



Superoxide dismutase activity confers (p)ppGpp-mediated antibiotic tolerance to stationary-phase *Pseudomonas aeruginosa*

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Metabolically quiescent bacteria represent a large proportion of those in natural and host environments, and they are often refractory to antibiotic treatment. Such drug tolerance is also observed in the laboratory during stationary phase, when bacteria face stress and starvation-induced growth arrest. Tolerance requires (p)ppGpp signaling, which mediates the stress and starvation stringent response (SR), but the downstream effectors that confer tolerance are unclear. We previously demonstrated that the SR is linked to increased antioxidant defenses in *Pseudomonas aeruginosa*. We now demonstrate that superoxide dismutase (SOD) activity is a key factor in SR-mediated multidrug tolerance in stationary-phase *P. aeruginosa*. Inactivation of the SR leads to loss of SOD activity and decreased multidrug tolerance during stationary phase. Genetic or chemical complementation of SOD activity of the $\Delta reLA spoT$ mutant (ΔSR) is sufficient to restore antibiotic tolerance to WT levels. Remarkably, we observe high membrane permeability and increased drug internalization upon ablation of SOD activity. Combined, our results highlight an unprecedented mode of SR-mediated multidrug tolerance in stationary-phase *P. aeruginosa* and suggest that inhibition of SOD activity may potentiate current antibiotics.

(p)ppGpp stringent response | *Pseudomonas aeruginosa* | antibiotic tolerance | superoxide dismutase | stationary phase

Treatment of chronic bacterial infections in the clinic often results in failure or relapses. A drug refractory state, commonly referred to as antibiotic tolerance, occurs even when the infecting organisms harbor no genotypic (heritable) resistance to the antibiotic, and renders many chronic infections difficult to eradicate (1–3). Medically important examples include chronic *Pseudomonas aeruginosa* lung infections in individuals with the genetic disease cystic fibrosis. Tolerance can also promote the emergence of genotypic drug resistance, thereby posing a major public health challenge (4). Bacteria develop drug tolerance during growth-limiting conditions when they adopt a slow or nonreplicating state, and a fraction of the population survives bactericidal drugs (5). In fact, a large proportion of microbes found in natural environments and in vivo during chronic human infections are likely metabolically quiescent (2, 6).

Laboratory stationary-phase bacteria provide a valuable window into the metabolically quiescent organisms widely observed in nature. The physiology of exponentially growing bacteria change remarkably as they enter stationary phase, yet little is known about the survival strategies of slow or nongrowing cells (7, 8). Stationary-phase bacteria must respond and adapt to a variety of growth-limiting stress and starvation cues (e.g., nutrient exhaustion, pH changes, oxidative or nitrosative stress) through processes regulated by the alternative σ -factor RpoS and (p)ppGpp signaling in *Escherichia coli* and *P. aeruginosa* (9, 10). The alarmone (p)ppGpp accumulates upon stress and starvation, leading to a global reorganization of cellular and metabolic

functions that promote stress adaptation and cell survival, a process termed the stringent response (SR) (11, 12).

Antibiotic tolerance among metabolically quiescent bacteria is widely attributed to the notion that drug targets are unavailable or inactive when cellular replication and macromolecule synthesis are shut down. Although antibiotic killing typically correlates with bacterial growth rate (13, 14), the lack of replication alone in the absence of (p)ppGpp signaling and downstream adaptive responses is often insufficient to confer tolerance (15–17). The downstream cellular processes that protect against antibiotic toxicity remain poorly understood. We previously observed that SR inactivation in the (p)ppGpp-null $\Delta reLA spoT$ mutant of *P. aeruginosa* (ΔSR) impairs multidrug tolerance in nutrient-limited, biofilm and stationary-phase bacteria (16, 18). Notably, the ΔSR mutant exhibited impaired superoxide dismutase (SOD) and catalase activities, leading us to propose that SR-mediated multidrug tolerance is linked to enhanced antioxidant defenses (16, 18).

Superoxide radicals are by-products of aerobic metabolism and a primary source of intracellular oxidative stress (19). Superoxide causes toxicity through direct damage of iron-containing enzymes, and indirectly through highly reactive hydroxyl radicals generated by Fenton chemistry (20). SODs rapidly disproportionate superoxide to oxygen and hydrogen peroxide, and the latter is detoxified by catalases and peroxidases. *P. aeruginosa* encodes two different SODs, SodA and SodB. The Fe-cofactored

Significance

Antibiotic tolerance causes antibiotic treatment failure and promotes the emergence of genotypic resistance in chronic infections, such as those caused by the pathogen *Pseudomonas aeruginosa*. Laboratory stationary-phase bacteria exhibit a slow growing and metabolically quiescent state associated with high levels of multidrug tolerance likely analogous to the in vivo environment during chronic infection. We demonstrate that superoxide dismutases confer multidrug tolerance in stationary-phase bacteria, and identify a link between (p)ppGpp-mediated stress responses, superoxide metabolism, and membrane permeability to antibiotics. Inhibition of superoxide dismutase activity may overcome multidrug tolerance and potentiate current bactericidal antibiotics in the treatment of *P. aeruginosa* chronic infections.

