



# The Type III Secretion System Effector SptP of *Salmonella enterica* Serovar Typhi

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**ABSTRACT** Strains of the various *Salmonella enterica* serovars cause gastroenteritis or typhoid fever in humans, with virulence depending on the action of two type III secretion systems (*Salmonella* pathogenicity island 1 [SPI-1] and SPI-2). SptP is a *Salmonella* SPI-1 effector, involved in mediating recovery of the host cytoskeleton postinfection. SptP requires a chaperone, SicP, for stability and secretion. SptP has 94% identity between *S. enterica* serovar Typhimurium and *S. Typhi*; direct comparison of the protein sequences revealed that *S. Typhi* SptP has numerous amino acid changes within its chaperone-binding domain. Subsequent comparison of  $\Delta sptP$  *S. Typhi* and *S. Typhimurium* strains demonstrated that, unlike SptP in *S. Typhimurium*, SptP in *S. Typhi* was not involved in invasion or cytoskeletal recovery postinfection. Investigation of whether the observed amino acid changes within SptP of *S. Typhi* affected its function revealed that *S. Typhi* SptP was unable to complement *S. Typhimurium*  $\Delta sptP$  due to an absence of secretion. We further demonstrated that while *S. Typhimurium* SptP is stable intracellularly within *S. Typhi*, *S. Typhi* SptP is unstable, although stability could be recovered following replacement of the chaperone-binding domain with that of *S. Typhimurium*. Direct assessment of the strength of the interaction between SptP and SicP of both serovars via bacterial two-hybrid analysis demonstrated that *S. Typhi* SptP has a significantly weaker interaction with SicP than the equivalent proteins in *S. Typhimurium*. Taken together, our results suggest that changes within the chaperone-binding domain of SptP in *S. Typhi* hinder binding to its chaperone, resulting in instability, preventing translocation, and therefore restricting the intracellular activity of this effector.

**IMPORTANCE** Studies investigating *Salmonella* pathogenesis typically rely on *Salmonella* Typhimurium, even though *Salmonella* Typhi causes the more severe disease in humans. As such, an understanding of *S. Typhi* pathogenesis is lacking. Differences within the type III secretion system effector SptP between typhoidal and nontyphoidal serovars led us to characterize this effector within *S. Typhi*. Our results suggest that SptP is not translocated from typhoidal serovars, even though the loss of *sptP* results in virulence defects in *S. Typhimurium*. Although SptP is just one effector, our results exemplify that the behavior of these serovars is significantly different and genes identified to be important for *S. Typhimurium* virulence may not translate to *S. Typhi*.

**KEYWORDS** *Salmonella enterica* serovar Typhi, SptP, type III secretion system

*Salmonella* species are diverse Gram-negative intracellular pathogens responsible for a range of diseases resulting in significant morbidity and mortality among both animals and humans worldwide (1). In humans, the outcome of infection with *Salmonella* primarily depends on the infecting serovar; while nontyphoidal serovars, such as

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*Salmonella enterica* serovar Typhimurium, typically cause self-limiting gastroenteritis, typhoidal serovars, such as *Salmonella enterica* serovar Typhi and *Salmonella enterica* serovar Paratyphi, result in the invasive systemic disease typhoid fever (1).

Central to *Salmonella* virulence is the action of two type III secretion systems (T3SS), encoded on *Salmonella* pathogenicity island 1 (SPI-1) and SPI-2, which secrete over 40 effectors to subvert host cell processes during infection (2–4). The SPI-1 T3SS is active when *Salmonella* is extracellular, where it functions to permit the invasion of nonphagocytic host cells (4), while the SPI-2 T3SS is activated upon internalization, where it functions to create a stable and permissive intracellular niche, termed the *Salmonella*-containing vacuole (SCV) (3, 4).

Although *S. Typhi* causes a more serious disease in humans, *S. Typhimurium* is often used as the model for understanding *Salmonella* pathogenesis, owing to biosafety concerns and, most importantly, the availability of viable *in vivo* models, since *S. Typhi* and other typhoidal serovars are strictly restricted to humans (5). Despite the widespread acceptance of *S. Typhimurium* as the *Salmonella* model organism, significant genomic differences exist between *S. Typhimurium* and *S. Typhi*. Comparison of *S. Typhimurium* strain LT2 to *S. Typhi* reference strain CT18 revealed that 89% of genes were shared; approximately 480 genes were unique to LT2 and approximately 600 genes were unique to CT18 (6). Significant variations in prophage, pathogenicity island, and plasmid elements were observed between the two serovars (6).

Importantly, many known SPI-1 and SPI-2 T3SS effectors are absent or pseudogenes in *S. Typhi*, and this may be associated with genome degradation linked with host restriction (6) and could contribute to the differences in pathogenicity observed between typhoidal and nontyphoidal serovars. A notable example is the *S. Typhimurium* effector GtgE, which is absent in *S. Typhi* (6); the introduction of GtgE into *S. Typhi* permits its growth in nonpermissive mouse cells, strongly linking the absence of this effector to typhoidal host restriction (7). Another example which highlights the limitations of using *S. Typhimurium* to investigate *Salmonella* pathogenesis is the fact that the *Salmonella* virulence plasmid (pSLT), encoding the effectors SpvB and SpvC, is required for full *S. Typhimurium* virulence in mice (8) but absent in *S. Typhi* and other typhoidal serovars (5, 9). Additional effectors that are either absent or pseudogenes in *S. Typhi* (CT18) include SopA, CigR, SopE2, SlrP, SseJ, SopD2, AvrA, SteB, GogB, Ssel/SrfH, SseK1, SseK2, SseK3, and SspH1 (6, 10). Other effectors appear to be differentially evolved between human-restricted and generalist serovars, including SipD, SseC, SseD, SseF, SifA, and SptP (11), which could reflect functional differences. Further evidence of the degradation of the T3SS effector repertoire has been reported within the highly successful and globally dominant *S. Typhi* haplotype 58 (H58), which, owing to a premature stop codon at position 185, lacks both the GTPase-activating phosphatase (GAP) and tyrosine phosphatase domains of SptP and therefore lacks a functional copy of this effector (12).

SptP is an SPI-1 effector encoded within the SPI-1 pathogenicity island immediately downstream of its chaperone, SicP (13). Like other effectors, SptP requires its chaperone for stability within the bacterial cytosol and to direct secretion (13, 14). SptP is a modular protein consisting of three distinct domains: a chaperone-binding domain, a GAP domain, and a tyrosine phosphatase domain. SptP is best characterized in its role as a GTPase-activating protein, acting antagonistically to another SPI-1 effector, SopE/SopE2, a guanine nucleotide exchange factor (GEF) which manipulates the host cytoskeleton to permit membrane ruffle formation and *Salmonella* internalization (2, 15). SptP deactivates the Rho GTPases Rac1 and Cdc42 to reverse cytoskeletal changes and return the host membrane to a steady-state condition (16). Importantly, extended activation of Rho GTPases by SopE can be sensed by host cells by NOD1, resulting in NF- $\kappa$ B activation and proinflammatory signaling (17); antagonism of SopE by SptP therefore represents an important mechanism to prevent the activation of host immune defenses. SptP has also been reported to directly inhibit the activation of the mitogen-activated protein kinase pathway and subsequently downregulate tumor necrosis factor alpha secretion from infected J774A.1 macrophages (18). The less well

characterized tyrosine phosphatase domain of SptP has been shown to be involved in SCV biogenesis and bacterial replication up to 8 h postinfection (19). SptP is also important for virulence *in vivo*, as a  $\Delta sptP$  *S. Typhimurium* SL1344 strain demonstrates attenuated virulence in BALB/c mice (20) and disruption of *sptP* in *S. Typhimurium* ST4/74 also results in reduced intestinal colonization of chicks, pigs, and cattle (21).

Most of the characterization of SptP has been performed with *S. Typhimurium* (16, 22–24), with the underlying assumption being that *S. Typhi* behaves similarly. The phenotypes attributed to SptP in *S. Typhimurium*, in particular, the virulence defects reported during *in vivo* infection, would seemingly contradict the global success of H58 *S. Typhi* strains, which lack a functional copy of SptP (12). This suggests that the function of this effector in *S. Typhi* may be different from that reported in *S. Typhimurium*.

