



Synergistic Quinolone Sensitization by Targeting the *recA* SOS Response Gene and Oxidative Stress

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ABSTRACT Suppression of the *recA* SOS response gene and reactive oxygen species (ROS) overproduction have been shown, separately, to enhance fluoroquinolone activity and lethality. Their putative synergistic impact as a strategy to potentiate the efficacy of bactericidal antimicrobial agents such as fluoroquinolones is unknown. We generated *Escherichia coli* mutants that exhibited a suppressed $\Delta recA$ gene in combination with inactivated ROS detoxification system genes ($\Delta sodA$, $\Delta sodB$, $\Delta katG$, $\Delta katE$, and $\Delta ahpC$) or inactivated oxidative stress regulator genes ($\Delta oxyR$ and $\Delta rpoS$) to evaluate the interplay of both DNA repair and detoxification systems in drug response. Synergistic sensitization effects, ranging from 7.5- to 30-fold relative to the wild type, were observed with ciprofloxacin in double knockouts of *recA* and inactivated detoxification system genes. Compared to *recA* knockout, inactivation of an additional detoxification system gene reduced MIC values up to 8-fold. In growth curves, no growth was evident in mutants doubly deficient for *recA* gene and oxidative detoxification systems at subinhibitory concentrations of ciprofloxacin, in contrast to the *recA*-deficient strain. There was a marked reduction of viable bacteria in a short period of time when the *recA* gene and other detoxification system genes (*katG*, *sodA*, or *ahpC*) were inactivated (using absolute ciprofloxacin concentrations). At 4 h, a bactericidal effect of ciprofloxacin was observed for $\Delta katG \Delta recA$ and $\Delta ahpC \Delta recA$ double mutants compared to the single $\Delta recA$ mutant ($\Delta 3.4 \log_{10}$ CFU/ml). Synergistic quinolone sensitization, by targeting the *recA* gene and oxidative detoxification stress systems, reinforces the role of both DNA repair systems and ROS in antibiotic-induced bacterial cell death, opening up a new pathway for antimicrobial sensitization.

KEYWORDS resistance reversion, quinolones, SOS response, *recA*, oxidative stress, detoxification systems

Widespread and indiscriminate use of antibiotics in human and veterinary medicine and agriculture has become a global public health problem (1, 2), and the need to take measures to avoid returning to the preantibiotic era is now urgent (3). Efforts to overcome the problem of resistance have focused mainly on modifying existing antibiotics by circumventing molecular mechanisms conferring resistance (4). Various strategies are being studied to combat this emergence, including searching for new antibiotics, use of adjuvants that increase sensitivity, genome editing technologies, and phage therapy (4–9). While such efforts might be efficacious against resistant strains, new resistance mechanisms often arise in the process of adaptation to new

Citation Diaz-Diaz S, Recacha E, Machuca J, García-Duque A, Docobo-Pérez F, Blázquez J, Pascual A, Rodríguez-Martínez JM. 2021. Synergistic quinolone sensitization by targeting the *recA* SOS response gene and oxidative stress. *Antimicrob Agents Chemother* 65:e02004-20. <https://doi.org/10.1128/AAC.02004-20>.

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Received 18 September 2020

Returned for modification 19 October 2020

Accepted 28 January 2021

Accepted manuscript posted online

1 February 2021

Published 18 March 2021

antimicrobial agents (10, 11). Hence, it becomes necessary to discover how to reverse resistance or to sensitize bacteria to old antibiotics and block their evolution (9).

Enterobacteriales bacteria such as *Escherichia coli* are among the most common causes of community and nosocomial infections. Fluoroquinolones are used for empirical and directed therapy in infections caused by *E. coli* (12). Fluoroquinolones are broad-spectrum antimicrobials whose mechanism of action is related to the inhibition of two enzymes essential for bacterial viability, DNA gyrase and topoisomerase type IV, which are involved in DNA synthesis. Target mutations in the topoisomerases cause resistance to these antimicrobial agents (13). Plasmid-mediated quinolone resistance mechanisms have also been described (14).

Microorganisms have the ability to evolve rapidly to ensure survival, which has led to high levels of antibiotic resistance. The SOS response is a coordinated cellular response to genotoxic damage, and it can contribute to this evolution (low-fidelity polymerases). Some antibiotics induce SOS by a variety of molecular mechanisms. Specifically, fluoroquinolones activate the SOS response caused by DNA damage to arrest replication forks (15, 16). A number of studies have recently demonstrated the role of RecA protein activation, which is associated with the SOS response in *E. coli*, in the emergence of mutations in *gyrA* and *parC* that lead to resistance (15, 17–19). LexA protein represses the SOS regulon, which consists of more than 50 genes, many associated with different repair mechanisms (16, 20, 21). When DNA damage occurs, single-strand DNA activates RecA, which forms nucleofilaments around it and induces autoproteolysis of LexA and activation of the SOS response. Sequential induction of genes could facilitate the transition from high-fidelity to lower-fidelity DNA repair mechanisms (16, 21). Recent studies have demonstrated that suppression of the SOS response enhances the bactericidal activity of antimicrobials such as quinolones in bacteria with specific resistant mechanisms (17, 22).