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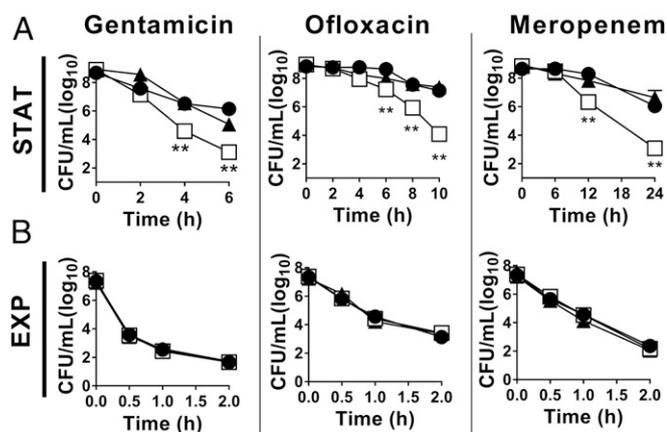


Fig. 1. SR inactivation impairs multidrug tolerance in stationary phase *P. aeruginosa*. (A) Stationary phase (STAT) or (B) exponential phase (EXP) cells of (●) WT, (□) ΔSR, and (▲) +SR challenged with 50 μg/mL gentamicin, 5 μg/mL ofloxacin, or 300 μg/mL meropenem in antibiotic killing assays. Note the different time scale in A and B. Results are mean ± SEM ($n = 6$). ** $P < 0.01$ vs. WT.

SodB is the most abundant in iron-replete conditions, while the Mn-cofactored SodA is under iron-dependent repression and only expressed under iron limitation (21, 22). In this study, we demonstrate that SOD activity is a critical effector of SR-mediated multidrug tolerance in stationary-phase *P. aeruginosa*, and that SOD activity is correlated with membrane permeability and drug internalization. Our data demonstrate a link between antioxidant defenses, drug permeability, and SR-mediated drug tolerance when *P. aeruginosa* are metabolically quiescent.

Materials and Methods

Experimental details can be found in the *SI Appendix*.

Bacterial Strains and Plasmids. All strains and plasmids used are listed in *SI Appendix*, Tables S1 and S2. The *P. aeruginosa* laboratory strain PAO1 is the parental WT strain. The (p)ppGpp-null isogenic ΔSR mutant carries unmarked deletions of both (p)ppGpp synthetases *relA* and *spoT*, and the +SR strain is the ΔSR mutant complemented for the *relA* and *spoT* genes (16).

Media and Growth Conditions. Bacteria were grown in LB Miller medium as described in *SI Appendix*. Exponential phase cells were grown for ~2 h to an $OD_{600} = 0.2$, and stationary-phase cells for 16 h to an $OD_{600} \approx 3.5$.

SOD Activity Assays. Total SOD and Sod-specific activities of cell lysates were measured using an in-solution and an in-gel activity assay respectively as described previously (23).

Antibiotic Killing Assays. Exponential or stationary-phase cultures were challenged with antibiotics without addition of fresh medium, and incubated in 96-well plates at 37 °C with shaking at 250 rpm. At specific time points, cells were mixed with 1:1 activated charcoal to bind free drug, and viable bacteria were measured by serial microdilution and overnight growth of CFU on LB agar plates.

Results

(p)ppGpp Signaling Is Required for the Multidrug Tolerance of Stationary-Phase *P. aeruginosa*. Stationary-phase bacteria are highly drug tolerant compared with their exponentially growing counterparts. To examine the contribution of (p)ppGpp signaling in *P. aeruginosa* stationary-phase drug tolerance, we challenged the (p)ppGpp-null ΔSR mutant to multiple distinct classes of bactericidal antibiotics and compared it to its WT isogenic parental strain. Stationary-phase ΔSR mutant cells are highly impaired for tolerance compared with WT (Fig. 1A), with 3–4 log₁₀ greater antibiotic killing by the aminoglycoside gentamicin (5.9- vs. 2.9-log₁₀ killing of ΔSR vs. WT, respectively, at $t = 6$ h), the fluoroquinolone ofloxacin (4.9- vs. 1.9-log₁₀ killing at $t =$

10 h), and the β-lactam meropenem (6.0- vs. 2.9-log₁₀ killing at $t = 24$ h). In contrast, exponential phase WT and ΔSR bacteria are equally susceptible to all three drugs and undergo rapid killing (Fig. 1B). Multidrug tolerance is fully restored to WT levels upon complementation of the ΔSR mutant with the *relA* and *spoT* genes (+SR), confirming that the loss of tolerance is attributable to *relA* and *spoT* mutations. Notably, the bacterial viability in stationary phase and growth rate in exponential phase are similar between the WT, ΔSR, and +SR strains (*SI Appendix*, Fig. S1A and B). Deletion of *relA* and *spoT* in two additional *P. aeruginosa* clinical strains to abrogate (p)ppGpp synthesis also results in loss of stationary-phase multidrug tolerance, although the magnitude of this effect differs in different *P. aeruginosa* genetic backgrounds (*SI Appendix*, Fig. S2A).