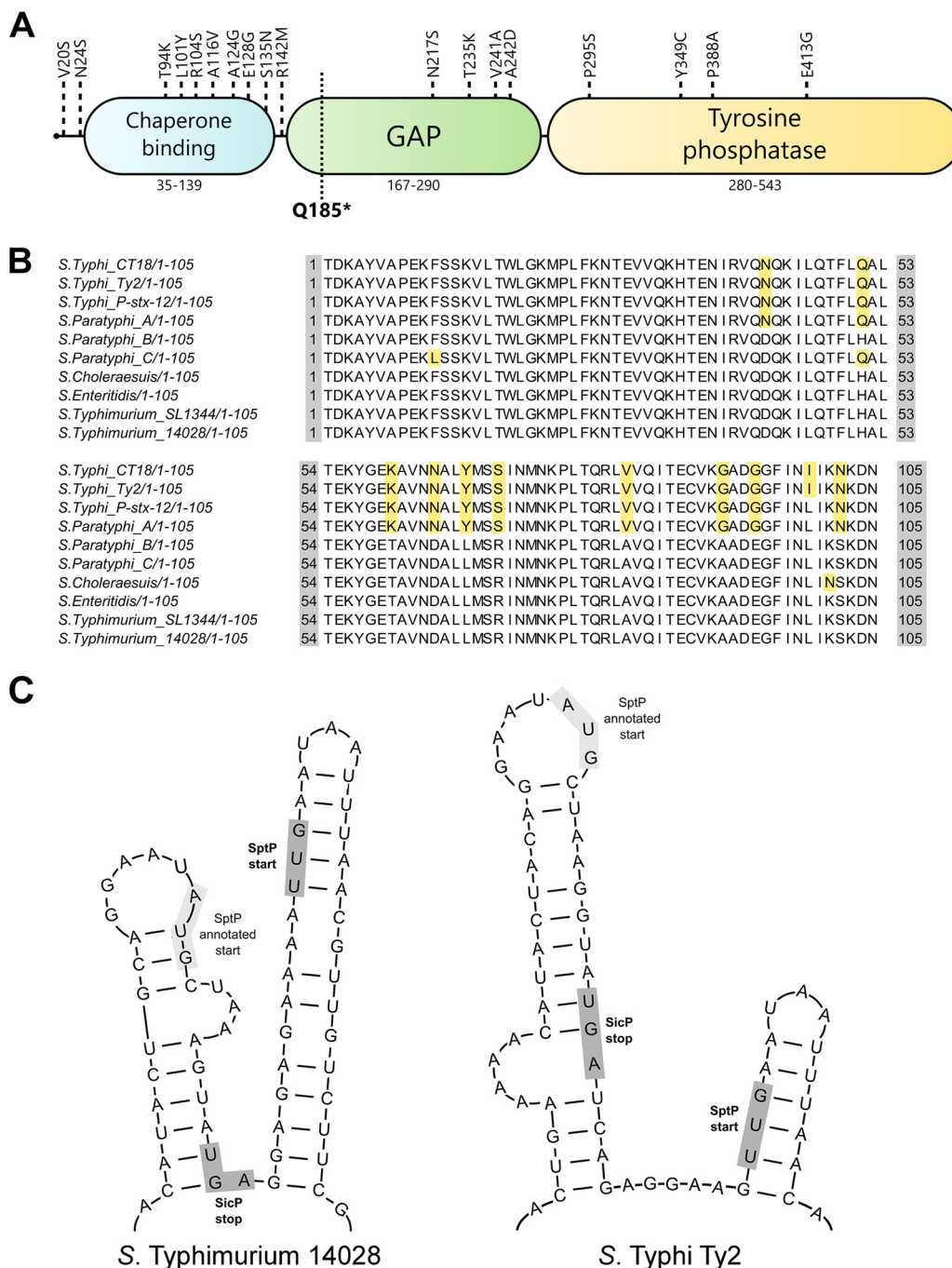
## RESULTS

**Comparison of SptP from different serovars.** SptP is relatively conserved between *S. Typhimurium* and *S. Typhi*, sharing 94% amino acid sequence identity between the two serovars. Alignment of the amino acid sequences of SptP from *S. Typhi* Ty2 and SptP from *S. Typhimurium* 14028 revealed that a large number of amino acid changes (10/31) are concentrated within the chaperone-binding domain of SptP (residues 35 to 139 [25]) (Fig. 1A), despite SicP being highly conserved between serovars (97% identity). Direct comparison of the chaperone-binding domains of SptP from various *Salmonella* serovars shows that these amino acid changes are conserved between *S. Typhi* and *S. Paratyphi* A and that these amino acids are distinct from those in nontyphoidal serovars, including *S. Typhimurium*, *S. Enteritidis*, and *S. Choleraesuis* (Fig. 1B).

Further differences between *sptP* of *S. Typhimurium* and *S. Typhi* can be found outside the protein-coding sequence. It has previously been demonstrated in *S. Typhimurium* that efficient translation of SptP following transcription from its endogenous locus depends on disruption of a stem-loop approximately 20 nucleotides upstream of the start codon of SptP, which is mediated by the upstream translation of SicP (23). Elongation of this stem-loop by just 3 nucleotides decreases the level of translation of SptP by approximately 80% (23). Comparison of the predicted RNA structure (by use of the Mfold server) (26) of this region between the two serovars reveals that *S. Typhi* possesses an elongated stem-loop structure relative to the sequence of *S. Typhimurium* (40 bp and 27 bp, respectively) (Fig. 1C), suggesting that the expression of SptP at the level of translation may differ between the two serovars. Given these differences, we investigated if SptP in *S. Typhi* has the same functions reported during *S. Typhimurium* infection.

**The role of SptP in *Salmonella* invasion and cytoskeletal recovery.** In *S. Typhimurium*, the loss of *sptP* abolishes the ability of infected cells to recover the host cytoskeleton, with cells displaying extended actin disruption up to 3 h postinfection; conversely, cells infected with wild-type (WT) *S. Typhimurium* regain a normal cytoskeletal structure as soon as 80 min postinfection (16). To assess if the same phenotype was seen during *S. Typhi* infection, we compared the ability of HeLa cells to regain a normal actin cytoskeleton following infection with WT and  $\Delta sptP$  *S. Typhi* strains. The equivalent *S. Typhimurium* strains were used as controls.