It is also known that bactericidal agents are involved in oxidative stress since, under aerobic conditions, they produce reactive oxygen species (ROS). Downstream of their target-specific interactions, bactericidal antibiotics induce complex redox alterations that contribute to cellular damage and death, thus supporting their involvement in antibiotic lethality (23–25). *In vitro* studies have shown a protective effect against ROS, produced by antibiotic, of some antioxidant molecules, such as ascorbic acid and *N*-acetylcysteine (25, 26). Bacteria also have multiple oxidative detoxification systems, such as superoxide dismutase (SOD) and catalase, to combat oxidative stress (27, 28). Specifically, *E. coli* has three types of SOD (SodA, SodB, and SodC), two types of catalase (KatG and KatE), which remove O₂⁻ and H₂O₂, respectively, and an alkyl hydroperoxide reductase (made of two subunits, AhpF and AhpC), which reduces NADH while removing H₂O₂. Transcriptional regulation is controlled by oxidative stress regulators such as OxyR or general stress response regulators such as RpoS. During oxidative stress, OxyR increases the expression of *katG* and *ahpC* genes, while RpoS regulates the expression of *katE* and *sodC* (29).

Hence, suppression of the SOS response and overproduction of ROS have been shown, separately, to enhance the activity and lethality of fluoroquinolones. Nevertheless, the use of their putative synergistic effect as a strategy to reinforce the efficacy of fluoroquinolones remains an open question. To address this point, we generated a wide collection of *E. coli* mutants that exhibited a suppressed SOS response ($\Delta recA$) in combination with inactivated ROS detoxification system genes ($\Delta sodA$, $\Delta sodB$, $\Delta katG$, $\Delta katE$, $\Delta sodA$, and $\Delta ahpC$) or inactivated oxidative stress regulatory genes ($\Delta oxyR$ and $\Delta rpoS$) to evaluate the interplay of both DNA repair/recombination and detoxification systems in the antimicrobial drug response. Our comprehensive analysis opens up a new strategy for bacterial sensitization by double-targeting SOS response and oxidative stress.

RESULTS

Quinolone sensitization by targeting the *recA* SOS response gene and oxidative stress response. We generated *E. coli* mutants that exhibited the deleted *recA* gene in combination with suppressed ROS detoxification system genes ($\Delta sodA$, $\Delta sodB$, $\Delta katG$, $\Delta katE$, and $\Delta ahpC$) or genes regulating oxidative stress ($\Delta oxyR$ and

TABLE 1 Heat map of the fold decrease in ciprofloxacin MICs of different *recA* and detoxification system inactivations^a

	BW25113	$\Delta katG$	$\Delta katE$	$\Delta sodA$	$\Delta sodB$	$\Delta ahpC$	$\Delta oxyR$	$\Delta rpoS$	$\Delta katG / \Delta katE$	$\Delta sodB / \Delta sodA$	$\Delta recA$	$\Delta katG / \Delta recA$	$\Delta katE / \Delta recA$	$\Delta sodA / \Delta recA$	$\Delta sodB / \Delta recA$	$\Delta ahpC / \Delta recA$	$\Delta oxyR / \Delta recA$	$\Delta rpoS / \Delta recA$	$\Delta katG / \Delta katE / \Delta recA$
BW25113	1																		
$\Delta katG$	1	1																	
$\Delta katE$	1	1	1																
$\Delta sodA$	1	1	1	1															
$\Delta sodB$	1	1	1	1	1														
$\Delta ahpC$	1	1	1	1	1	1													
$\Delta oxyR$	1	1	1	1	1	1	1												
$\Delta rpoS$	1	1	1	1	1	1	1	1											
$\Delta katG / \Delta katE$	1	1	1	1	1	1	1	1	1										
$\Delta sodB / \Delta sodA$	1	1	1	1	1	1	1	1	1	1									
$\Delta recA$	3,75	3,75	3,75	3,75	3,75	3,75	3,75	3,75	3,75	3,75	1								
$\Delta katG / \Delta recA$	15	15	15	15	15	15	15	15	15	15	4	1							
$\Delta katE / \Delta recA$	7,5	7,5	7,5	7,5	7,5	7,5	7,5	7,5	7,5	7,5	2	0,5	1						
$\Delta sodA / \Delta recA$	15	15	15	15	15	15	15	15	15	15	4	1	2	1					
$\Delta sodB / \Delta recA$	15	15	15	15	15	15	15	15	15	15	4	1	2	1	1				
$\Delta ahpC / \Delta recA$	15	15	15	15	15	15	15	15	15	15	4	1	2	1	1	1			
$\Delta oxyR / \Delta recA$	30	30	30	30	30	30	30	30	30	30	8	2	4	2	2	2	2	1	
$\Delta rpoS / \Delta recA$	7,5	7,5	7,5	7,5	7,5	7,5	7,5	7,5	7,5	7,5	2	0,5	1	0,5	0,5	0,5	0,25	1	
$\Delta katG / \Delta katE / \Delta recA$	15	15	15	15	15	15	15	15	15	15	4	1	2	1	1	1	0,5	2	1

^aMIC values represent the means of at least three independent determinations performed on separate days. The raw MIC values of all strains are shown in Table S1 in the supplemental material.

$\Delta rpoS$) to evaluate the interplay of both DNA repair and detoxification systems in anti-microbial drug response (Table 1; see Table S1 in the supplemental material).