SOD Activity Is Induced During Stationary Phase and Requires (p)ppGpp Signaling. Our recent observation that ΔSR mutant biofilms exhibit low SOD activity led us to examine whether SOD activity requires (p)ppGpp signaling in planktonic stationary-phase bacteria. We first note a fourfold increase in SOD activity upon transition of the WT from exponential to stationary phase. However, this induction is largely abrogated in the ΔSR mutant and the SOD activity in stationary-phase ΔSR cells is reduced to 35% of WT levels (40 vs. 14.5 U/mg, $P < 0.01$) (Fig. 2A). We also observed similar effects upon inactivation of the *relA* and *spoT* genes in two *P. aeruginosa* clinical isolates (*SI Appendix*, Fig. S1B). Notably, differences in SOD activity are not due to disparities in bacterial growth rate, viability, or total cellular protein content (*SI Appendix*, Fig. S1B and C). In addition, complementation of the ΔSR mutant with the *relA* and *spoT* genes restores SOD activity to WT levels, and confirms that the SOD defect is attributable to the *relA* and *spoT* mutations. These observations thus indicate that (p)ppGpp signaling is required for full SOD activity during stationary phase.

SodB Is the Dominant SOD in Stationary Phase and Is Regulated by the SR and RpoS. We showed by in-gel SOD activity assay that SodB confers all SOD activity in stationary-phase WT, ΔSR, and +SR cells, while SodA activity is undetectable under these conditions (Fig. 2B). Mirroring SodB activity, *sodB* expression, as measured by a *sodB-lacZ* transcriptional reporter, is induced in WT cells but not in the ΔSR mutant once cells enter stationary phase ($t = 8$ h). For example, *sodB* expression is 2.2- to 2.8-fold lower in stationary phase ΔSR compared with WT cells (Fig. 2C) in the absence of any differences in growth rate (*SI Appendix*, Fig. S1D).

The SR and the alternative σ-factor RpoS regulatory networks significantly overlap to control gene expression during stationary

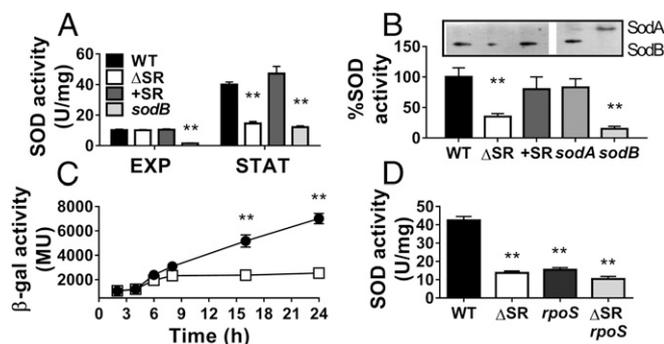


Fig. 2. SOD activity and *sodB* expression are induced during stationary phase in the WT but not the ΔSR mutant. (A) Total SOD activity of exponential (EXP) or stationary phase (STAT) WT, ΔSR, +SR, and *sodB* strains. (B) SodA and SodB specific activities in stationary phase cells measured by in-gel SOD activity assays, with a representative gel image shown. SodB activity in the *sodA* mutant and SodA activity in the *sodB* mutant were confirmed as control. (C) *sodB-lacZ* reporter activities in the WT (●) or ΔSR (□) cells during growth in LB medium. (D) Total SOD activity in stationary-phase WT, ΔSR, *rpoS*, and ΔSR *rpoS* strains measured by in-solution assay. All results are shown as mean ± SEM ($n \geq 5$). ** $P < 0.01$ vs. WT in the corresponding growth phase.

