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Phage-Mediated Acquisition of a Type III Secreted Effector Protein Boosts Growth of *Salmonella* by Nitrte Respiration

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**ABSTRACT** Information on how emerging pathogens can invade and persist and spread within host populations remains sparse. In the 1980s, a multidrug-resistant *Salmonella enterica* serotype Typhimurium clone lysogenized by a bacteriophage carrying the *sopE* virulence gene caused an epidemic among cattle and humans in Europe. Here we show that phage-mediated horizontal transfer of the *sopE* gene enhances the production of host-derived nitrate, an energetically highly valuable electron acceptor, in a mouse colitis model. In turn, nitrate fuels a bloom of *S. Typhimurium* in the gut lumen through anaerobic nitrate respiration while suppressing genes for the utilization of energetically inferior electron acceptors such as tetrathionate. Through this mechanism, horizontal transfer of *sopE* can enhance the fitness of *S. Typhimurium*, resulting in its significantly increased abundance in the feces.

**IMPORTANCE** During gastroenteritis, *Salmonella enterica* serotype Typhimurium can use tetrathionate respiration to edge out competing microbes in the gut lumen. However, the concept that tetrathionate respiration confers a growth benefit in the inflamed gut is not broadly applicable to other host-pathogen combinations because tetrathionate respiration is a signature trait used to differentiate *Salmonella* serotypes from most other members of the family *Enterobacteriaceae*. Here we show that by acquiring the phage-carried *sopE* gene, *S. Typhimurium* can drive the host to generate an additional respiratory electron acceptor, nitrate. Nitrate suppresses genes for the utilization of energetically inferior electron acceptors such as tetrathionate while enhancing the luminal growth of *S. Typhimurium* through anaerobic nitrate respiration. Pathways for anaerobic nitrate respiration are widely conserved among members of the family *Enterobacteriaceae*, thereby making our observations relevant to other enteric pathogens whose relative abundance in the intestinal lumen increases during infection.

The first human outbreak of a multidrug-resistant (MDR) pathogen was recorded between 1964 and 1966 in Great Britain (1). The outbreak was caused by a *Salmonella enterica* serotype Typhimurium strain which circulated in cattle and had acquired resistance to furazolidone, ampicillin, chloramphenicol (Cm), kanamycin (Kan), neomycin, streptomycin, sulfonamides, and tetracyclines. Differentiation of *S. Typhimurium* isolates by phage typing identified the outbreak strain as a distinct clone with definitive phage type 29 (DT29) (1). Subsequent surveillance by the Central Veterinary Laboratory in the United Kingdom revealed the emergence of several MDR clones over time. Each epidemic clone, after dominating for a period of time in its bovine animal reservoir, was eventually replaced, as indicated by the dominance of a new phage type (reviewed in reference 2). However, the rise and fall of epidemic *S. Typhimurium* isolates, including phage types DT204, DT204c, and DT49 and laboratory strain SL1344 (4). Based on the association of *SopE* with an epidemic *S. Typhimurium* clone, it has been speculated that acquisition of the SopE protein by phage-mediated horizontal gene transfer might have increased the fitness of the resulting strain in its host population (5). However, experimental evidence for this hypothesis is lacking.

The *sopE* gene encodes a secreted substrate (effector protein) of the invasion-associated type III secretion system (T3SS-1) of *S. Typhimurium* (3, 6). SopE activates several innate immune signaling pathways (6–8) and is a potent inducer of intestinal inflammation (9, 10), but the molecular mechanisms by which SopE might enhance the fitness of an epidemic *S. Typhimurium* clone remain unknown. Here we investigated whether SopE increases the fitness of *S. Typhimurium* in a mouse colitis model and elucidated the underlying mechanism.