We confirmed first that the *recA* knockout produces the expected perturbation of the SOS response in both the wild-type BW25113 and the various detoxification system knockouts. Figure S1 represents the results obtained for each isogenic group used in this study. No fluorescence was observed when *recA* was inactivated, confirming the perturbation of the SOS response (together with DNA repair and recombination). In terms of MIC values, as expected, the ciprofloxacin MICs were 3.75-fold lower for the $\Delta recA$ mutant compared to wild-type BW25113 (Table 1, Table S1). No reductions in ciprofloxacin MIC were observed when any of the detoxification system genes were inactivated, including single ($\Delta katG$, $\Delta katE$, $\Delta sodA$, $\Delta sodB$, $\Delta ahpC$, $\Delta oxyR$, and $\Delta rpoS$) and double knockouts ($\Delta katG \Delta katE$ and $\Delta sodB \Delta sodA$) compared to wild-type BW25113.

Notably, in general, a synergistic sensitization effect was observed in double knockouts of *recA* (with SOS inactivation deficient for DNA repair and recombination) and the different ROS detoxification system genes that were inactivated (see Table S1). Fractional inhibitory concentration (FIC) values were below 0.5 for double and triple gene deletion mutants, 0.16 ($\Delta oxyR \Delta recA$) and 0.32 ($\Delta katG \Delta recA$, $\Delta sodA \Delta recA$, $\Delta sodB \Delta recA$, $\Delta ahpC \Delta recA$, and $\Delta katG \Delta katE \Delta recA$). Only $\Delta katE \Delta recA$ and $\Delta rpoS \Delta recA$ show an additive effect, as their FIC value was 0.63; the value was located in the range of 0.5 to ≤ 1 . The process of sensitization was similarly efficient across all double combinations, including the effector detoxification genes (*sodA*, *sodB*, *katG*, *katE*, *sodA*, and *ahpC*) and genes regulating oxidative stress (*oxyR* and *rpoS*), which ranged from 7.5-fold to 30-fold compared to wild-type BW25113 (Table 1). The highest sensitization was found for the $\Delta recA \Delta oxyR$ double knockout, with a ciprofloxacin MIC value of 0.0005 mg/liter (Table 1, Table S1). For most of the combinations, sensitization increased about 15-fold compared to wild-type BW25113. Compared to values for the *recA* knockout, inactivation of an additional detoxification system gene reduced MIC

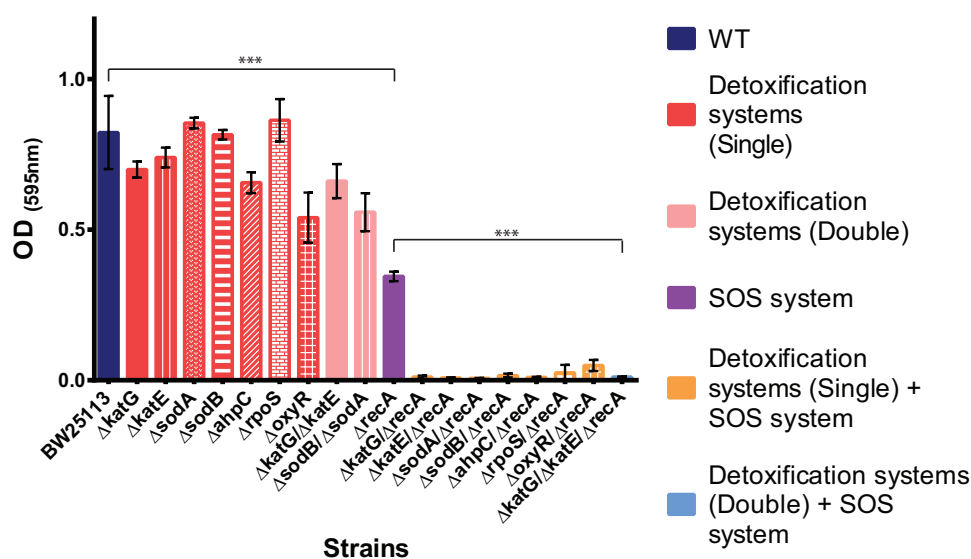


FIG 1 The impact of *recA* gene inactivation and detoxification system suppression on cell growth in the presence of sublethal quinolone concentrations. OD values for *E. coli* BW25113 and all genetic combinations with ciprofloxacin concentrations of 0.001 mg/liter at 20 h. Data are represented as the mean of at least four independent measurements. The OD of all constructions was similar in the nonantibiotic control (data not shown). Standard deviations are indicated by error bars. Significant *P* values are noted (***, *P* < 0.001).

values between 2-fold and 8-fold. Here, we show that, in terms of MIC, *recA* gene suppression combined with inactivation of multiple detoxification system genes sensitizes to ciprofloxacin in a synergistic way.

As expected, the effect of synergistic sensitization to ciprofloxacin following *recA* gene suppression in combination with inactivation of detoxification system genes was not evident under anaerobic conditions (Table S1).

In addition, we also tried to evaluate the impact of this strategy on an evolved *E. coli* mutant strain harboring acquired quinolone resistance mechanisms (BW15 strain in Table S1). This mutant harbored changes affecting the *gyrA*, *gyrB*, *parE*, and *marR* genes, including a D87G substitution in the *gyrA* gene. The modifications produced a ciprofloxacin-resistant phenotype with a MIC value of 2 mg/liter according to CLSI and EUCAST breakpoints (30, 31). As expected, the *recA*-deficient BW15 derivative involved a change of category to susceptible-intermediate or areas of technical uncertainty (ATU) according to CLSI and EUCAST breakpoints (MIC of 0.5 mg/liter). Unexpectedly, no further sensitization was apparent when the *recA* gene and detoxification system genes (*katG* or *sodA*) were inactivated in this resistant mutant (Table S1).