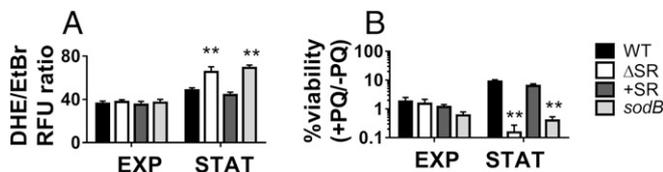


Fig. 3. Inactivation of the SR is associated with increased superoxide levels and paraquat killing. (A) Relative intracellular superoxide levels using the DHE/EtBr fluorescence ratio. (B) Paraquat (PQ) killing of exponential (EXP) or stationary-phase (STAT) cells, calculated as percent bacterial survival after 6-h challenge with 10 mM PQ, compared with similar conditions without PQ. Results are shown as mean \pm SEM ($n \geq 6$). ** $P < 0.01$ vs. WT.

phase (24), and here we show that SodB activity is under both SR and RpoS control. The Δ SR and *rpoS* mutants show equally reduced SOD activity compared with WT (~ 15 vs. 42 U/mg, $P < 0.01$) (Fig. 2D). Furthermore, the modest reduction in SOD activity in the Δ SR *rpoS* triple mutant compared with the *rpoS* mutant (10 vs. 14 U/mg, respectively, $P < 0.05$) further supports the overlapping SR- and RpoS-dependent regulation of SodB activity, and presumably *sodB* expression.

Stationary-Phase Δ SR Cells Have Higher Superoxide Levels and Lower Paraquat Tolerance than WT Cells. Because of reduced SOD activity, we predicted that Δ SR cells would be impaired in superoxide detoxification, leading to elevated superoxide levels and greater susceptibility to paraquat, a superoxide-generating compound. We therefore monitored relative intracellular superoxide levels using dihydroethidium (DHE), a cell-permeable probe that, when oxidized by superoxide, is converted to the fluorescent product 2-hydroxyethidium (25). DHE fluorescence intensity is ~ 2.5 -fold higher in the Δ SR and *sodB* mutants compared with WT and +SR strains in stationary, but not exponential phase (SI Appendix, Fig. S3 A and B). To account for potential differences in DHE probe loading, we used structurally similar ethidium bromide (EtBr) as a loading control and calculated DHE/EtBr fluorescence ratios as an estimate of relative superoxide levels. The DHE/EtBr ratio is 1.4-fold higher in the Δ SR and *sodB* mutants compared with the WT and +SR in stationary phase, but no differences are seen in exponential phase (Fig. 3A). Because aerobic respiration is a major source of superoxide (26), we also estimated the respiratory activity using resazurin reduction (27) and found no differences between the strains (SI Appendix, Fig. S1E). This leads us to infer that the excess accumulation of superoxide in the Δ SR mutant is likely attributable to its SOD defect, although other nonrespiratory sources of superoxide cannot be excluded. Additionally, we tested susceptibility to paraquat, a superoxide-generating compound, and found that killing by paraquat correlated with the SOD activity profiles across strains and growth phases (Fig. 3B). Combined, these results suggest that inactivation of the SR impairs protection against superoxide-mediated toxicity during stationary phase.

Genetic or Biochemical Restoration of SOD Activity Rescues Stationary-Phase Multidrug Tolerance. We first expressed pBAD-*sodB* in the Δ SR mutant (Δ SR +*sodB*), which restores its SOD activity (Fig. 4A) and DHE fluorescence levels (Fig. 4B) to WT levels. Notably, *sodB* expression in the Δ SR mutant also restores ofloxacin (Fig. 4C), gentamicin, and meropenem tolerance (SI Appendix, Fig. S4 A and B) to WT levels. Expression of *sodA*, the Mn-dependent SOD, similarly restores SOD activity and drug tolerance (SI Appendix, Fig. S5). Next, we chemically complemented the Δ SR mutant with 100 μ M Mn^{III}TMPyP, a cell-permeable SOD mimetic that catalytically dismutates superoxide at rates ~ 10 -fold lower than Mn-SodA and Fe-SodB (4×10^7 M⁻¹s⁻¹ vs. 7×10^8 M⁻¹s⁻¹) (28, 29). Mn^{III}TMPyP confers protection against antibiotic killing, with 2- to 3- \log_{10} higher viable CFU counts following challenges with ofloxacin (Fig. 4D), gentamicin, and meropenem (SI Appendix, Fig. S4 C and D). These results thus suggest that restoring

SOD activity in the Δ SR mutant is sufficient to rescue its stationary-phase tolerance defect.

SOD Activity Correlates with Multidrug Tolerance in a Dose-Dependent Manner. We exploited the biological variability of SOD activity in independent replicate batch cultures to assess the dose-dependent relationship between SOD activity and antibiotic tolerance. We measured SOD activity of a culture before antibiotic challenge and its bacterial viability following antibiotic challenge, and found a strong positive correlation with ofloxacin ($R^2 = 0.60$, $P < 0.01$) (Fig. 4E), gentamicin ($R^2 = 0.62$, $P < 0.01$), and meropenem tolerance ($R^2 = 0.64$, $P < 0.01$) (SI Appendix, Fig. S4 E and F).