**RESULTS**

*SopE* enhances growth in the inflamed intestine. The *sopE* gene increases the severity of intestinal inflammation (8, 10), and this host response enhances the luminal growth of *S. Typhimurium*...
The sopE gene induces iNOS expression and enhances the growth of S. Typhimurium in the intestine. Groups of streptomycin-pretreated mice were infected with the SL1344 wild type (CAL63, sopE+) or a sopE mutant (CAL88, sopE). Groups of mice were euthanized at the indicated time points. For all groups, n = 4 except for the day 2 sopE+ group (n = 3) (a) Analysis of bacterial numbers in the colon contents. Boxes in whisker plots represent the second and third quartiles, while lines indicate the first and fourth quartiles. (B) Nos2 mRNA levels in the mucosa were quantified by real-time PCR. Bars represent geometric mean Nos2 copy numbers per 100 copies of glyceraldehyde 3-phosphate dehydrogenase (Gapdh) mRNA ± the standard errors. (C) Levels of iNOS protein in the mucosa from mice infected with sopE+ (lanes 1 to 4) or sopE mutant (lanes 5 to 8) strains were determined by Western blotting (top panel). Detection of tubulin by Western blotting served as a loading control (bottom panel). Each lane represents protein extracts isolated from one animal. Molecular masses of standard proteins are indicated on the right. (D) Quantification of iNOS levels in Western blot assays by densitometry. Each square represents the intensity of the iNOS signal divided by the intensity of the tubulin signal from one animal. Significance is based on the two-tailed Student t test. *, P < 0.05.
FIG 2 Inhibition of tetrathionate respiration by nitrate and SopE. (A) Mucin broth containing the indicated concentrations of nitrate and tetrathionate was inoculated with an equal mixture of the SL1344 wild-type strain (wt, CAL63) and a ttrA mutant (CAL66). Bacterial numbers were determined after 16 h of anaerobic growth. Bars represent geometric means; the competitive index ± standard errors. Results are averages of three independent experiments. (B) Groups of streptomycin-pretreated mice (n = 4 per group) were infected with the SL1344 wild type (CAL63, sopE+) or a sopE mutant (CAL88, sopE−), and bacterial RNA was extracted from cecal contents 4 days after infection. The relative levels of ttrA mRNA were quantified by real-time PCR, normalized to the abundance of S. Typhimurium 16S rRNA, and expressed as n-fold differences from the ttrA mRNA levels present in the SL1344 wild type (CAL63, sopE+). Bars represent geometric means ± standard errors. Significance is based on the two-tailed Student t test. **, P < 0.01; *, P < 0.05; ns, P > 0.05.

FIG 3 Inactivation of nitrate reductases in S. Typhimurium. (A) Detection of nitrate reductase activity in the S. Typhimurium wild type (SL1344) and a narG narZ napA mutant (CAL59). Briefly, the nitrate reductase assay measures the reduction of nitrate to nitrite with methyl viologen as an electron donor. Nitrate is added to the reaction medium containing lyzed bacterial cells, and nitrite (from nitrate reductase activity) is measured on the basis of its formation of a colored azo compound, which is quantified with a spectrophotometer. (B) Mucin broth was inoculated with an equal mixture of the SL1344 wild-type strain (wt, CAL63) and a narG narZ napA mutant (CAL64). Competitive indices for anaerobic growth in mucin broth with (+) or without (−) nitrate are shown. Bars represent geometric means ± standard errors. The data are from three independent experiments. Significance is based on the two-tailed Student t test. **, P < 0.01.

To further characterize the role of SopE during infection, we analyzed the cecal immune response elicited by S. Typhimurium strain SL1344 in the mouse colitis model. We noticed that the SL1344 wild type (CAL63) induced Nos2 gene expression at significantly higher levels than a sopE mutant (CAL88) at day 3 after the infection of streptomycin-pretreated C57BL/6 mice (Fig. 1B), although the Nos2 mRNA levels in the two groups were similar at day 4 after infection. Nos2 encodes inducible nitric oxide synthase (iNOS), which was present at significantly (P < 0.05) higher levels in the cecal mucosa 3 days after infection with the wild type (CAL63) than after infection with a sopE mutant (CAL88), as shown by immunoblotting (Fig. 1C) followed by quantification by densitometry (Fig. 1D). Thus, the SopE-mediated growth advantage observed 4 days after infection (Fig. 1A) was preceded by a SopE-mediated increase in iNOS expression at day 3 after infection (Fig. 1B to D).