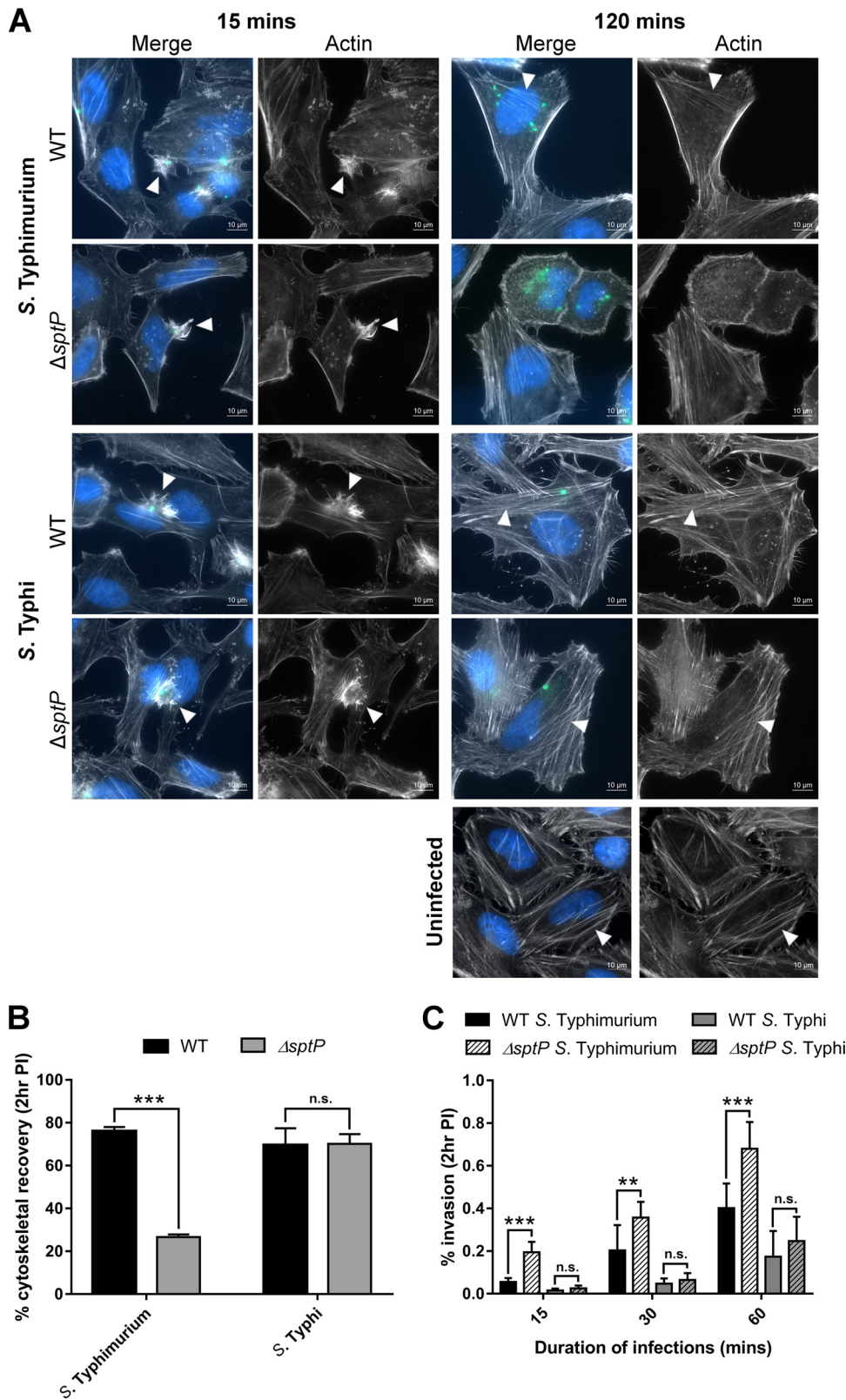
During epithelial cell infection, both *S. Typhimurium* and *S. Typhi* manipulate the host cytoskeleton, with membrane ruffles associated with invasion events clearly being visible at 15 min postinfection (Fig. 2A). By 2 h, cells infected with WT *Salmonella* display an actin architecture similar to that seen for uninfected cells, visible as the formation of stress fibers (Fig. 2A). In line with previous findings (16), cells infected with *S. Typhimurium*  $\Delta sptP$  did not demonstrate cytoskeletal recovery and continued to exhibit membrane ruffling and an absence of stress fibers. In contrast, cells infected with *S. Typhi*  $\Delta sptP$  displayed a recovered cytoskeleton at 2 h postinfection. To quantify this phenotype, we determined the proportion of infected cells displaying an actin architecture akin to that of uninfected cells at 2 h postinfection. While 76% of cells infected with WT *S. Typhimurium* ( $n = 309$ ) demonstrated cytoskeletal recovery, only 27% of



**FIG 1** SptP of *Salmonella* Typhi. (A) Schematic of SptP showing amino acid changes (weakly similar or dissimilar amino acids) between *S. Typhimurium* 14028 and *S. Typhi* Ty2. The location of the single nucleotide polymorphism (Q185\*) which generates a premature stop codon in H58 isolates is also shown. (B) Alignment of the sequence of the chaperone-binding domain (positions 35 to 139) of SptP from various *Salmonella* serovars. Nonconserved amino acids are highlighted. (C) Comparison of the mRNA structures upstream of the start codon of SptP present in *S. Typhimurium* and *S. Typhi*, predicted by use of the Mfold server. Both the annotated ATG (AUG) start codon and the proposed TTG (UUG) start codon of SptP are highlighted.

cells infected with *S. Typhimurium*  $\Delta$ sptP ( $n = 315$ ) demonstrated a recovered actin cytoskeleton. In comparison, 70% of cells infected with either WT *S. Typhi* ( $n = 294$ ) or *S. Typhi*  $\Delta$ sptP ( $n = 301$ ) exhibited cytoskeletal recovery (Fig. 2B).

Although SptP has not previously been linked with invasion in *S. Typhimurium* (20, 27), SptP, via its antagonism of SopE, should directly govern the size and duration of membrane ruffling and therefore influence the efficiency of invasion. As such, we



**FIG 2** Role of SptP in invasion and cytoskeletal recovery following exponential growth. *Salmonella* strains, grown aerobically to late exponential phase, were used to infect HeLa cells at an MOI of 100. (A) Representative images of HeLa cells infected with WT or  $\Delta sptP$  *S. Typhimurium* or *S. Typhi* at 15 min and 2 h postinfection. Actin was stained with phalloidin (white), bacteria were stained with the CSA-1 antibody (green), and nuclei were stained with Hoechst (blue). Arrowheads, invasion-associated membrane ruffles (15 min) or stress fibers (120 min). (B) HeLa cells infected with *Salmonella* strains were fixed at 2 h postinfection (PI) and stained with phalloidin to determine the

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compared the levels of invasion of WT and  $\Delta sptP$  *S. Typhimurium* and *S. Typhi* strains into HeLa cells. While deletion of *sptP* from *S. Typhimurium* resulted in a significantly increased level of invasion relative to that of the WT strain, deletion of *sptP* from *S. Typhi* did not affect the invasion efficiency (Fig. 2C). Increasing the length of time that *Salmonella* was incubated with host cells improved the invasion efficiency of all strains but did not alter the trends observed (Fig. 2C). SPI-1-deficient *S. Typhimurium* ( $\Delta prgH$ ) and *S. Typhi* ( $\Delta invA$ ) strains, used as controls, failed to invade HeLa cells, as expected (see Fig. S1 in the supplemental material).

**Growth under different SPI-1-inducing conditions influences SopE activity in *S. Typhi*.** A recent study comparing *S. Paratyphi* A and *S. Typhimurium* following growth under different SPI-1-inducing conditions demonstrated that *S. Paratyphi* was significantly less invasive, had reduced levels of SPI-1 expression, and had reduced levels of SptP and SopE2 following aerobic growth to late exponential phase compared to the results obtained by microaerobic growth to stationary phase (28). Since our initial comparisons between WT and  $\Delta sptP$  *Salmonella* strains were performed following aerobic exponential growth (Fig. 2), we were interested in whether SopE, the guanine exchange factor which SptP antagonizes (16), was active during *S. Typhi* infection after growth under these conditions.

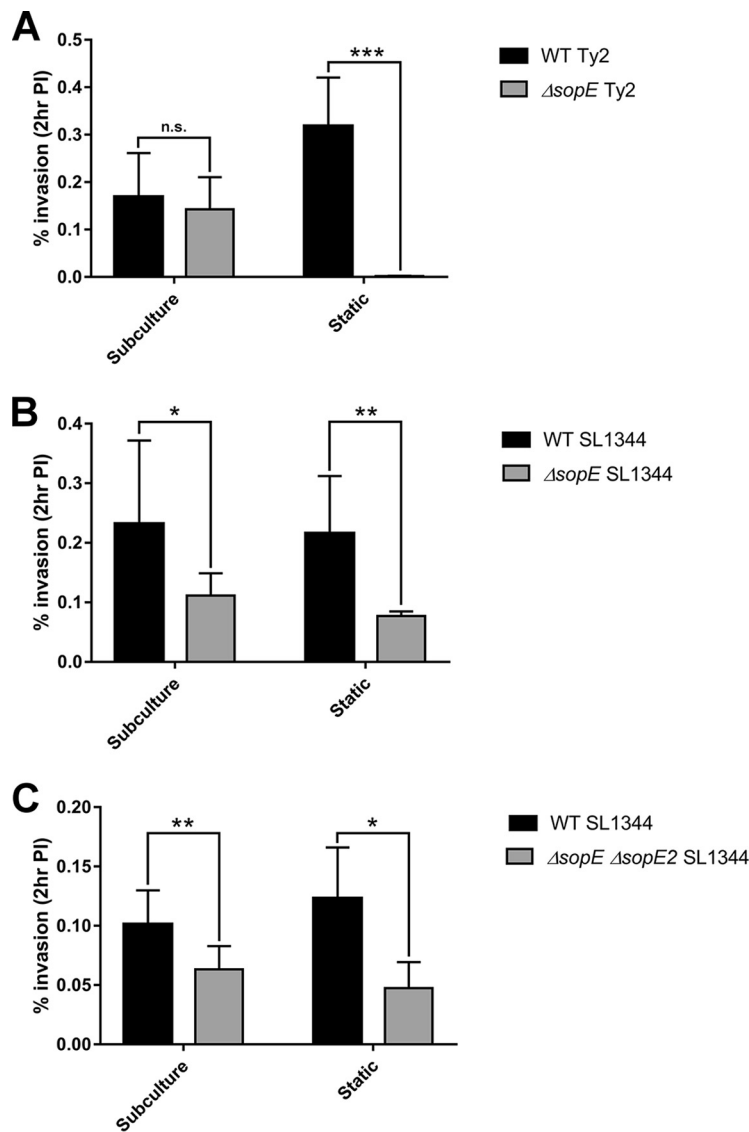
To determine if environmental conditions impact SopE activity, we assessed the invasiveness of WT or  $\Delta sopE$  *S. Typhi* and *S. Typhimurium* strains following aerobic growth to late exponential phase or microaerobic growth to stationary phase. Since *S. Typhimurium* 14028 expresses SopE2 but lacks SopE (29), while *S. Typhi* Ty2 expresses SopE but not SopE2 (6, 30), we instead used the *S. Typhimurium* strain SL1344, which has both SopE and SopE2 (31), for the comparisons in order to account for any differences in activity or regulation between SopE and SopE2 (31). Following growth to late exponential phase, deletion of *sopE* in *S. Typhi* resulted in a modest decrease in the rate of invasion into HeLa cells, with a relative invasion rate of 84.4% of that of the WT (Fig. 3A). When grown to stationary phase, however, invasion of the *S. Typhi*  $\Delta sopE$  strain was significantly attenuated (0.002% invasion efficiency, with a relative invasion rate of 0.5% of that of the WT) (Fig. 3A). A  $\Delta invA$  *S. Typhi* strain was unable to invade HeLa cells following growth to either exponential phase or stationary phase, demonstrating that invasion under both conditions is SPI-1 dependent (Fig. S1). In contrast, *S. Typhimurium* SL1344 did not demonstrate this marked phenotypic difference between growth conditions, as the  $\Delta sopE$  (Fig. 3B) and  $\Delta sopE \Delta sopE2$  (Fig. 3C) strains demonstrated similar reductions in the rates of invasion following growth to either late exponential or stationary phase.