Stationary-Phase Δ SR Cells Have Increased Membrane Permeability, Which Correlates with Antibiotic Killing. Internalization EtBr, which was used as a loading control for the DHE probe, also provides a well-established measure of global membrane permeability (30, 31). As expected, we first observed that EtBr internalization is 5- to 10-fold higher in exponential vs. stationary phase for all strains (Fig. 5A and SI Appendix, Fig. S3C). However, EtBr internalization is \sim twofold higher in the Δ SR and *sodB* mutants compared with WT and +SR strains during stationary phase, while no differences are seen during exponential phase. Furthermore, genetic restoration of SOD activity with pBAD-*sodB* expression in the Δ SR mutant restores its EtBr internalization to WT levels (Fig. 5B). Given that EtBr is a substrate for efflux pumps, which are highly expressed in *P. aeruginosa*, we wanted to confirm that differences in EtBr internalization were not due to differential efflux. To estimate of efflux activity, we measured the ratio of EtBr fluorescence with and without CCCP, a proton ionophore that inactivates efflux pumps, and found no differences between WT, Δ SR, and +SR cells (SI Appendix, Fig. S6 A and B). Finally, to further assess the outer membrane permeability, we carried out a periplasmic β -lactamase leakage assay (32) and found that Δ SR cells have leakier outer membranes than WT cells (SI Appendix, Fig. S6C), consistent with the EtBr results. Together, our results demonstrate that membrane permeability decreases in the WT

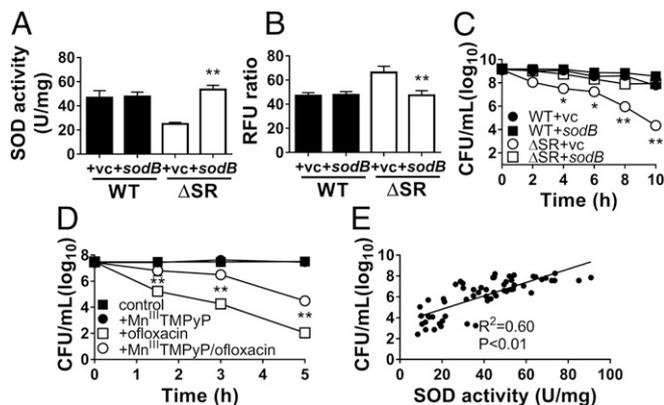


Fig. 4. Complementation of SOD activity restores superoxide levels and rescues ofloxacin tolerance in the Δ SR mutant. Stationary-phase WT and Δ SR cells carrying the pBAD-*sodB* (+*sodB*) or control (+vc) vector were assayed for (A) total SOD activity, (B) relative superoxide levels (DHE/EtBr fluorescence ratio), or (C) killing by 5 μ g/mL ofloxacin. (D) Stationary-phase Δ SR cells pretreated \pm the SOD mimetic Mn^{III}TMPyP before challenge with 5 μ g/mL ofloxacin. All results are shown as mean \pm SEM ($n \geq 6$). * $P < 0.05$ and ** $P < 0.01$ vs. WT +vc (in A to C) or vs. "+ofloxacin" (in D). (E) Correlation between bacterial viability of stationary-phase cells after ofloxacin challenge and SOD activity. The SOD activity was measured in stationary-phase cultures before ofloxacin challenge, and bacterial viability of the same culture was measured after 10-h challenge of 5 μ g/mL ofloxacin. Data from different strains (WT, Δ SR, Δ SR +*sodB*, Δ SR +*sodA*, +SR) were combined and each data point represents an independent replicate ($n \geq 40$). The correlation coefficient R^2 was calculated using linear regression.

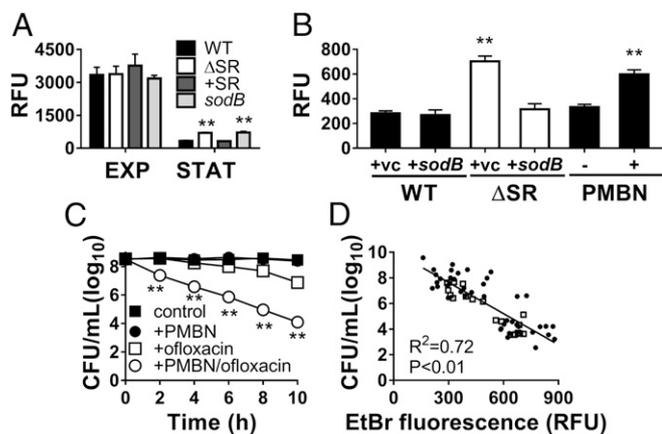


Fig. 5. The SR and SOD activity confers membrane impermeability, a key determinant of stationary-phase tolerance. EtBr internalization in the presence of CCCP in **A** of exponential (EXP) or stationary-phase (STAT) cells, **(B)** stationary-phase WT and Δ SR cells carrying the *pBAD-sodB* (+*sodB*) or control (+vc) vector, or in stationary-phase WT cells \pm 50 μ g/mL PMBN. **(C)** Killing of stationary-phase WT with 5 μ g/mL ofloxacin \pm PMBN pretreatment. All results are shown as mean \pm SEM ($n \geq 6$). $**P < 0.01$ vs. "WT+vc" or vs. "+PMBN". **(D)** Correlation between bacterial viability of stationary-phase cells after ofloxacin challenge and membrane permeability, as measured by the EtBr internalization assay of the same culture before ofloxacin challenge (5 μ g/mL for 10 h). Data from different strains (WT, Δ SR, Δ SR +*sodB*, Δ SR +*sodA*, +SR) noted as \bullet or WT +50 μ g/mL PMBN noted as \square , were combined and each data point ($n \geq 40$) represents an independent replicate. The correlation coefficient R^2 was calculated using linear regression.

upon transition from exponential to stationary phase, and this process is impaired in the Δ SR and *sodB* mutants.