Nitrate is a negative regulator of tetrathionate respiration. The activity of iNOS generates NO, which reacts with reactive oxygen species (ROS) to form peroxynitrite (ONOO−), a bactericidal reactive nitrogen species that quickly isomerizes to harmless nitrate (NO3−) (16). Nitrate is the preferred anaerobic electron acceptor of Salmonella serotypes (17), presumably because the relatively high standard redox potential (E0) of the nitrate-nitrite electron acceptor (17, 19). We therefore reasoned that the intesti- nal growth advantage confered by SopE (Fig. 1A) might be due to anaerobic nitrate respiration. To test this hypothesis, we generated a nitrate respiration-deficient mutant of SL1344. Three operons, including narGHJI, narZYYV, and napFDAGHBC, contribute to nitrate reductase activity in S. Typhimurium (20, 21). We disrupted the narG, narZ, and napA genes in SL1344 to create a mutant (CAL55) that completely lacked respiratory nitrate reductase activity (Fig. 3A). Anaerobic coculture of the Kanr SL1344 wild type (CAL63) with the isogenic Carbr narG narZ napA mutant (CAL64) in mucin broth resulted in enrichment for the wild type when the medium was supplemented with nitrate but not when nitrate was absent (Fig. 3B).

SopE changes ttrA gene expression during infection. One possible scenario to explain the above observations is that SopE-induced expression of iNOS in the cecal mucosa generates nitrate, which in turn suppresses the expression of tetrathionate utilization genes (ttrBAC) in the inflamed gut. To test this idea, we extracted bacterial RNA from the intestinal contents of mice 4 days after infection with the SL1344 wild-type strain (CAL63) or with an isogenic sopE mutant (CAL88) (mouse colitis model). Relative levels of expression of the ttrA mRNA, which encodes the large subunit of the tetrathionate reductase complex, were determined by quantitative real-time PCR and normalized to the abundance of S. Typhimurium 16S rRNA (15). Remarkably, expression of ttrA was significantly lower in the SopE-expressing SL1344 wild type (CAL63) than in the sopE mutant (CAL88) (Fig. 2B). These data suggested that the presence of the sopE gene markedly reduced the expression of ttrA in luminal S. Typhimurium in vivo.

SopE enhances nitrate respiration-dependent growth of S. Typhimurium. S. Typhimurium can utilize nitrate as a terminal electron acceptor (17, 19). We therefore reasoned that the intestinal growth advantage confered by SopE (Fig. 1A) might be due to anaerobic nitrate respiration. To test this hypothesis, we generated a nitrate respiration-deficient mutant of SL1344. Three operons, including narGHJI, narZYYV, and napFDAGHBC, contribute to nitrate reductase activity in S. Typhimurium (20, 21). We disrupted the narG, narZ, and napA genes in SL1344 to create a mutant (CAL55) that completely lacked respiratory nitrate reductase activity (Fig. 3A). Anaerobic coculture of the Kanr SL1344 wild type (CAL63) with the isogenic Carbr narG narZ napA mutant (CAL64) in mucin broth resulted in enrichment for the wild type when the medium was supplemented with nitrate but not when nitrate was absent (Fig. 3B).

To investigate whether nitrate respiration genes confer a growth benefit in the inflamed intestine, we used the mouse colitis model and infected C57BL/6 mice with an equal mixture of the SL1344 wild-type strain (CAL63) and the isogenic, nitrate respiration-deficient mutant (narG narZ napA mutant, CAL64). S. Typhimurium infection resulted in acute intestinal inflammation (Fig. 4A and B) and markedly increased levels of mRNAs for genes that encode proinflammatory cytokines (keratinocyte-
derived chemokine [KC], tumor necrosis factor alpha, and gamma interferon [IFN-\(\gamma\)] (Fig. 4C). Four days after infection, the SL1344 wild-type strain (CAL63) was recovered in higher numbers than the \(\text{narG narZ napA}\) mutant (CAL64) (Fig. 4D). Similar results were obtained in the absence of streptomycin pretreatment when genetically resistant (CBA/J) mice were infected with an equal mixture of CAL63 and CAL64. Outgrowth of the nitrate respiration-competent strain (CAL63) was observed starting day 16 after infection in this model (see Fig. S2 in the supplemental material).