Besides synergistic sensitization found for ciprofloxacin, in terms of MICs, was not always reproduced for other fluoroquinolones in the multiple genetic combinations assayed (Table S1). Based on these data, the rest of the study focused on the synergistic effects of sensitization to ciprofloxacin as a result of *recA* gene suppression in combination with inactivation of detoxification systems.

Monitoring bacterial *in vitro* growth in the presence of quinolones. *In vitro* growth curves were analyzed at subinhibitory concentrations of ciprofloxacin (0.001 mg/liter, 1/15× MIC based on the susceptibility of the BW25113 strain with the unmodified *recA* gene or ROS detoxification system genes) (Fig. 1). At this concentration, growth (after 20 h) was observed for the wild type and mutants with deficient detoxification systems (optical density [OD] values ranged from 0.54 to 0.86). As expected, reduced growth was observed for the *recA*-deficient strain (OD value, 0.35) (*P* < 0.001 compared to wild-type BW25113; OD value, 0.82). Interestingly, no growth was found for mutants doubly deficient in the *recA* gene and oxidative detoxification systems (OD of less than 0.05), including effector detoxification genes (*sodA*, *sodB*,

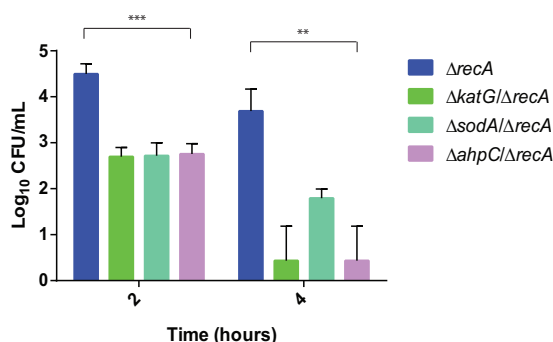


FIG 2 Combination of *recA* gene inactivation and detoxification system suppression inactivation enhances synergistically bactericidal activity of quinolones. Viable bacterial counts of *E. coli* $\Delta recA$, *E. coli* $\Delta katG \Delta recA$, *E. coli* $\Delta sodA \Delta recA$, and *E. coli* $\Delta ahpC \Delta recA$ in time-kill assays at ciprofloxacin (CIP) concentrations of $1 \times$ MIC $\Delta recA$ (0.004 mg/liter) at 2 and 4 h. Data are represented as the means of at least three independent measurements. *E. coli* BW25113 was not included because the antibiotic concentration used was under its MIC, so its behavior was similar to the nonantibiotic control. Viable bacterial counts of all constructions were similar in the nonantibiotic control (data not shown). Error bars represent standard deviations. Significant *P* values are noted (**, $P < 0.01$; ***, $P < 0.001$).

katG, *katE*, *sodA*, and *ahpC*) and regulatory genes of oxidative stress (*oxyR* and *rpoS*) ($P < 0.001$ compared to *recA*-deficient BW25113) (Fig. 1).

Figure S2 shows full monitoring of *in vitro* growth curves at subinhibitory concentrations (0.0005 to 0.002 mg/liter) of ciprofloxacin for each isogenic group. Under these conditions, no growth or a very marked delay in growth was observed in mutants doubly deficient in the *recA* gene and oxidative detoxification systems compared to *recA*-deficient BW25113. For example, for effector detoxification genes such as the *katE* group (Fig. S2B), at concentrations of 0.001 mg/liter of ciprofloxacin, only the BW25113 strain and $\Delta katE$ grew regularly at both 8 h and 16 h (this behavior was stable after 24 h). Significantly delayed growth was observed for $\Delta recA$, and no growth was observed for the $\Delta katE \Delta recA$ combination. A similar pattern was obtained for the rest of the effector and regulator detoxification genes (Fig. S2).

These data support the efficacy of combining the inactivation of the *recA* gene with detoxification system genes in order to prevent bacterial growth in the presence of sublethal concentrations of ciprofloxacin over shorter and longer periods of time.

The *recA* gene and oxidative stress response suppression enhances ciprofloxacin bactericidal activity. To show the synergistic impact of suppressing the *recA* gene and the oxidative stress response in terms of bacterial viability, time-kill curves were obtained for each isogenic group (*recA*, *katG*, *sodA*, and *ahpC*) (Fig. 2 and Fig. S3). At $1 \times$ MIC relative to the $\Delta recA$ MIC value of ciprofloxacin (0.004 mg/liter), a marked reduction in viable bacteria was observed in a short period of time (2 and 4 h) when the *recA* gene and other ROS detoxification system genes (*katG*, *sodA*, or *ahpC*) were inactivated (Fig. 2). At 2 h, a bacteriostatic effect (drop of $<3 \log_{10}$ CFU/ml) was observed for the three detoxification system genes (although a marked difference was observed compared to the single $\Delta recA$ mutant, up to $\Delta 1.8 \log_{10}$ CFU/ml for the $\Delta katG \Delta recA$ genes) ($P < 0.001$). At 4 h, however, a bactericidal effect (drop of $>3 \log_{10}$ CFU/ml) was observed for the double mutants $\Delta katG \Delta recA$ and $\Delta ahpC \Delta recA$ compared to the single $\Delta recA$ mutant ($\Delta 3.4 \log_{10}$ CFU/ml for the $\Delta katG \Delta recA$ and $\Delta ahpC \Delta recA$ genes) ($P < 0.01$) (Fig. 2).