To further validate the contribution of membrane permeability to drug tolerance, we permeabilized stationary-phase cells using a chemical approach with polymyxin B nonapeptide (PMBN). PMBN is a cationic peptide that binds outer membrane lipopolysaccharides, leading to permeabilization of the outer membrane without intrinsic bactericidal activity (33). Stationary-phase WT cells challenged with 50 μ g/mL PMBN alone show an \sim twofold increase in EtBr fluorescence (Fig. 5*B*) without loss of viability (Fig. 5*C*). Importantly, pretreatment of stationary-phase WT cells with PMBN significantly enhances ofloxacin killing (Fig. 5*C*) (4.4- vs. 1.4- \log_{10} killing at $t = 10$ h), as well as gentamicin and meropenem killing (*SI Appendix, Fig. S7 A and B*). Finally, we observed a significant linear correlation between EtBr fluorescence and tolerance to all three drugs (Fig. 5*D* and *SI Appendix, Fig. S7 C and D*), suggesting that membrane permeability is a major determinant of SR- and SOD-dependent antibiotic tolerance in stationary-phase *P. aeruginosa*.

Drug Penetration Is Enhanced by the Δ SR and *sodB* Mutant Permeability Defect.

Membrane permeability is a major determinant of drug uptake and internalization. We thus directly assessed intracellular drug levels in stationary-phase cells using ofloxacin (which is intrinsically fluorescent) (*SI Appendix, Fig. S6 A and B*), Texas Red-labeled gentamicin (*SI Appendix, Fig. S8 A and B*), and FITC-labeled meropenem (*SI Appendix, Fig. S8 C and D*). Stationary-phase Δ SR and *sodB* cells exhibit 2- to 2.5-fold higher drug uptake than WT or +SR complemented cells for all three drugs: ofloxacin (Fig. 6*A*), Texas Red gentamicin (*SI Appendix, Fig. S8 A*), and FITC-meropenem (*SI Appendix, Fig. S8 C*). Furthermore, *sodB* expression in the Δ SR mutant restores drug accumulation to WT levels (Fig. 6*B* and *SI Appendix, Fig. S8 B and D*). As controls, we confirmed that uptake of unconjugated Texas Red and FITC fluorophores is minimal and that the bactericidal activity of labeled drugs is similar to that of unconjugated ones (*SI Appendix, Fig. S9*).

Taking these data together, we demonstrate that ablation of (p)ppGpp signaling or deletion of *sodB* compromises membrane permeability specifically during stationary phase, leading to enhanced drug penetration and thus drug killing. Because restoration of SOD activity in the Δ SR mutant is sufficient to rescue both membrane impermeability and drug accumulation, this suggests that SOD activity is critical to SR-mediated membrane impermeability and multidrug tolerance in stationary-phase cells.

Loss of SOD Activity Abrogates the Emergence of Drug Resistance.

Genotypically resistant mutants likely arise from tolerant bacterial populations that survive sustained antibiotic exposure (1, 4). Hence, the loss of tolerance should abrogate the emergence of genotypic resistance. To test this, we measured the emergence of drug-resistant colonies from cell suspensions ($\sim 10^{11}$ CFU per plate) of stationary-phase bacteria spread and incubated for 5 d on agar plates containing ofloxacin. We enumerated newly emerging ofloxacin-resistant colonies after 72-h drug exposure to exclude those stemming from preexisting resistant cells. Consistent with our previous observations (16), we find that ablation of (p)ppGpp signaling all but eliminates the emergence of ofloxacin resistance, and this defect is restored in the complemented +SR strain (Fig. 6*C*). The development of ofloxacin resistance is also abrogated in *sodB* cells, while *sodB* expression in the Δ SR mutant restores the rate of ofloxacin-resistant mutants to WT levels (Fig. 6*D*). Hence, perturbations in SOD activity are sufficient to suppress the emergence of ofloxacin resistance.