Interestingly, when streptomycin-pretreated C57BL/6 mice were infected with an equal mixture of a Kan’ \(\text{invA spiB}\) mutant (CAL86) and a Carb’ \(\text{sopE narG narZ napA}\) mutant (CAL85). As expected, iNOS expression was detected in wild-type mice but not in iNOS-deficient mice (Fig. 5A) but both groups exhibited similarly severe pathological changes 4 days after infection (Fig. 5B and C). While a growth advantage of the wild type (CAL63) over the \(\text{narG narZ napA}\) mutant (CAL64) was observed in wild-type mice, no enrichment for the S. Typhimurium wild-type strain was observed in \(\text{Nos2}\)-deficient mice (Fig. 5D; \(P < 0.01\)). These data suggested that host-derived

**FIG 4** The \textit{sopE} gene boosts nitrate respiration-dependent growth \textit{in vivo}. Groups of streptomycin-pretreated mice (\(n = 4\) per group) were inoculated with sterile medium (mock) or with equal mixtures of S. Typhimurium SL1344 derivatives (relevant genotypes are indicated), and organs were collected for analysis 4 days after infection. (A) Blinded histopathology scoring of cecal inflammation showing average scores (bars) and scores of individual animals (squares). (B) Representative images of sections from the cecal mucosa. (C) Expression of proinflammatory markers in the cecal mucosa was determined by quantitative real-time PCR analysis. Bars represent geometric mean KC-, tumor necrosis factor alpha-, and IFN-\(\gamma\)-encoding (\(\text{Kc, Tnfa, and Ifng}\), respectively) mRNA copy numbers per 100,000 copies of glyceraldehyde 3-phosphate dehydrogenase (\(\text{GAPDH}\)) mRNA ± the standard errors. (D) Competitive indices of S. Typhimurium strains recovered from the colon contents of infected mice. Bars represent geometric means ± standard errors. Significance is based on the two-tailed Student \(t\) test. **, \(P < 0.01\).
nitrate supports the nitrate respiration-dependent luminal growth of S. Typhimurium.

**Horizontal transfer of the SopEΦ prophage enhances growth by nitrate respiration.** SopE is encoded by a mobile genetic element, lysogenic SopEΦ, that can be transferred horizontally between different strains (5, 25). To recapitulate phage-mediated horizontal transfer of the sopE gene, we determined how intestinal growth of S. Typhimurium IR715 is altered by lysogenization with SopEΦ. We infected mice with an equal mixture of IR715 with an isogenic nitrate respiration-deficient mutant (IR715 narG narZ napA mutant, CAL50) (mouse colitis model). Four days after infection, mice exhibited acute intestinal inflammation with SopE is a T3SS-1 effector protein (26) that enhances intestinal growth by nitrate respiration, we repeated the experiment after deleting the sopE gene. Mice were infected with an equal mixture of IR715 carrying a SopEΦ prophage derivative in which the sopE gene had been deleted (IR715 SopEΦΔsopE mutant, CAL102) and an isogenic nitrate respiration-deficient mutant (IR715 SopEΦΔsopE narG narZ napA mutant, CAL103) (mouse colitis model). Deletion of the sopE gene (CAL102 versus CAL103) reduced the luminal growth advantage conferred by nitrate respiration to levels observed in strains lacking the SopEΦ prophage (IR715 versus, CAL50) (Fig. 6D). The reduced competitive advantage was not due to lessened inflammation, as both the histopathology scores and levels of inflammatory cytokines of the groups were similar (Fig. 6A to C).

Taken together, these results indicated that SopEΦ-mediated horizontal transfer of the sopE gene could increase the luminal abundance of S. Typhimurium by boosting growth through nitrate respiration.