Since these results suggest that SopE is nonessential for *S. Typhi* invasion following aerobic subculture to late exponential phase, it follows that SptP would also be dispensable, potentially explaining the lack of phenotypes previously observed for *S. Typhi*  $\Delta sptP$  compared to *S. Typhimurium*  $\Delta sptP$  (Fig. 2). The role of SptP in invasion and cytoskeletal recovery postinfection was therefore reassessed as before, but with strains instead being grown under microaerobic conditions to stationary phase. While significant differences in invasion and cytoskeletal recovery were observed between the WT and  $\Delta sptP$  *S. Typhimurium* strains, no significant differences between the WT and  $\Delta sptP$  *S. Typhi* strains were observed (Fig. 4).

**Complementation of *S. Typhimurium*  $\Delta sptP$  with *S. Typhi* *sptP*.** Since the loss of SptP does not result in phenotypes during *S. Typhi* infection the same as those observed during *S. Typhimurium* infection, it suggests that functional differences exist

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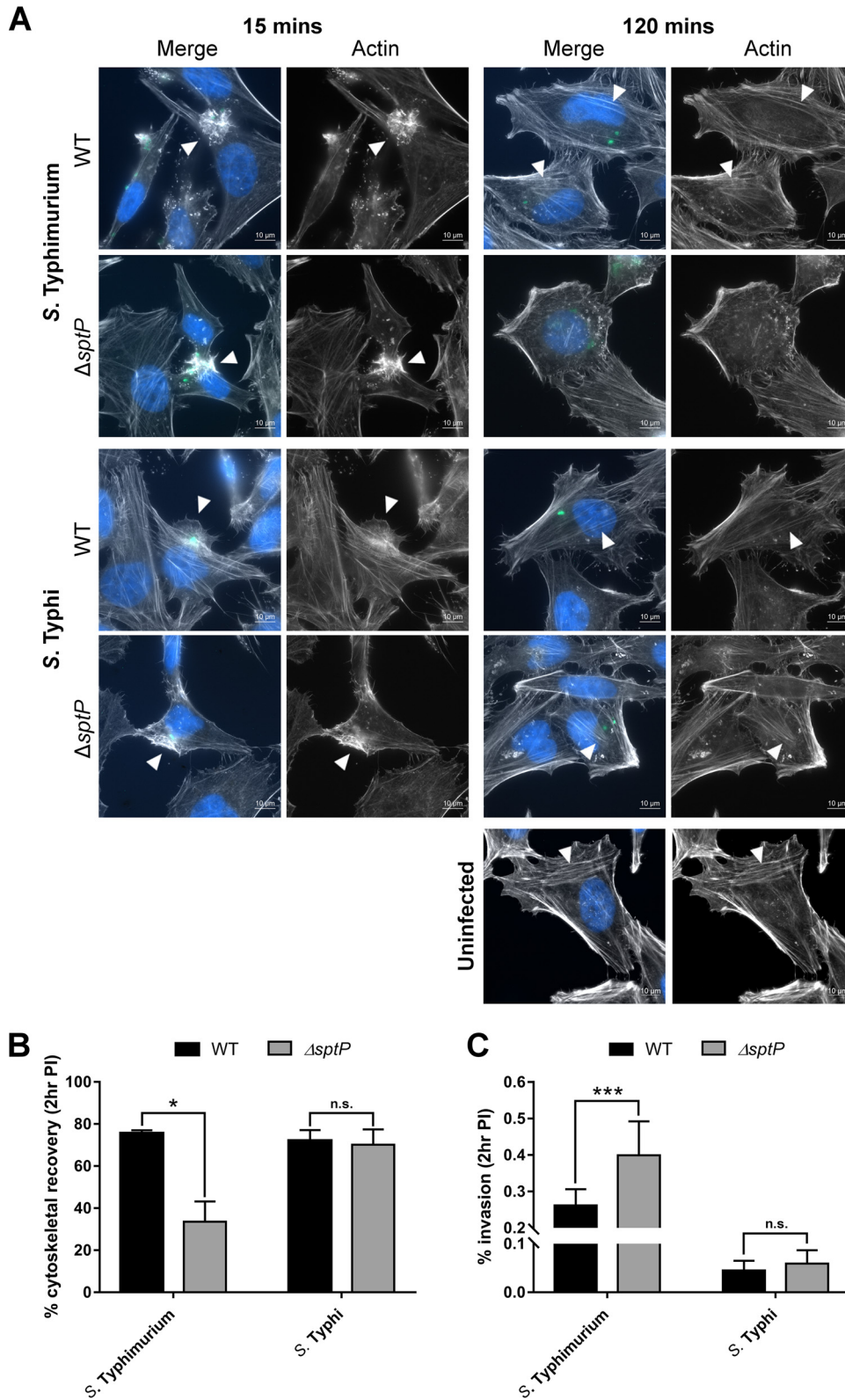
proportion of infected cells displaying a normal actin cytoskeleton. At least 100 infected cells were counted per strain and per biological repeat. Error bars show SDs ( $n = 3$ ). The levels of cytoskeletal recovery of WT and  $\Delta sptP$  strains were compared by *t* test (\*\*\*,  $P < 0.001$ ; n.s., no significant difference). (C) HeLa cells were infected with WT or  $\Delta sptP$  *S. Typhimurium* 14028 or *S. Typhi* Ty2 for 15, 30, or 60 min. The percentage of intracellular bacteria at 2 h postinfection relative to the number of bacteria added in the inoculum is shown. Error bars show SDs ( $n = 3$ ). The invasion rates of the strains were compared by *t* test (\*\*,  $P < 0.01$ ; \*\*\*,  $P < 0.001$ ).



**FIG 3** Growth conditions alter the dependency on SopE for invasion in *S. Typhi*. *Salmonella* strains were grown either aerobically to late exponential phase (subculture) or to stationary phase under microaerobic conditions (static overnight culture) and added to HeLa cells at an MOI of 100. Cells were infected for 1 h with WT and  $\Delta$ sopE *S. Typhi* (A) and 15 min for WT,  $\Delta$ sopE, and  $\Delta$ sopE  $\Delta$ sopE2 *S. Typhimurium* SL1344 (B and C). The percentage of intracellular bacteria at 2 h postinfection relative to the number of bacteria added in the inoculum is shown. Error bars show SDs ( $n = 3$ ). The invasion rates of the strains were compared by *t* test (\*,  $P < 0.05$ ; \*\*,  $P < 0.01$ ; \*\*\*,  $P < 0.001$ ; n.s., no significant difference).

between the SptP proteins from these serovars. SptP was one of several *Salmonella* T3SS proteins identified as being differentially evolved between typhoidal and nontyphoidal serovars (11). We therefore investigated if expression of *S. Typhi* SptP would be able to cross complement *S. Typhimurium*  $\Delta$ sptP. As shown above (Fig. 2 and 4), deletion of *sptP* from *S. Typhimurium* results in a significant increase in the level of invasion of HeLa cells. While complementation with *S. Typhimurium* *sptP* restored the level of invasion back to WT levels, *S. Typhimurium*  $\Delta$ sptP expressing *sptP* from *S. Typhi* was as invasive as both the  $\Delta$ sptP strain and the  $\Delta$ sptP strain carrying the empty pWSK29 vector (Fig. 5). Furthermore, a  $\Delta$ sptP *S. Typhimurium* strain expressing the truncated SptP present within H58 strains was as invasive as the strain expressing Ty2 SptP (Fig. 5).

