body vary from extremely low levels in the skin (0.05  $\mu\text{M}$ ) to 30-fold-higher levels in the blood (1.5 to 2.4  $\mu\text{M}$ ) and yet-higher levels in the central nervous system (160 to 180  $\mu\text{M}$ ) (41, 52). *S. aureus* will encounter varying concentrations of manganese depending on its location during infection, and these encompass the concentrations over which PerR is active, since titration experiments show that in vitro



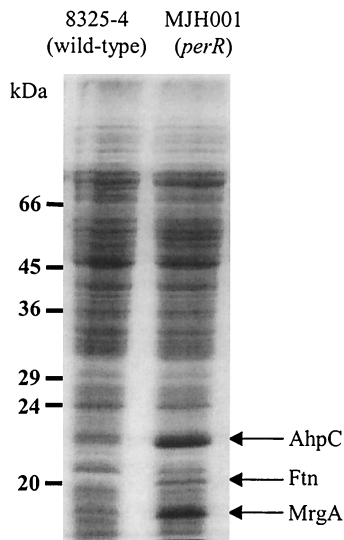


FIG. 6. Sodium dodecyl sulfate-polyacrylamide gel electrophoresis of total proteins from wild-type and MJH001 (*perR*) strains after growth to stationary phase in CLR at 37°C. The AhpC, Ftn, and MrgA proteins identified by N-terminal sequencing are shown.

manganese-dependent PerR repression occurs at and above 1 to 2  $\mu$ M concentrations (data not shown).

In *B. subtilis*, PerR was identified as a metal limitation and peroxide-sensing repressor of the genes encoding catalase, MrgA, AhpC, and heme biosynthesis enzymes (13, 17, 18, 19). The *S. aureus* PerR regulon was shown here to be metal regulated; however, not all of the genes were induced by hydrogen peroxide. Induction was observed only at early and mid-log phase and not the stationary phase of growth. The lack of induction of transcription for *perR* and *fur* suggests that some promoters may be noninducible with hydrogen peroxide, a feature that would indicate a complex response to peroxide stress for members of the PerR regulon. Transcription of the *perR* gene, for example, is autoregulated. The complex regulation of some of the PerR-dependent loci is highlighted by the effect of iron. In some cases, such as *kataA*, the level of transcription after addition of iron was as high as that in a *perR* mutant background. We have previously demonstrated that *kataA* is positively regulated by Fur in an iron-dependent manner (37).

The intracellular redox-regulating molecules thioredoxin, glutaredoxin, and protein disulfide isomerase maintain cellular redox status and catalyze the formation and reduction of disulfide bonds. PerR-dependent regulation of thioredoxin reductase (TrxB) and a thiol peroxidase (Bcp) further demonstrates the central role of *S. aureus* PerR in protection from reactive oxygen species. The thiol peroxidase (Bcp) recently characterized by Jeong et al. (39) was shown to reduce linoleic acid hydroperoxide and hydrogen peroxide with the use of thioredoxin as an in vivo immediate electron donor. We note that a second thiol peroxidase encoded in the *S. aureus* 8325 genome has a putative PerR box (AAGTATTATTATTATTATTATT) with a high level of identity to those in this study and is located in the likely promoter region of this gene. It would thus appear that *S. aureus* PerR, like OxyR of *E. coli* (2), has a regulatory role in transcription of genes for the thioredoxin

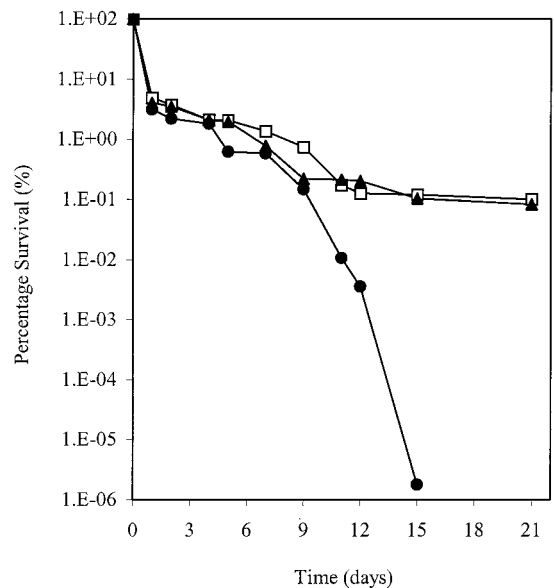


FIG. 7. Starvation survival of 8325-4 (wild type) ( $\square$ ), MJH001 (*perR*) ( $\blacktriangle$ ), and ST16 (*kataA*) ( $\bullet$ ) after growth and incubation in CDM containing 0.1% (wt/vol) glucose (glucose limiting). Samples were aseptically removed at the times indicated, and viability was assessed by dilution and counting on BHI agar.