**DISCUSSION**

SopE is a T3SS-1 effector protein (26) that enhances intestinal inflammation during S. Typhimurium infection (9, 10) by activating caspase 1, which in turn cleaves the immature forms of interleukin-1β (IL-1β) and IL-18 into biologically active cytokines (7, 8). During S. Typhimurium infection, IL-18 contributes to the early production of IFN-γ (27), a cytokine that is a potent inducer of iNOS expression (28–31). Here we show that the presence of the sopE gene resulted in a significant increase in mucosal iNOS production 3 days after S. Typhimurium infection in the mouse colitis model. NO generated by iNOS can react with ROS to form peroxynitrite (ONOO−), which isomerizes to nitrate
(NO$_3^-$) (16), a terminal electron acceptor that can promote the growth of S. Typhimurium by anaerobic nitrate respiration (17, 19). The elevated mucosal iNOS production observed 3 days after infection was followed the next day by a nitrate respiration-dependent luminal growth advantage of sopE-positive S. Typhimurium. This nitrate respiration-dependent luminal growth advantage was abrogated in iNOS-deficient mice. The picture emerging from these data is that SopE-dependent iNOS expression generates host-derived nitrate, which in turn boosts the luminal growth of S. Typhimurium in the inflamed intestine (Fig. 7).

The sopE gene is carried by a bacteriophage, termed SopEΦ (3), that is absent from most S. Typhimurium isolates (4). The presence of SopEΦ in an S. Typhimurium clone that caused an epidemic in cattle and humans in Europe in the 1980s suggests that the fitness of this epidemic clone might have been increased by phage-mediated horizontal transfer of the sopE gene (5). Here we show that transfer of SopEΦ into a SopE-negative S. Typhimurium isolate (ATCC 14028) increased its fitness by conferring a nitrate respiration-dependent luminal growth advantage.

In Escherichia coli, genes for anaerobic respiration are subject to a hierarchical control that ensures that the respiratory electron acceptor with the most positive $E_0$ is used preferentially (32–38). The nitrate-nitrite redox couple is a terminal electron acceptor with a higher $E_0$ (433 mV) (39) than the tetrathionate-thiosulfate couple ($E_0 = 170$ mV) (18). Here we show that nitrate is a negative regulator of anaerobic tetrathionate respiration in vitro. Furthermore, the presence of the sopE gene significantly reduced the expression of tetrathionate respiration genes in the luminal S. Typhimurium population in vivo. This hierarchical control of genes involved in anaerobic respiration might ensure that the energetically preferred terminal electron acceptor is used preferentially to maximize the growth rate in the competitive environment of the large bowel.

The observation that tetrathionate provides a growth advantage in the inflamed intestine (40) is not germane to host interactions of the majority of the members of the family Enterobacteriaceae, because tetrathionate respiration is a signature trait historically used to differentiate Salmonella serotypes from most other bacteria (41). In contrast, the concept that nitrate respiration can provide a growth benefit during intestinal inflammation is relevant to other host-microbe interactions, since nitrate reduc-
and stored at 4°C. To transfer mutations between strains, diluted phage lysate containing the mutation of interest was incubated for 1 h at room temperature with an overnight culture of the recipient bacterial strain. The bacteria were then spread onto plates with the appropriate antibiotic. Resulting colonies were streaked onto Evans blue uranine agar plates (10 g/liter tryptone, 5 g/liter yeast extract, 5 g/liter NaCl, 2.5 g glucose, 15 g/liter Difco agar, 0.00125% Evans blue solution, 0.5% K$_2$HPO$_4$, 0.0025% sodium fluorescein) to select for nonlysogenic colonies. The sensitivity of the resulting strains to reinfection with P22 was determined by cross-streaking of the bacterial strain with P22 H5.