At $1 \times$ MIC relative to the wild-type MIC of ciprofloxacin, 0.015 mg/liter (Fig. S3), a marked reduction in viable bacteria was also observed after a short and long period of time (2 to 8 h) when the *recA* gene together with other detoxification system genes (*sodA* or *ahpC*) was inactivated, compared to the single *recA* mutant. Nevertheless, this concentration appeared to be too high to demonstrate differences in the case of the *katG* gene group (BW25113, $\Delta katG$, $\Delta recA$, and $\Delta katG \Delta recA$). Under these conditions, a drop of $1.79 \log_{10}$ CFU/ml was observed at 4 h for the $\Delta ahpC \Delta recA$ double mutant

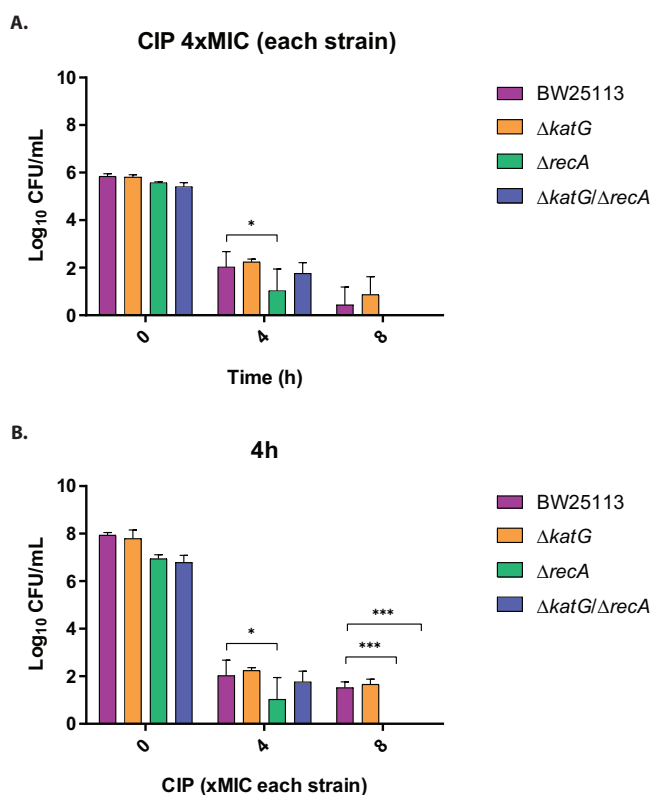


FIG 3 (A and B) Viable bacterial counts of *E. coli* BW25113 with *recA* gene and *katG* detoxification system inactivation alone or in combinations in time-kill assays at (A) ciprofloxacin (CIP) concentrations of 4× MIC of each strain (normalized MICs, 0.06 mg/liter for BW25113 and $\Delta katG$, 0.015 mg/liter for $\Delta recA$, and 0.004 mg/liter for $\Delta katG \Delta recA$) at 0 h, 4 h, and 8 h and (B) ciprofloxacin concentrations of 4× MIC and 8× MIC of each strain (normalized MICs, 0.125 mg/liter for BW25113 and $\Delta katG$, 0.03 mg/liter for $\Delta recA$, and 0.008 mg/liter for $\Delta katG \Delta recA$) and the nonantibiotic control at 4 h. Data are represented as the means of at least three independent measurements. Standard deviations are shown. CIP, ciprofloxacin. Significant *P* values are noted (*, *P* < 0.05; ***, *P* < 0.001).

compared to the single $\Delta recA$ mutant (Fig. S3). Interestingly, after 8 h, no viable bacteria were detected for the single $\Delta recA$ mutant and the $\Delta katG \Delta recA$, $\Delta soda \Delta recA$, and $\Delta ahpC \Delta recA$ double mutants compared to wild-type *E. coli* BW25113 and single mutants for detoxification system genes.

From another point of view, we also determined bacterial viability, specifically for the isogenic group KatG, at normalized drug concentrations according to the MICs. At a ciprofloxacin concentration of 4× MIC relative to each strain (0.06 mg/liter for BW25113 and $\Delta katG$; 0.015 mg/liter for $\Delta recA$ and 0.004 mg/liter for $\Delta katG \Delta recA$) at 0 h, 4 h, and 8 h, no significant differences were observed between strains, with the exception of $\Delta recA$ relative to control strain BW25113 (*P* < 0.05) (Fig. 3A). Also, at 8× MIC relative to each strain (0.125 mg/liter for BW25113 and $\Delta katG$, 0.03 mg/liter for $\Delta recA$, and 0.008 mg/liter for $\Delta katG \Delta recA$) at 4h, significant differences were observed between $\Delta recA$ and $\Delta katG \Delta recA$ relative to control strain BW25113 (*P* < 0.001) (Fig. 3B), but not between them. These data support that observed differences in lethality could be due to initial differences observed in terms of MIC values.