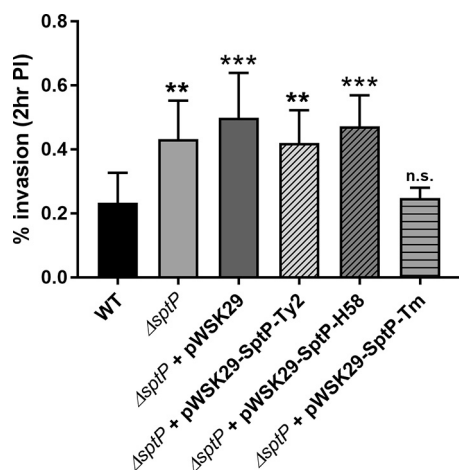
The inability of *S. Typhi* SptP to complement *S. Typhimurium*  $\Delta$ sptP could be due to either the amino acid changes within the GAP and tyrosine phosphatase domains



**FIG 4** Role of SptP in invasion and cytoskeletal recovery following growth to stationary phase. *Salmonella* strains, grown under microaerobic conditions to stationary phase, were used to infect HeLa cells at an MOI of 100. (A) Representative images of HeLa cells infected with WT or  $\Delta sptP$  *S. Typhimurium* 14028 or *S. Typhi* Ty2 at 15 min and 2 h postinfection. Actin was stained with phalloidin (white), bacteria were stained with the CSA-1 antibody (green), and nuclei were stained with Hoechst (blue). Arrowheads, invasion-associated membrane ruffles (15 min) or stress fibers (120 min). (B) HeLa cells infected with *Salmonella* strains were fixed at 2 h postinfection and stained with phalloidin to determine the proportion of infected cells displaying a normal actin cytoskeleton. At least 100

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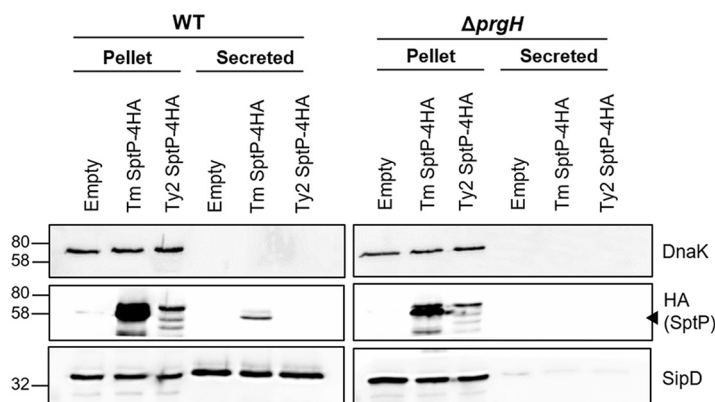
**FIG 5** Expression of *S. Typhi* SptP fails to complement the deletion in *S. Typhimurium*. WT and  $\Delta sptP$  *S. Typhimurium* strains complemented with either empty pWSK29-Spec or plasmids expressing *S. Typhimurium* (Tm) or *S. Typhi* Ty2 SptP from a constitutive promoter were grown aerobically to late exponential phase and added to HeLa cells at an MOI of 100. Strains were left to infect the cells for 15 min, and the number of intracellular bacteria was determined at 2 h postinfection. The percentage of intracellular bacteria relative to the number of bacteria added in the inoculum is shown. Error bars show SDs ( $n = 3$ ). The invasion rates of the strains were compared to the rate for the WT by one-way ANOVA, followed by Tukey's *post hoc* test (\*\*,  $P < 0.01$ ; \*\*\*,  $P < 0.001$ ; n.s., no significant difference).

rendering it functionally inactive or the numerous substitutions within the chaperone-binding region compromising the ability of *S. Typhi* SptP to bind the SicP chaperone, therefore destabilizing SptP intracellularly and preventing its secretion by the SPI-1 T3SS (13, 14) (Fig. 1). We tested these two possibilities by performing an SPI-1 secretion assay with hemagglutinin (HA)-tagged SptP expressed in *S. Typhimurium*. While SptP from both *S. Typhimurium* and *S. Typhi* was detected intracellularly in WT *S. Typhimurium*, *S. Typhi* SptP was observed at lower levels and only *S. Typhimurium* SptP was detected in culture supernatants, with secretion being dependent on the presence of a functional SPI-1 T3SS (Fig. 6). Similar levels of the SPI-1 translocon component SipD were secreted by the WT strains, suggesting that the lack of *S. Typhi* SptP secretion relates specifically to differences between the two SptP variants, likely from changes within the chaperone-binding domain.

**The chaperone-binding domain of SptP is essential for stability.** Although SicP is highly similar between *S. Typhimurium* and *S. Typhi* (97% identity), the inability of *S. Typhi* SptP to function within *S. Typhimurium* might reflect its inability to efficiently bind *S. Typhimurium* SicP, rather than indicating a general inability to bind its chaperone. As SicP binding of SptP also prevents the degradation of SptP within the bacterial cytosol prior to secretion (13), the ability of SptP to bind SicP can be assessed by detecting the total levels of HA-tagged *S. Typhi* and *S. Typhimurium* SptP in an *S. Typhi* SPI-1 mutant background ( $\Delta invA$ ), where secretion of effectors is prevented. Two constructs were used to assess SptP stability for both *S. Typhimurium* and *S. Typhi* SptP: one constitutively expressing SptP from the T3 promoter of pWSK29 and one expressing SptP from the endogenous promoter of each serovar, taken to be 700 bp upstream from the SptP start codon, on the basis of the predicted transcriptional start site from *S. Typhimurium* transcriptome sequencing data (32). While the T3 promoter accounts for any regulatory differences between *S. Typhi* and *S. Typhimurium*, expression from

#### FIG 4 Legend (Continued)

infected cells were counted per strain and per biological repeat. Error bars show SDs ( $n = 3$ ). The levels of cytoskeletal recovery of WT and  $\Delta sptP$  strains were compared by *t* test (\*,  $P < 0.05$ ; n.s., no significant difference). (C) HeLa cells were infected with WT or  $\Delta sptP$  *S. Typhimurium* 14028 or *S. Typhi* Ty2 for 30 min. The percentage of intracellular bacteria at 2 h postinfection relative to the number of bacteria added in the inoculum is shown. Error bars show SDs ( $n = 3$ ). The invasion rates of the strains were compared by *t* test (\*\*\*,  $P < 0.001$ ).

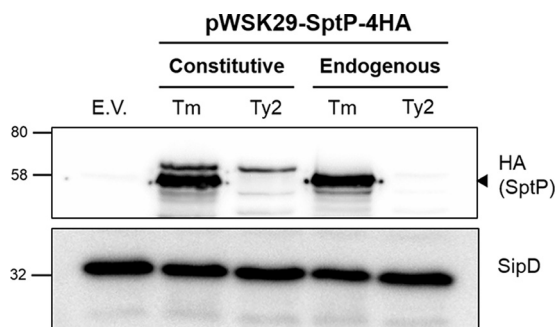


**FIG 6** *S. Typhimurium* cannot secrete *S. Typhi* SptP. WT and SPI-1 mutant *S. Typhimurium* ( $\Delta prgH$ ) strains expressing C-terminally 4HA-tagged SptP were grown to late exponential phase. The presence of *S. Typhi* Ty2 or *S. Typhimurium* (Tm) SptP in bacterial pellets and the supernatant was detected using an anti-HA antibody; the cytoplasmic protein DnaK was used as control for bacterial lysis, and the SPI-1 translocon component SipD was used as a positive control for SPI-1 secretion. A representative blot of three independent repeats is shown. The numbers on the left are molecular masses (in kilodaltons). Arrowhead, the band which corresponds to SptP.