**Construction of S. Typhimurium mutants.** To construct the napA mutant CAL46, as well as the narZ mutant CAL40, approximately 1-kb regions upstream and downstream of either napA or narZ were PCR amplified (see Table S2 in the supplemental material) and digested with XbaI (New England Biolabs). The two regions flanking each target gene were ligated and PCR amplified to obtain an approximately 2-kb region that was inserted into pCR2.1 with a TOPO cloning kit (Invitrogen), creating plasmids pCAL3 and pCAL1, respectively. The insertions were then sequenced. The plasmids were next digested with SpI to excise the napA or narZ flanking regions, which were then inserted into suicide plasmid pRDH10 to obtain pCAL9 and pCAL10, respectively. These plasmids were propagated in *E. coli* DH5α *apir*. A Kan^r^ cassette from pUCKSAC was inserted into the Xbal restriction site between the flanking regions to create pCAL11 and pCAL12, respectively. *E. coli* S17-1 *apir* was transformed with the resulting plasmids and subsequently used for conjuga-
sion. Single homologous recombination events were selected for by growth on Kan and Cm plates. To select for strains that had lost the integrated plasmid, leaving the Kan^r^ sequence in place of the target gene, we used sucrose selection. For sucrose selection, overnight cultures of bacteria were grown in LB broth without antibiotics, digested, spread onto sucrose plates (5% sucrose, 8 g/liter nutrient broth, 15 g/liter agar), and then incubated for 16 h at 30°C. Sucrose-sensitive and Kan^r^ Cm^r^ strains in which the napA or narZ gene had been replaced with the Kan^r^ cassette were termed CAL42 and CAL34, respectively. To generate nonpolar mu-
tants, *E. coli* S17-1 *apir* was transformed with either pCAL9 or pCAL10 and conjugated with CAL42 and CAL34, respectively. Sucrose selection was performed as described above, and Kan^r^ Cm^r^ strains in which the napA or narZ gene had been deleted were named CAL46 and CAL40, respectively. Nonpolar mutations were confirmed by PCR. The ∆napA and ∆narZ mutations were transiently marked with Cm^r^ by reintroducing pCAL9 or pCAL10 into the chromosome of CAL46 and CAL40 to create CAL117 and CAL45, respectively.

To construct the narG mutant CAL27, an approximately 500-bp region within the narG sequence was PCR amplified and digested with the restriction enzymes XbaI and SpI. The digested fragment was subsequently cloned into pGPT04 and propagated in *E. coli* S17-1 *apir*. The resulting plasmid, pCAL5, was introduced into the IR715 chromosome by conjugation, selecting for single homologous recombination events by plating on plates containing Carb. Mutations were confirmed via PCR.

To generate a nonpolar deletion in *spiB*, *E. coli* S17-1 *apir* was transformed with pSPN56 and conjugated with SPN450. Integration of pSPN56 into SPN450 was selected for by plating on LB containing Cm and Nal. Sucrose selection was used to select for the loss of pSPN56. Colonies were screened for sensitivity to Cm and Kan, and a colony exhibiting both phenotypes was confirmed by PCR to contain an unmarked ∆*spiB* locus, yielding SPN456. To enable transduction of the unmarked *spiB* deletion, pSPN56 was integrated into SPN456 by conjugation as described above, resulting in a merodiploid ∆*spiB* state flanking the integrated plasmid, a state that was maintained by selection for Cm^r^, a transconjugant exhibiting Cm^r^ and loss of the PCR product from unmarked ∆*spiB* amplification was termed SPN458 (∆*spiB*:pSPN56).

To construct the sopE mutant SW976, DNA regions upstream and downstream of sopE were PCR amplified and digested with XbaI. The two flanking regions were ligated, PCR amplified, and cloned into pRDH10 using BamHI restriction enzyme sites to create pSW245. To create

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**MATERIALS AND METHODS**

**Bacterial culture conditions.** For the *Escherichia coli* and *S. Typhimurium* strains used in this study, see Table S1 in the supplemental material. Strains were routinely grown in LB broth (10 g/liter tryptone, 5 g/liter yeast extract, 10 g/liter) or on LB plates (15 g/liter agar) with the appropriate antibiotic at the following concentrations: Carb, 0.1 mg/ml; Cm, 0.03 mg/ml; Kan, 0.1 mg/ml; Nal, 0.05 mg/ml; tetracycline (Tet), 0.02 mg/ml.