Additionally, we determined how antioxidants, such as L-ascorbic acid, affects drug lethality when detoxification systems are intact or suppressed in a *recA*-deficient background (Fig. S4). We found that pretreatment with L-Ascorbic acid provided at least 1 log₁₀ of protection from cell death induced by ciprofloxacin (1× MIC of $\Delta recA$) at 4 h posttreatment in a *recA*-deficient background. The protection was higher when, in addition to a *recA*-deficient background, other detoxification system genes (*katG* or

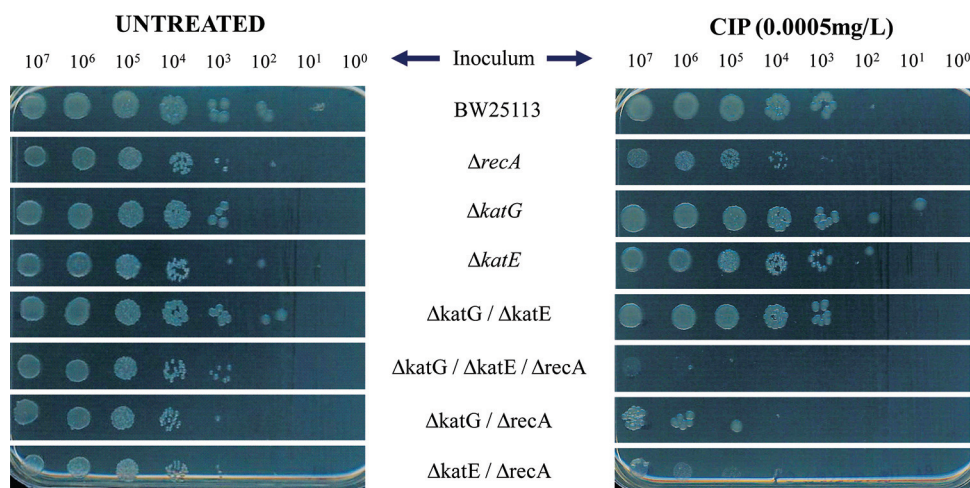


FIG 4 Survival of *E. coli* mutants by spot test assays treated with ciprofloxacin under aerobic conditions. Overnight liquid cultures were serially diluted and plated on LB agar with or without ciprofloxacin at concentrations of 0.0005 mg/liter. The results shown are representative of at least three independent experiments.

ahpC) were inactivated (almost 2 log₁₀ of protection). Minor differences were observed for the $\Delta sodA \Delta recA$ double mutant compared to the $\Delta recA$ mutant. Notably, L-ascorbic acid slightly increased the MIC against ciprofloxacin (2- to 4-fold, data not shown), supporting again that differences in lethality could be due in part to differences observed in MIC values.

Bacterial survival of *E. coli* mutants after treatment with ciprofloxacin under aerobic conditions was also evaluated using spot test assays with multiple inocula (Fig. 4). Under these conditions, the $\Delta recA$ mutant was more sensitive than wild-type BW25113 to the presence of ciprofloxacin. Furthermore, double and triple mutants with a combination of the *recA* gene and detoxification system genes ($\Delta katG \Delta recA$, $\Delta katE \Delta recA$, and $\Delta katG \Delta katE \Delta recA$) were more sensitive than the single $\Delta recA$ mutant. Higher susceptibility was not observed for mutants with deletions only in the detoxification system genes (Fig. 4).

Under these conditions, the *recA* gene and oxidative stress response suppression lead to a higher bactericidal effect in *E. coli* compared to *recA* gene inactivation alone.

Increased ROS production in the *recA* gene and oxidative stress response-suppressed backgrounds. At the test concentration (0.004 mg/liter of ciprofloxacin, 1× MIC $\Delta recA$), only minor, nonsignificant differences in ROS production were observed for single detoxification system gene mutants compared to the wild-type BW25113 (Fig. S5). The tested concentration was too low to show obvious differences in ROS production. However, as expected, at this concentration of ciprofloxacin, strong significant differences ($P < 0.001$) were observed in the $\Delta recA$ mutant versus wild-type BW25113. In general, a trend toward increased ROS production was observed with the combination of *recA* gene and detoxification system suppression in double and triple mutants compared to the single $\Delta recA$ mutant. This comparison was significant in the case of $\Delta katG \Delta recA$ ($P < 0.05$), $\Delta rpoS \Delta recA$ ($P < 0.001$), and $\Delta katG \Delta katE \Delta recA$ ($P < 0.05$) mutants (Fig. S5). This association between increased ROS production and inactivation of the *recA* gene-detoxification systems following treatment with a bactericidal antibiotic support that ROS may actively contribute to the bactericidal activity of fluoroquinolones.

DISCUSSION

Bacterial stress responses, such as the SOS response, play an important role in adaptation and acquired bacterial resistance to antibiotics. The key regulators of the SOS response (LexA and RecA) have been proposed as an attractive target for increasing bacterial sensitivity to antibiotics and combating the emergence of resistance.

Systematically altering the bacterial SOS activity, mainly through *recA* gene inactivation, has been revealed as a possible therapeutic strategy for potentiating bactericidal antibiotics such as quinolones against susceptible and resistant *E. coli*. Beyond this, *recA* inactivation modifies DNA repair and recombination events (17, 22, 32). Several compounds have also been shown to inhibit the ATPase activity of RecA *in vitro* (26, 33, 34). Phthalocyanine tetrasulfonates were recently characterized as an *in vivo* RecA inhibitor (35). ROS, on the other hand, are formed in cells exposed to three major classes of bactericidal antibiotics: quinolones, β -lactams, and aminoglycosides. This observation led to the theory that downstream formation of ROS is a common part of the killing mechanism of bactericidal antimicrobials (36). According to this line of thought, a secondary effect of exposure to these antibiotics is ROS formation through a common chain of events, which is an equally important and possibly more lethal part of the killing mechanism than the primary effect (25, 37–39). Hence, given this new way of thinking about the response of bacterial populations to lethal stress, efforts to target oxidative stress pathways as adjuvant antimicrobial therapy could increase lethality (40). In light of this literature, and even though much clinical investigation remains to be delivered on ROS therapy, *in vitro* work on infection models and early clinical evaluations looks promising (41, 42).