Discussion

Our group and others have shown that the SR and (p)ppGpp signaling mediate antibiotic tolerance, likely through several different mechanisms (15, 16, 34, 35). Among these, the most extensively studied mechanism is the formation of specialized drug-tolerant cells termed persister cells (36). In *E. coli*, (p)ppGpp signaling is central to the regulation of toxin-antitoxin module-dependent pathways involved in persister formation (15, 36). In *P. aeruginosa*, Verstraeten et al. (34) reported that the Obg GTPase induces generation of aminoglycoside-tolerant cells, but through mechanisms yet unclear in *P. aeruginosa*. Although such recent studies provide important insights into the molecular basis of persister formation, it remains unclear to what degree persister mechanisms overlap or differ from those implicated in stationary-phase tolerance.

P. aeruginosa remains viable in a nonreplicating state for prolonged periods of time (8) and displays a high level of multidrug

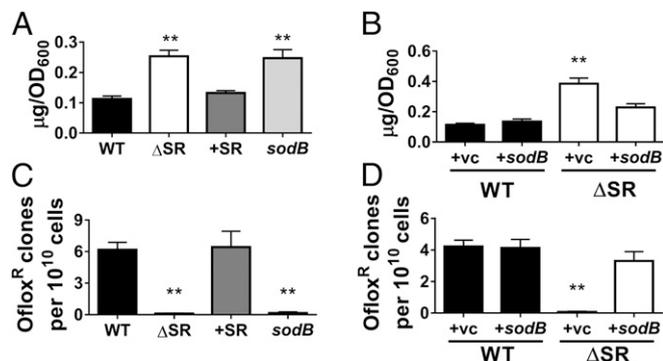


Fig. 6. Loss of (p)ppGpp signaling and SOD activity increase ofloxacin internalization and abrogates emergence of genotypic resistance to ofloxacin. Intracellular ofloxacin levels (RFU, Ex/Em 292/496 nm) in **(A)** stationary-phase WT, Δ SR, +SR, and *sodB* strains or **(B)** WT and Δ SR expressing the *pBAD-sodB* (+*sodB*) or control (+vc) vector. The number of ofloxacin-resistant colonies that arose between 3 and 5 d after incubation of 10^{11} cells of each strain on 12 μ g/mL ofloxacin LB agar plates in **(C)** WT, Δ SR, +SR, and *sodB* strains and **(D)** WT and Δ SR expressing the *pBAD-sodB* (+*sodB*). Results are mean \pm SEM ($n \geq 6$). $**P < 0.01$ vs. WT (for **A** and **C**) or WT +vc (for **B** and **D**).

tolerance during stationary phase. The SR acts as a major global regulator of bacterial stationary-phase physiology and stress responses that exerts widespread direct and indirect effects on gene transcription (11, 12). We show here that the SR modulates *sodB* transcription and total SOD activity in stationary-phase *P. aeruginosa* primarily through RpoS, which underscores the significant overlap between SR- and RpoS-dependent gene regulation, as we and others have previously noted (18, 37, 38). The link between the SR and SOD regulation is not widely conserved across different bacterial species (39–41) but SodB expression in *E. coli* is SR-dependent (42), and carbon starvation induces both SodA and SodB synthesis (43).

Despite the pleiotropic effects of the SR, genetic and chemical complementation of SOD activity is sufficient to restore multidrug tolerance of the Δ SR mutant to WT levels. This suggests that SOD activity is protective against antibiotic toxicity and plays a key role in (p)ppGpp-dependent multidrug tolerance during stationary phase. We observed a strong correlation between SOD activity and antibiotic survival to all three classes of drugs, across different isogenic strains constructed in the PAO1 genetic background. The role of the SR and SOD appears conserved in other *P. aeruginosa* clinical strains, although the magnitude of their effect likely varies based on the genetic background.

We recognize that our measurements of SOD activity and stationary-phase survival following antibiotic challenge reflect population rather than single-cell phenotypes. Gene expression is highly heterogeneous during stationary phase (8), and phenotypic heterogeneity allows bacterial populations to persist in the face of fluctuating environments and lethal stress, including antibiotics (17). Future studies with single-cell analyses would be informative in demonstrating whether variations in SOD activity confers survival to antibiotic stress at the individual cell level.

We used DHE to probe superoxide levels and acknowledge its lack of specificity because it can also react with hydroxyl radicals (44). However, in our experiments DHE fluorescence intensity varied with SOD activity in the *sodB* mutant (which has ~30% of the SOD activity of WT cells) (Fig. 2A) and in the *sodB* expression constructs (Fig. 4B), suggesting that the DHE signal reflects relative superoxide levels.

Although superoxide radicals are considered a primary source of intracellular oxidative stress and likely contribute to the lethal effects of antibiotics (45–47), the biological consequences of elevated superoxide are not fully understood. Whether SOD activity confers tolerance directly through its ability to detoxify superoxide radicals, or indirectly through downstream cellular processes responsive to SOD activity, remains to be determined. The relationship between SOD activity, SOD-dependent effects, and superoxide metabolism is complex. SODs appear to have moonlighting functions in eukaryotic cells. For example, SOD1 in the yeast *Saccharomyces cerevisiae* also functions in signal transduction as its SOD activity stabilizes kinases involved in oxidative and metabolic responses (48), and it can act as a nuclear transcription factor (49). Whether such unorthodox SOD functions exist in bacteria remains unknown.