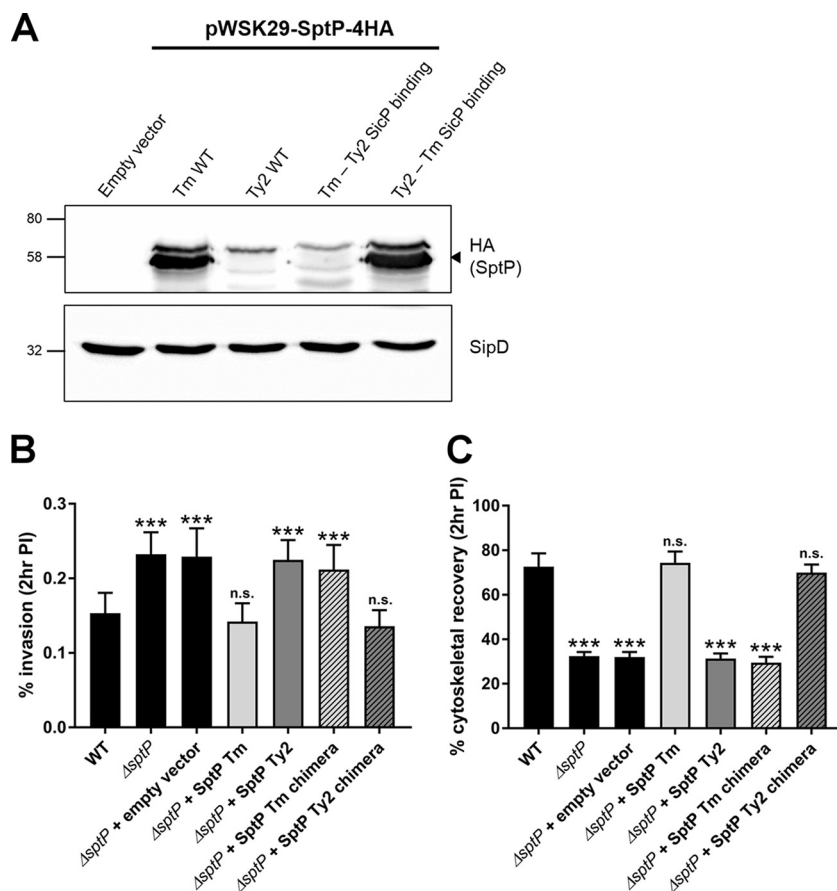
the endogenous promoter, which also encodes SicP, ensures the correct coregulation of SicP and SptP at both the transcriptional and translational levels (23).

*S. Typhi* SptP-4HA was not detectable within bacteria following expression from either the constitutive or endogenous promoter, while SptP-4HA from *S. Typhimurium* was readily detectable following expression from either construct (Fig. 7). The stability of *S. Typhimurium* SptP-4HA within *S. Typhi* indicates that SicP is still functional in this serovar, suggesting that the instability of *S. Typhi* SptP results from changes within SptP alone. It is worth highlighting that the doublet produced from the constitutive SptP-4HA construct likely arises from forced expression from the current annotated ATG start codon of SptP, as well as expression from a downstream TTG codon, reported to be the true start site for SptP (23). Given that only the lower band was secreted (Fig. 6) and the size of the *S. Typhimurium* product expressed by the endogenous promoter also corresponds to that of the lower band, it is probable that the smaller band represents the correct form of SptP.

In order to further confirm that the instability of SptP in *S. Typhi* results from changes within the chaperone-binding domain, constitutive HA-tagged SptP chimeras were constructed in which the first 139 residues of *S. Typhimurium* SptP were swapped with those of *S. Typhi* and vice versa, and the proteins were again expressed in an *S.*



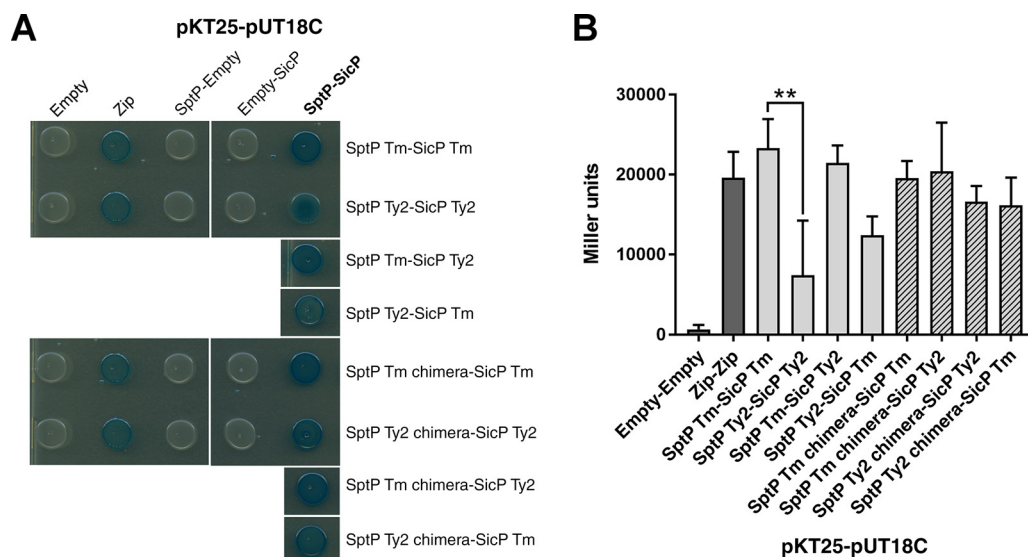
**FIG 7** SptP is unstable within *S. Typhi*. SPI-1 mutant ( $\Delta invA$ ) *S. Typhi* bacteria expressing *S. Typhimurium* (Tm) or *S. Typhi* Ty2 SptP-4HA from either a constitutive T3 promoter or the endogenous promoter of *sptP* were grown aerobically to late exponential phase. The presence of intracellular SptP was detected using an anti-HA antibody. SipD was used as both a loading control and a positive control for SPI-1 expression. A representative blot of three independent repeats is shown. The numbers on the left are molecular masses (in kilodaltons). Arrowhead, the band which corresponds to SptP; E.V., empty vector.



**FIG 8** The chaperone-binding domain of *S. Typhi* SptP is responsible for its instability. (A) SPI-1 mutant ( $\Delta invA$ ) *S. Typhi* bacteria constitutively expressing either WT SptP-4HA from *S. Typhimurium* (Tm) or *S. Typhi* Ty2 or SptP-4HA chimeras with the SicP binding domain switched with that of the other serovar were grown aerobically to late exponential phase. The presence of SptP was determined using an antibody against the HA tag. SipD was used as both a loading control and a positive control for SPI-1 expression. A representative blot of three independent repeats is shown. The numbers on the left are molecular masses (in kilodaltons). Arrowhead, the band which corresponds to SptP. (B and C) HeLa cells were infected with WT or  $\Delta sptP$  *S. Typhimurium* strains constitutively expressing either WT SptP-4HA from *S. Typhimurium* (Tm) and *S. Typhi* (Ty2) or the SptP-4HA chimeras for 15 min. The percentage of intracellular bacteria at 2 h postinfection relative to the number of bacteria added in the inoculum is shown (B), and the proportion of infected cells displaying normal actin cytoskeleton was enumerated (C). At least 100 infected cells were counted per strain and per biological repeat. Error bars show SDs ( $n = 3$ ). The invasion rates of the strains and cytoskeletal recovery were compared to those for the WT by one-way ANOVA, followed by Tukey's *post hoc* test (\*\*\*,  $P < 0.001$ ; n.s., no significant difference).

*S. Typhi* SPI-1 null background ( $\Delta invA$ ). As before, WT *S. Typhi* SptP was undetectable within bacteria; however, when the SicP binding domain of *S. Typhi* SptP was replaced with that of *S. Typhimurium*, *S. Typhi* SptP could be detected intracellularly (Fig. 8A). In contrast, *S. Typhimurium* SptP was destabilized when given the *S. Typhi* SicP binding domain, while the WT SptP remained readily detectable (Fig. 8A).