In this study, we used a wide collection of mutants to evaluate the impact of the *recA* gene (affecting the SOS response and DNA repair) and multiple ROS detoxification systems on the activity of quinolones such as ciprofloxacin. We showed that the combination of *recA* inactivation in combination with detoxification system gene inactivation produced a synergistic sensitization effect in terms of FIC index and susceptibility to ciprofloxacin. Furthermore, combined suppression of the *recA* gene and oxidative stress response leads to a higher bactericidal effect in *E. coli* than *recA* gene inactivation alone. Our results represent an important advance in the design and improvement of possible strategies to generate adjuvants to treatment with bactericidal antimicrobials.

The process of sensitization, using this double targeting strategy, was equally efficient across all genes, including effector detoxification genes and those regulating oxidative stress, and ranged from 7.5-fold to 30-fold relative to wild-type BW25113 (Table 1). Although these differences were moderate, they could play a significant role in therapeutic failure, bearing in mind the concentration-dependent character of these antimicrobials, whose predictors of efficacy *in vivo* are C_{max}/MIC and AUC/MIC (43, 44).

In terms of kinetic assays, both growth curves and killing curves supported the synergistic effect of sensitization to ciprofloxacin through *recA* gene suppression in combination with inactivation of detoxification system genes (Fig. 1 and 2). Rapid quinolone-mediated killing was increased in a *recA*-deficient *E. coli* background after detoxification system inactivation (for *katG*, *sodA*, and *ahpC*) when absolute concentrations were used (not normalized to MICs). A bactericidal effect (drop of $>3 \log_{10}$ CFU/ml) was observed for the $\Delta katG \Delta recA$ and $\Delta ahpC \Delta recA$ double mutants relative to the single $\Delta recA$ mutant (Fig. 2). However, these differences were not evident when high killing concentrations normalized to MIC values were used (Fig. 3), indicating that mechanistic differences are not attributable to the lethality process. In addition, these data support a synergistic sensitization effect for ciprofloxacin, confirming that ROS are relevant factors in the quinolone-mediated lethality mode (25, 40, 45). More importantly, synergistic lethality was enhanced in the absence of DNA repair systems like those under the control of the SOS response (Fig. 5). Hence, even highly lethal fluoroquinolones can be improved by finding ways to increase lethality through the SOS response and ROS accumulation. ROS-stimulating and SOS response inhibitor adjuvants may enhance the lethality of quinolones and possibly other antimicrobials. Accordingly, some antioxidants, such as L-ascorbic acid, provide protection against antibiotic lethality (25).

It is important to mention that a wide range of ciprofloxacin concentrations were initially used in this study, some of them being excessively lethal (Fig. S6 and S7). The

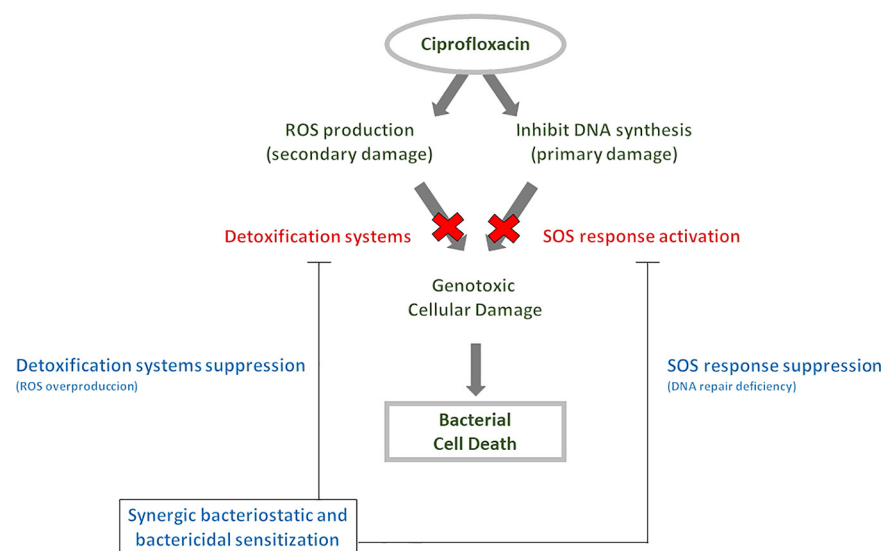


FIG 5 Scheme describing synergistic quinolone sensitization by targeting the SOS response and oxidative stress.

results of the conditions that showed the greatest differences were selected to elaborate this study.

In summary, we highlight synergistic quinolone sensitization by targeting the *recA* gene and oxidative detoxification stress systems. These data reinforce the role of DNA repair systems and ROS in antibiotic-induced bacterial cell death and how both pathways can act synergistically in the quinolone sensitization process. Our detailed analysis opens up a new strategy for sensitizing drug resistance by targeting the *recA* gene and oxidative stress.