pathway and thiol peroxidases. The gene encoding 3-phosphoglycerate dehydrogenase (*pdh*) is located in the same operon as *bcp*, such that it will be under PerR control. The reason for this is unclear but may be a metabolic mechanism for regenerating

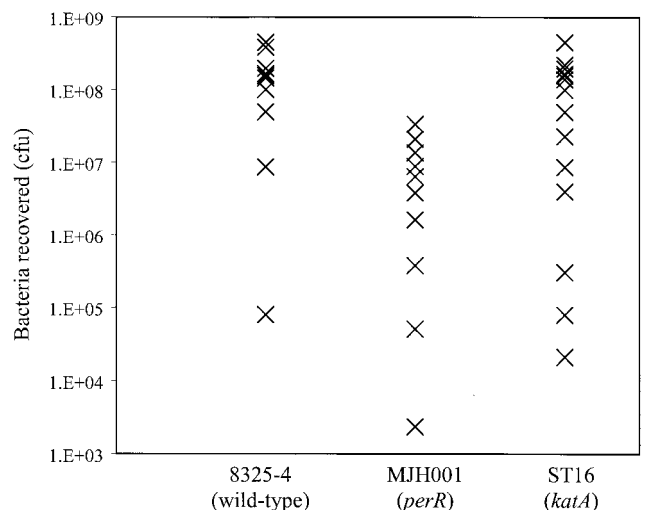


FIG. 8. Pathogenicity of *S. aureus* strains in a murine skin abscess model of infection. Approximately  $10^8$  CFU of each strain was inoculated subcutaneously into 6- to 8-week-old BALB/c mice; the strains used were 8325-4 (wild type) ( $n = 10$ ), MJH001 (*perR*) ( $n = 10$ ), and ST16 (*kataA*) ( $n = 16$ ). Seven days after infection, mice were euthanized, lesions were removed and homogenized, and viable bacteria were counted after dilution and growth on BHI agar plates. The mean percent recovery of each strain and the Student *t* test *P* values are as follows, respectively: 8325-4 (wild type), 143%; MJH001 (*perR*), 10.5% and  $P < 0.005$ ; and ST16 (*kataA*), 86.3% and  $P = 0.124$ .

the reduced NAD that is consumed when a cell is subjected to oxidative stress.

The ability of *S. aureus* to survive intracellularly in granulocytes has been noted in several studies (25, 49, 70), and recently this was shown to contribute to infection (30). The intracellular location of the staphylococci was shown to be in macropinosome-like vacuoles similar to those seen with salmonellae in epithelial cells and macrophages. *S. aureus* has also been shown to escape the endosome of epithelial cells (8). It is not yet clear what role oxidative stress resistance has in these intracellular life cycles. However, the ability of PerR to sense hydrogen peroxide and to regulate antioxidant defense and iron storage may be important for coordinating a survival response. The mechanism by which PerR senses hydrogen peroxide has not been determined, although it has been proposed that metal ion oxidation may be involved (13). We are currently characterizing PerR from *S. aureus* to determine how oxidative stress is perceived by this protein and how this signal is transduced to DNA binding.

Catalase has long been implicated as a virulence determinant in *S. aureus* (40, 46). We have shown definitively that it has no role in the murine skin abscess model of infection. This, together with our previous work with SodA (21), suggests that these determinants are not important for infection, at least not in the well-characterized skin abscess model that we have studied. We note that the previous studies used undefined clinical isolates and that the levels of both catalase and SOD were reduced in their avirulent isolates. We have not observed any correlation between disruption of *katA* and *sodA* levels; in fact, disruption of *sodA* increased catalase levels. *Staphylococcus aureus* subsp. *anaerobius* is an organism closely related to *S. aureus* and is the etiological agent of lymphadenitis and abscess formation in young sheep and goats (57). The main phenotypic differences between the strains are the lack of catalase and weak aerobic growth in the former (22). Weak aerobic growth was not observed in *S. aureus* ST16 (*katA*), and we suggest that the altered characteristics of *S. aureus* subsp. *anaerobius* are not due solely to its dysfunctional catalase gene (57).

The ability to survive long-term starvation and desiccation has been implicated as an important factor in the nosocomial transmission of *S. aureus* and is likely to exacerbate the problem of methicillin-resistant *S. aureus* (23). We have previously identified a number of starvation survival components required for nutrient recycling, cellular repair, and oxidative stress resistance (65). The identification of catalase as a critical component for maintaining viability in long-term starvation reinforces the importance of protection from reactive oxygen species in this process.

This study has identified oxidative stress resistance and/or metal ion homeostasis to be important in the ability of *S. aureus* to cause disease. The role of the many PerR-regulated genes and the complex coregulatory processes linking stress resistance, metal ion homeostasis, and pathogenicity await investigation.

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