To date, studies that examined the role of SODs in antibiotic lethality in various bacteria report divergent conclusions. *Sod* mutants show increased susceptibility to bactericidal antibiotics in *Campylobacter jejuni* (50), *Staphylococcus aureus* (51), and *Enterococcus faecalis* (52), but not *Acinetobacter baumannii* (53). A *sodA sodB* mutant of *E. coli* is reportedly no more susceptible than WT to ampicillin, gentamicin, and norfloxacin killing (54, 55). Furthermore, expression of *sodA* or *sodB* did not mitigate ampicillin and ofloxacin killing of *E. coli* (56). However, several important biological and experimental differences likely account for the different observations on the relationship between SOD and antibiotic susceptibility. First, many metabolic and antioxidant responses differ between *E. coli* and *P. aeruginosa*, including RpoS (57) and SoxR (58), particularly during growth arrest (8). Second, experimental conditions to assess antibiotic lethality differ significantly between the different studies. Most notably, antibiotic killing assays in the *E. coli* studies (54–56) were carried out on actively replicating (exponential phase) cells, or cells that exit stationary phase to resume growth upon dilution into fresh medium. In contrast, we challenged stationary-phase cells

under nongrowing conditions where new nutrients are not supplied, and our results demonstrate that SOD-mediated tolerance is growth phase-specific. Others have also reported that stationary-phase *sodA sodB* mutant *E. coli* is more susceptible to gentamicin killing when maintained in a nongrowing state (59). Combined, these results suggest that SOD is particularly important to antibiotic survival during stationary phase.

Interestingly, Dukan et al. (43) previously observed that SOD-deficient *E. coli* mutants exhibit increased protein oxidation, but only in stationary-phase cultures. They proposed that superoxide stress was a hallmark of respiring but nonreplicating stationary-phase cells (60). More recently, studies in *E. coli* and *S. aureus* reported that the antibiotic-tolerant stationary-phase cells and persister cells have low cellular ATP levels, and that manipulations that increase ATP levels also enhance antibiotic killing (61–63). Although increased respiration increases both ATP levels and superoxide generation, it is not known if SOD activity in turn alters ATP levels. Our results show that the differences in SOD activity and tolerance cannot be attributable to differences in respiration.

Here, we show that the *P. aeruginosa* cell envelope becomes less permeable during stationary phase, thus limiting drug penetration, and this process is SR- and SOD-dependent. Loss of cell permeability likely represents a common adaptive strategy for bacteria to survive under growth-limiting conditions (2). Alterations in cell envelope composition and permeability have been described in bacteria that transition into growth-limiting conditions, and those in stationary phase (64, 65). For example, the cell wall structure of stationary-phase *S. aureus* displays reduced cross-linking and increased peptidoglycan mass (66). Nutrient-starved and nonreplicating *Mycobacterium tuberculosis* have altered cell walls that display decreased permeability to chemically distinct classes of drugs, including fluoroquinolones (67). How SOD activity directly or indirectly contributes to the alterations in the cell envelope remains to be determined.

We measured intracellular drug concentrations, which result from the net effect of drug uptake and efflux. Because our results showed comparable efflux activity in the Δ SR and *sodB* mutants, this implies that increased drug accumulation is likely due to increased internalization. In Gram-negative bacteria, the outer membrane is widely recognized as the primary barrier to drug internalization, although the molecular mechanisms of drug uptake are often still poorly understood (68). We tested gentamicin, ofloxacin, and meropenem, three distinct classes of bactericidal antipseudomonal antibiotics targeting the bacterial ribosomal translational machinery, DNA gyrase, and peptidoglycan cell wall synthesis, respectively. We recognize that these chemically distinct drugs are likely internalized through different pathways. Aminoglycosides, such as gentamicin, require active uptake through a proton-gradient dependent process (69), while meropenem and ofloxacin likely diffuse passively through the membrane bilayer via outer membrane porins, such as *oprD* (70) and *ompF* (71), respectively. The mechanisms by which SR- and SOD-dependent tolerance influence these uptake processes remain unknown.

Our study uncovers an unprecedented link between SOD activity, SR-mediated multidrug tolerance, membrane permeability, and antibiotic internalization in stationary-phase *P. aeruginosa*. We also demonstrate that deletion of SOD-dependent pathways enhances antibiotic lethality and abrogates the emergence of genotypic resistance in stationary-phase *P. aeruginosa*. Thus, targeting the (p)ppGpp and SOD-dependent pathways may improve bactericidal activity against slow-growing bacteria and prevent the emergence of drug resistance in chronic infections.

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