**Transduction using phage P22.** Once single mutations were obtained in an *S. Typhimurium* strain, phage lysates were generated. We incubated 1 ml of an overnight culture of the strain of interest with 4 ml of P22 broth (LB broth supplemented with 1 × E minimal medium [46], 0.2% glucose, and 10$^{10}$ to 10$^{11}$ PFU/ml P22 HT-int) overnight at 37°C. Debris was removed by centrifugation at 10,000 × g for 5 min, and the phage-containing supernatant was mixed with chloroform (20% final volume) and stored at 4°C. To transfer mutations between strains, diluted phage lysate containing the mutation of interest was incubated for 1 h at room temperature with an overnight culture of the recipient bacterial strain. The bacteria were then spread onto plates with the appropriate antibiotic. Resulting colonies were streaked onto Evans blue uranine agar plates (10 g/liter tryptone, 5 g/liter yeast extract, 5 g/liter NaCl, 2.5 g glucose, 15 g/liter Difco agar, 0.00125% Evans blue solution, 0.5% K$_2$HPO$_4$, 0.0025% sodium fluorescein) to select for nonlysogenic colonies. The sensitivity of the resulting strains to reinfection with P22 was determined by cross-streaking of the bacterial strain with P22 H5.

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To construct the sopE mutant SW976, DNA regions upstream and downstream of sopE were PCR amplified and digested with XbaI. The two flanking regions were ligated, PCR amplified, and cloned into pRDH10 using BamHI restriction enzyme sites to create pSW245. To create

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**FIG 7** Model of the chain of events by which SopE boosts nitrate respiration-dependent growth of *S. Typhimurium* in the inflamed intestine. For an explanation, see the text. DUOX2, dual-oxidase 2 of epithelial cells; NOX1, NADPH oxidase 1 of epithelial cells; PHOX, NADPH oxidase of phagocytes.
pSW246, a Kan’ cassette was cloned into the XbaI site between the two sopE flanking regions in pSW245. These plasmids were conjugated from E. coli S17-1 Apir into S. Typhimurium. Plasmid integration into the chromosomal was verified by PCR. The sopE gene in SL1344 was deleted by introducing pSW246 into SL1344(pSW172) by conjugation, selecting for exconjugants on LB plates containing Carb and Kan. To ensure stable replication of the temperature-sensitive plasmid pSW172, conjugation experiments involving SL1344(pSW172) were performed at 30°C. pSW172 was subsequently cured by growth at 37°C. A Kan’ Cmr’ strain indicated that sopE had been replaced with Kan’ and was termed SW975.

To create a nonpolar deletion of sopE, pSW245 was integrated into the chromosome of SW975 by conjugation, selecting for concomitant Kan’ and Cmr’. Sucrose selection was then used to obtain a Kan’ Cmr’ strain named SW976. The ΔsopE deletion was marked with Cmr’ by conjugating SW976 with E. coli S17-1 Apir containing pSW245. The Cmr’ strain was termed CAL74.

The narG::pCAL5, ΔnarZ::pCAL10, and ΔnapA::pCAL9 mutations were transduced from CAL27, CAL45, and CAL117, respectively, into IR715. pCAL10 and pCAL9 were removed via sucrose selection, and the resulting strain was designated CAL50. The ΔnapA::Tetr and ΔspiB:: pSPN56 mutations were transduced from SW562 and SPN458, respectively, into CAL50, pSPN56 was removed via sucrose selection, and the resulting strain was designated CAL87. The sopE::pCAL9 mutations from CAL50 into SL1344. The narG, napA, and napZ mutations from CAL27, CAL45, and CAL117 were transduced into SL1344 and CAL63, and plasmids pCAL10 and pCAL9 were removed via sucrose selection to yield either mutants defective for a single nitrate reductase (designated CAL65, CAL51, and CAL67, respectively) or nitrate reductase-deficient mutants (CAL59 and CAL64). The invA::Tet and invA::spiB::pSPN56 mutations were transduced from SW562 and SPN458 into strains CAL63 and CAL64, pSPN56 was removed via sucrose selection, and the resulting strains were designated CAL86 and CAL85, respectively. Additionally, the sopE:: pSW245 mutation was transduced from CAL74 into strains CAL63, CAL64, CAL98, and CAL99 and the integrated plasmid pSW245 was removed via sucrose selection to create strains CAL68, CAL89, CAL102, and CAL103, respectively. The trtA::pSW171 mutation was transduced from SW661 into CAL63 to yield CAL66.