MATERIALS AND METHODS

Strains, growth conditions, and antimicrobial agents. Wild-type *E. coli* BW25113 was used as the starting strain for all constructions (Table S1). *E. coli* BW25113 (wild-type) single-gene deletion mutants ($\Delta recA$, $\Delta katG$, $\Delta katE$, $\Delta sodA$, $\Delta sodB$, $\Delta ahpC$, $\Delta oxyR$, and $\Delta rpoS$) were selected from the KEIO collection (46). Double-gene deletion mutants ($\Delta katG \Delta recA$, $\Delta katE \Delta recA$, $\Delta sodA \Delta recA$, $\Delta sodB \Delta recA$, $\Delta ahpC \Delta recA$, $\Delta oxyR \Delta recA$, $\Delta rpoS \Delta recA$, $\Delta katG \Delta katE$, and $\Delta sodB \Delta sodA$) and the triple-gene deletion mutant ($\Delta katG \Delta katE \Delta recA$) were generated by P1vir phage transduction and by excision of the kanamycin resistance cassette with the plasmid pCP20 expressing the Flp recombinase (47, 48). Although some triple knockouts such as $\Delta sodA \Delta sodB \Delta recA$ were planned at the start of the study, they proved not to be viable after several attempts.

Liquid or solid Luria broth (LB) medium, Mueller-Hinton broth (MHB), and M9 minimal medium were used. Strains were grown at 37°C. Ciprofloxacin was used in all assays (Sigma-Aldrich, Madrid, Spain). Kanamycin (Sigma-Aldrich) at 30 mg/liter was used for plasmid maintenance. Expression of ciprofloxacin-induced GFP into the kanamycin-resistant pMS*recA*-gfp vector was used to detect *recA* transcription (see Table S2) (17, 49).

MICs. MICs were determined in triplicate for each bacterial strain under conditions of aerobiosis and anaerobiosis, using two techniques, the gradient strip technique and broth microdilution following CLSI reference methods (30).

FIC index analysis. The FIC index was calculated for a double gene deletion mutant with the equation $FIC = MIC(\text{double gene deletion mutant})/MIC(\text{recA gene deletion mutant}) + MIC(\text{double gene deletion mutant})/MIC(\text{ROS detoxification gene deletion mutant})$. In the case of the triple gene deletion mutant, the FIC index was calculated with the equation $FIC = MIC(\text{triple gene deletion mutant})/MIC(\text{recA gene deletion mutant}) + MIC(\text{triple gene deletion mutant})/MIC(\text{double gene deletion mutant } \Delta katG \Delta katE)$. The combination of two systems was considered to be synergistic when the FIC value was ≤ 0.5 , additive when it was 0.5 to ≤ 1 , indifferent when it was 1 to 4, and antagonistic when it was > 4 , following the criteria of (50).

Bacterial growth rates. Transparent 96-well flat-bottom plates containing 200 μ l of sublethal concentrations ($1/8 \times MIC$ to $1/32 \times MIC$ of wild-type) of ciprofloxacin in MHB were prepared as previously described using an Infinite 200 PRO plate reader (Tecan, Madrid, Spain) (17). At least four biological replicates were measured for each condition in at least two independent assays.

Time-kill curve assays. To show the synergistic effect of suppression of RecA protein and oxidative detoxification systems on bacterial viability when challenged by ciprofloxacin, time-kill assays were performed with each isogenic group, based on RecA inactivation, specific detoxification system suppression, or combinations. LB was used with ciprofloxacin concentrations of $1\times$ MIC of the wild type and $1\times$ MIC of $\Delta recA$ (ciprofloxacin concentrations were relative to MICs for strains harboring the unmodified *recA* gene or $\Delta recA$ mutant, respectively). Additionally, bacterial viability, specifically for the isogenic group *katG*, at normalized ciprofloxacin concentrations according to the MICs ($4\times$ MIC and $8\times$ MIC relative to the MICs of each strain, normalized MICs) were also evaluated. Growth in drug-free broth was evaluated in parallel as a control. Cultures were incubated at 37°C with shaking at 180 rpm. An initial inoculum of 10^6 CFU/ml was used in all experiments, and viable cells were determined at 0, 2, 4, 6, 8, and 24 h by colony counting.

Spot test. Fresh overnight cultures in LB were diluted to achieve an optical density at 625 nm ($\text{OD}_{625\text{nm}}$) of 0.1 (ca. 10^8 CFU/ml) and then serially diluted in saline solution (1%). A $5\text{-}\mu\text{l}$ volume of each dilution was spotted on LB agar plates with $1/8\times$ MIC (0.0005 mg/liter) of ciprofloxacin (relative to the $\Delta recA$ mutant) under aerobic conditions. After 20 h of incubation at 37°C , spots were checked for growth and compared with spots on the control LB agar plates.

Statistical analysis. All statistical analyses were performed using GraphPad Prism 6 software. Student's *t* test was used for statistical evaluation when two groups were compared. Differences were considered significant when *P* values were <0.05 .

SUPPLEMENTAL MATERIAL

Supplemental material is available online only.

SUPPLEMENTAL FILE 1, PDF file, 0.7 MB.

ACKNOWLEDGMENTS

This work was supported by the Plan Nacional de I+D+i 2013-2016 and the Instituto de Salud Carlos III (projects PI14/00940 and PI17/01501), Subdirección General de Redes y Centros de Investigación Cooperativa, Ministerio de Economía, Industria y Competitividad, Spanish Network for Research in Infectious Diseases (REIPI; RD16/0016/0001 and REIPI RD16/0016/0009) cofinanced by the European Development Regional Fund "A Way to Achieve Europe," operative program Intelligent Growth 2014-2020. Sara Díaz-Díaz is supported by a PFIS grant from the Instituto de Salud Carlos III (FI18/00086).

We have no transparency declarations to declare.

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