Since replacement of the chaperone-binding domain of SptP confers stability to *S. Typhi* SptP, we were interested to see if these changes would also influence the activity of *S. Typhi* SptP. Therefore,  $\Delta sptP$  *S. Typhimurium* was transformed with both WT and chimera *S. Typhimurium* and *S. Typhi* HA-tagged SptP, and invasion and cytoskeletal recovery assays were performed. While expression of *S. Typhimurium* SptP-4HA decreased the level of invasion back to WT levels, a strain expressing *S. Typhi* SptP-4HA was found to be as invasive as both the  $\Delta sptP$  strain and the  $\Delta sptP$  strain carrying the empty pWSK29 vector (Fig. 8B), in line with our previous findings (Fig. 5). The same pattern was observed with cytoskeletal recovery, with the *S. Typhimurium* SptP complemented strain achieving recovery rates akin to those achieved with WT infection,



**FIG 9** *S. Typhi* SptP interacts weakly with its chaperone, SicP. The interaction of SptP with its chaperone was assessed via a bacterial two-hybrid assay. The adenylate cyclase T25 and T18 fragments were fused to the N terminus of SptP and SicP of *S. Typhimurium* (Tm) and *S. Typhi* Ty2, respectively, and cotransformed into a *cyoA* mutant, *E. coli* DHM1. The levels of coexpression of the T25 and T18 fragments alone (Empty) or fused to leucine zipper motifs (Zip) were used as negative and positive controls, respectively. (A) Ten microliters of an overnight culture was spotted in triplicate on LB-X-Gal plates and incubated at 30°C for 24 h. Blue colonies indicate restored adenylate cyclase activity. A representative plate is shown. (B) Colonies were resuspended, and a  $\beta$ -galactosidase assay was performed in order to quantify the strength of the interaction between SptP and its chaperone, SicP (Miller units). Error bars show SDs ( $n = 3$ ). The levels of  $\beta$ -galactosidase expression induced by *S. Typhimurium* SptP and *S. Typhi* SptP were compared by *t* test (\*\*,  $P < 0.01$ ).

while the strain expressing *S. Typhi* SptP demonstrated significantly impaired cytoskeletal recovery at 2 h postinfection (Fig. 8C). However,  $\Delta$ *sptP* *S. Typhimurium* expressing the *S. Typhimurium* SptP chimera displayed significantly increased invasion and significantly reduced cytoskeletal recovery compared to WT *S. Typhimurium* (Fig. 8), demonstrating that the *S. Typhi* chaperone-binding domain abolishes the ability of *S. Typhimurium* SptP to complement *sptP* deletion. Conversely, the  $\Delta$ *sptP* *S. Typhimurium* strain was fully complemented by expression of the *S. Typhi* SptP chimera, indicating that the GAP domain of *S. Typhi* SptP is still functional.

**Binding of SptP to SicP.** Assessment of the secretion and stability of SptP within *S. Typhimurium* and *S. Typhi* represents only an indirect way to measure SptP-SicP binding. In order to directly determine the strength of the interaction between SptP and SicP of both serovars, a bacterial two-hybrid assay was performed. SptP from *S. Typhimurium* and *S. Typhi* was found to interact with SicP of both serovars, indicated by the blue coloration of the colonies on Lennox broth (LB)-5-bromo-4-chloro-3-indolyl- $\beta$ -D-galactopyranoside (X-Gal) indicator plates arising from the reconstitution of adenylate cyclase activity and subsequent induction of  $\beta$ -galactosidase expression (33). However, SptP-SicP interactions involving *S. Typhi* SptP resulted in a color change weaker than that in the equivalent cotransformations involving *S. Typhimurium* SptP (Fig. 9A). Since  $\beta$ -galactosidase expression is correlated with the strength of the interaction, a  $\beta$ -galactosidase assay was performed on the resuspended colonies in order to quantify the ability of SptP and SicP to interact between serovars. While *S. Typhimurium* SptP could interact strongly with SicP from *S. Typhimurium*, the equivalent interaction involving SptP and SicP from *S. Typhi* was significantly weaker (Fig. 9B). Interestingly, the strength of the interaction between *S. Typhimurium* SptP and *S. Typhi* SicP was equivalent to that of the interaction between *S. Typhimurium* SptP and its endogenous chaperone (Fig. 9B), further validating that SicP in *S. Typhi* is still functional.

Although replacement of the *S. Typhimurium* chaperone-binding domain with that of *S. Typhi* does not significantly decrease its interaction with SicP compared to that of

WT SptP, replacement of the chaperone-binding domain of *S. Typhi* SptP with that of *S. Typhimurium* increased the ability of *S. Typhi* SptP to bind both *S. Typhi* and *S. Typhimurium* SicP (Fig. 9B), reiterating that the reduced chaperone binding observed within *S. Typhi* SptP likely arises through changes within its SicP binding domain.

## DISCUSSION

The presence of a premature stop codon within the SPI-1 effector SptP in H58 strains (12) led us to characterize the function of this effector during *S. Typhi* infection and compare it to that of this effector during *S. Typhimurium* infection. To date, there has been very limited characterization of SptP or, indeed, any *Salmonella* T3SS effector using WT *S. Typhi*, with studies instead tending to use *S. Typhimurium* or attenuated *S. Typhi* strains.

Our data demonstrate that SptP of *S. Typhi* has reduced chaperone binding, resulting in intracellular instability and preventing secretion. Unsurprisingly, given the instability of SptP within the *Salmonella* cytosol, deletion of *sptP* from *S. Typhi* Ty2 did not affect the efficiency of invasion into HeLa cells, in line with the findings of a previous study (24), and did not affect cytoskeletal recovery postinfection. Conversely, *S. Typhimurium* SptP demonstrates strong binding to its chaperone, SicP, ensuring its stability and secretion; as such, deletion of *sptP* from *S. Typhimurium* resulted in significant phenotypic differences, including increased invasion and a failure to recover the host cytoskeleton postinfection, as previously reported (13, 16).

Although significant differences between the ability of *S. Typhimurium* and *S. Typhi* SptP to bind SicP were observed in our bacterial two-hybrid assay, this technique failed to discern any significant differences between the WT and chaperone-binding chimeras of SptP, even though marked phenotypes with the SptP chimera constructs were seen in *Salmonella* (Fig. 8). The use of *Escherichia coli* as a host, incubation at a lower temperature (30°C), and overexpression of both SptP and SicP likely promoted protein-protein interactions and reduced the overall sensitivity of the assay. However, experiments conducted in both *S. Typhimurium* and *S. Typhi* demonstrate that changes within the first 139 amino acids of *S. Typhi* SptP, corresponding to the SicP binding domain, are sufficient to compromise its activity.

Our results strongly suggest that SptP is nonfunctional in *S. Typhi*. Interestingly, *S. Paratyphi* A shares the same amino acid changes within the chaperone-binding domain, and SptP failed to be detected intracellularly by Western blotting from *S. Paratyphi* A grown to late exponential phase (28), suggesting that the instability of SptP may be a common feature of typhoidal serovars. This apparent loss of function may explain how a premature stop codon within SptP within strains belonging to the globally dominant *S. Typhi* H58 haplotype is tolerated (12), despite evidence from *in vivo* *S. Typhimurium* infections demonstrating attenuated virulence when *sptP* is deleted (20, 21). This suggests that *S. Typhimurium* is not always an appropriate model for typhoidal disease and that SptP, like the *spv* locus (8), may be required for systemic disease in mice or other hosts but not humans.

Although pseudogenization or the loss of known SPI-1 and SPI-2 effectors within *S. Typhi* has previously been reported (6, 10), the mechanism by which the function of SptP has been lost—via multiple amino acid changes, reducing its interaction with its chaperone—is unusual. A premature stop codon, such as that found in *S. Typhi* H58 strains (12), or even deletion of the gene would achieve the same end, without expending the energy invested in the transcription or translation of SptP. This could suggest that the loss of SptP function occurred gradually, maybe in conjunction with the acquisition and assimilation of a new effector with a similar function that was better suited to the induction of typhoidal disease; the presence of *S. Typhi*-specific effectors is not unlikely, given that *S. Typhi* CT18 has almost 600 genes not found in *S. Typhimurium* LT2 and effectors found only in *S. Typhimurium* and not *S. Typhi* are common (6). Alternatively, SptP is relatively unique among SPI-1 effectors, in that it is encoded within the SPI-1 pathogenicity island itself (4). Its central location within SPI-1 may not permit deletion or truncation without having deleterious effects on the

transcription or translation of neighboring genes, such as *hilA*, thereby driving the loss of function at the protein level.