Respiratory nitrate reductase assay. Bacterial overnight cultures were diluted 1:100 in LB broth at a final concentration of 108 CFU/mouse in 0.1 ml LB broth. Fecal pellets were collected between 1 and 4 days after infection as indicated. Cecal tissue and cecal contents were flash frozen and stored at −80°C. Colon contents were stored in phosphate-buffered saline (PBS) on ice. CFU counts were determined by plating dilutions of the inoculum or colon content on selective medium.

In the model of extended S. Typhimurium infection, wild-type CBA/J mice (Jackson Laboratories) were intragastrically provided an inoculum containing a 1:1 ratio of competing strains at a final concentration of 1 × 106 CFU/mouse in 0.1 ml LB broth. Fecal pellets were collected between 1 and 28 days postinfection as indicated and stored in PBS on ice. At the end of the experiment, mice were euthanized and colon contents were removed and stored in PBS on ice. CFU counts were determined by plating dilutions of the inoculum, fecal pellets, or colon content on selective medium.

Histopathology. The distal segment of the cecum was fixed in 10% buffered formalin and stained with hematoxylin and eosin. A veterinary pathologist scored histopathological changes by blinded scoring of sections by a scheme described previously (48).

Isolation of cecal RNA and proteins. Tissue was homogenized in a Mini-Beadbeater (BioSpec Products, Bartlesville, OK), and RNA and protein were isolated by the TRI-Reagent method (Molecular Research Center, Inc.) following the manufacturer’s protocol. RNA remaining in the RNA isolation portion was removed using the DNA-free kit (Applied Biosystems). RNA samples and protein extracts were stored at −80°C or −20°C, respectively.

Quantification of mRNA levels. Isolated RNA was normalized to 100 ng/μl and reverse transcribed using random hexamers and Moloney murine leukemia virus reverse transcriptase (Applied Biosystems). Quantitative real-time PCR was performed using SYBR green (Applied Biosystems) PCR mix and the appropriate primer sets (see Table S2 in the supplemental material) at a final concentration of 0.25 mM. In cases where n-fold changes were calculated, relative expression was compared to either mock-infected cecal tissue for murine gene expression experiments or to SopE-positive S. Typhimurium for bacterial gene expression analysis. In cases where absolute values were calculated, a standard curve was used. To create the standards, first we used a standard PCR to amplify the target gene fragment with the quantitative reverse transcription (qRT)-PCR primers and inserting it into pCR2.1 by using the TOPO cloning kit. The resulting plasmids were sequenced. The plasmids were quantified to create a set of standards ranging from 100 to 108 copies/μl diluted in a 0.02-mg/ml yeast RNA (Sigma) solution. Each qRT-PCR was performed in duplicate.

Analysis of cecal proteins. The concentration of mouse cecal protein was measured using a Micro BCA protein assay kit (Pierce) in accordance with the manufacturer’s instructions. A 0.05-μg sample of protein was separated by SDS-PAGE and transferred to a polyvinylidene fluoride membrane (Millipore). Three percent nonfat dried milk and 0.1% Tween 20 (Bio-Rad) in PBS solution were used as blocking agents. To detect iNOS, a 1:1,500 dilution of the primary antibody (anti-mouse; BD Transduction Laboratories) in PBS with 5% bovine serum albumin (BSA) was added to the membrane. As a loading control, α-β-tubulin was detected with a 1:5,000 dilution of the primary antibody (anti-rabbit; Cell Signaling) in PBS with 5% BSA. The horseradish peroxidase (HRP)-conjugated secondary antibody (Bio-Rad), either anti-mouse or anti-rabbit, was diluted 1:5,000 in blocking buffer and applied to the membrane. Protein bands were visualized by applying 0.5 ml HRP substrate solution and imaged with a UVP BioSpectrum Imaging System. Blot density quantification was performed with the Labworks Image Acquisition and Analysis Software (UVP version 4.6.00.0).

Statistics. The natural log of the data values was calculated before statistical analysis. We used Student’s t test to assess significance and considered P values below 0.05 to be significant.
SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at http://mbio.asm.org/lookup/suppl/doi:10.1128/mBio.00143-12/-/DCSupplemental.

Figure S1, PDF file, 0.1 MB.
Figure S2, PDF file, 0.1 MB.
Table S1, PDF file, 0.1 MB.
Table S2, PDF file, 0.1 MB.

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