If SptP is indeed not functional in *S. Typhi*, several key questions remain. Given that extended Rho GTPase activation mediated by SopE during infection would lead to proinflammatory signaling uncharacteristic of *S. Typhi* infection (17, 34), is another effector responsible for antagonizing SopE in *S. Typhi*? Perhaps most intriguingly, the question as to what function SicP, the chaperone of SptP, is performing in *S. Typhi* remains. SicP is relatively conserved between serovars (97% identity); since *S. Typhimurium* SptP is stable within *S. Typhi* and is able to strongly interact with *S. Typhi* SicP in a direct bacterial two-hybrid assay, we can be confident that SicP itself is still functional in *S. Typhi*. It is currently thought that SicP acts as a chaperone for SptP only (35); however, if it is not required to bind SptP in *S. Typhi*, it is unclear why its activity has been preserved. Although SicP is better conserved between serovars than SptP, there are amino acid changes. It is possible that, given time, these changes will accumulate and decrease the activity of SicP. An alternative explanation could be that SicP is simply acting as a chaperone for another, currently unknown effector.

Although the finding is not directly related to the function of SptP, we have also observed that different environmental conditions induced *S. Typhi*-specific regulation of the SPI-1 effector SopE. At present there are two accepted methods to induce SPI-1 expression, achieved either via aerobic growth to late exponential phase (subculture) or through growth to stationary phase under microaerobic conditions (static overnight culture) (28, 36). Our findings suggest that in comparison to *S. Typhimurium*, *S. Typhi* has additional regulatory mechanisms to control SPI-1 activity, likely in response to oxygen tension, acting at least at the level of SopE. Interestingly, a study comparing global SPI-1 expression in *S. Paratyphi* A between the different SPI-1-inducing conditions reported significant differences in SPI-1 expression between aerobic and microaerobic growth (28), while a similar study using *S. Typhimurium* SL1344 did not (36). Overall, this suggests that typhoidal serovars additionally use oxygen availability to tightly control virulence gene expression, an observation that may help explain the different diseases induced by typhoidal and nontyphoidal serovars.

Overall, this study demonstrates that the behavior of *S. Typhi* is significantly different from that of *S. Typhimurium*; the potential loss of function of SptP within *S. Typhi* exemplifies just how different these serovars are, even at the level of a single effector. This indicates that the findings of work performed in one serovar may not directly translate to the other, highlighting the need for further studies to elucidate the role of T3SS effectors in the context of *S. Typhi* infection.

## MATERIALS AND METHODS

**Bacterial strains, growth conditions, and cell culture.** The strains and plasmids used in this study are listed in Table 1. *Salmonella* strains were routinely grown in Lennox broth (LB; Sigma-Aldrich). Kanamycin (50  $\mu$ g/ml), ampicillin (100  $\mu$ g/ml), and spectinomycin (100  $\mu$ g/ml) were supplemented as required.

HeLa cells (ATCC) were maintained in Dulbecco's modified Eagle medium (DMEM) supplemented with 10% fetal bovine serum (FBS) (Sigma-Aldrich) in a 5% CO<sub>2</sub> incubator at 37°C. Cells were maintained at a maximum density of  $1 \times 10^6$  cells/ml. The HeLa cells used in this study were authenticated via short tandem repeat profiling in February 2016 (Microsynth).

**Construction of plasmids and strains.** A full list of the oligonucleotides used in this study can be found in Table S1 in the supplemental material.

To permit the use of pWSK29 in ampicillin-resistant *S. Typhi* strains, pWSK29-Spec and pWSK29-Spec-4HA were constructed by amplifying the backbone of pWSK29 (37) and pWSK29-4HA, without the ampicillin resistance cassette, with primers containing NcoI and AvrII restriction sites (primer pair 1 and 2); the *aadA* gene, which confers resistance to spectinomycin, was amplified with primers containing the same restriction sites (primer pair 3 and 4). In addition to digestion with NcoI and AvrII, both products were digested with DpnI prior to ligation.

To create SptP complementation plasmids, SptP from *S. Typhimurium* 14028 (STM14\_3477) and *S. Typhi* Ty2 (t2780) and H58 were amplified with primers containing BamHI and KpnI restriction sites (primer pair 5 and 6 and primer pair 7 and 8, respectively) and then cloned into these sites of pWSK29-Spec, generating constructs which constitutively express untagged SptP.

To create HA-tagged SptP, pWSK29-Spec-4HA was amplified with a reverse primer containing the PacI digestion site (primer pair 9 and 10). SptP was again amplified from both *S. Typhimurium* and *S. Typhi* with primers containing BamHI and PacI restriction sites (primer pair 11 and 13 and primer pair 12 and

**TABLE 1** Strains and plasmids used in this study

Strain or plasmid	Identifier	Genotype or comments	Source or reference
<b>Strains</b>			
<i>S. Typhimurium</i>			
14028	ICC797	WT	This study
14028	ICC1373	$\Delta sptP::Kan^r$	This study
14028	ICC796	$\Delta prgH::Tn5$	This study
SL1344	ICC314	WT	This study
SL1344	ICC1374	$\Delta sopE::Kan^r$	This study
SL1344	ICC1376	$\Delta sopE::FRT \Delta sopE2::Kan^r$	This study
<i>S. Typhi</i>			
Ty2	ICC1500	WT	
Ty2	ICC1522	$\Delta sptP::Kan^r$	This study
Ty2	ICC1556	$\Delta invA::Kan^r$	This study
Ty2	ICC1555	$\Delta sopE::Kan^r$	This study
<b>Plasmids</b>			
pWSK29-Amp			37
pWSK29-4HA			This study
pWSK29-Spec	pICC2489	Empty vector	This study
pWSK29-Spec	pICC2491	<i>S. Typhimurium</i> SptP, T3 promoter	This study
pWSK29-Spec	pICC2490	Ty2 SptP, T3 promoter	This study
pWSK29-Spec	pICC2535	H58 SptP, T3 promoter	This study
pWSK29-Spec-4HA	pICC2493	<i>S. Typhimurium</i> SptP-4HA, T3 promoter	This study
pWSK29-Spec-4HA	pICC2492	Ty2 SptP-4HA, T3 promoter	This study
pWSK29-Spec-4HA	pICC2494	<i>S. Typhimurium</i> SptP-4HA, endogenous promoter	This study
pWSK29-Spec-4HA	pICC2495	Ty2 SptP-4HA, endogenous promoter	This study
pWSK29-Spec-4HA	pICC2520	<i>S. Typhimurium</i> SptP-Ty2 SicP binding 4HA, T3 promoter	This study
pWSK29-Spec-4HA	pICC2521	Ty2 SptP- <i>S. Typhimurium</i> SicP binding 4HA, T3 promoter	This study
pKT25	pICC2531	Empty vector	Euromedex
pKT25	pICC2533	Zip	Euromedex
pKT25	pICC2498	<i>S. Typhimurium</i> SptP	This study
pKT25	pICC2499	Ty2 SptP	This study
pUT18C	pICC2532	Empty vector	Euromedex
pUT18C	pICC2534	Zip	Euromedex
pUT18C	pICC2504	<i>S. Typhimurium</i> SicP	This study
pUT18C	pICC2505	Ty2 SicP	This study
pKD4	pICC893	Kanamycin resistance cassette template plasmid	38
pKD46	pICC1298	Bacteriophage lambda red recombinase plasmid	38
pCP20	pICC1303	FLP recombinase plasmid	38

13, respectively). Both products were digested, and SptP was cloned into the existing BamHI site and the introduced PacI site of pWSK29-Spec-4HA, resulting in constitutively expressed C-terminally tagged SptP-4HA. SptP-4HA expressed from its endogenous promoter was constructed in the same way, with the exception that SptP was amplified from both *S. Typhimurium* and *S. Typhi* with an additional 700 bp upstream from the annotated start codon (primer pair 14 and 13) and ligated into pWSK29-Spec-4HA as described above.

To create SptP-4HA chimeras in which the chaperone-binding domain was switched with that of the other serovar, the first 417 bp of SptP, encompassing the SicP binding domain (amino acid positions 35 to 139), was amplified from the *S. Typhimurium* (primer pair 15 and 16) and *S. Typhi* (primer pair 17 and 18) chromosomes. The *S. Typhimurium* and *S. Typhi*/pWSK29-SptP-4HA plasmids were then amplified without the SicP binding domain (primer pair 19 and 21 and primer pair 20 and 21, respectively), and the two products (an *S. Typhimurium*/pWSK29-SptP-4HA backbone with *S. Typhi* SicP binding and vice versa) were assembled via Gibson assembly (50°C, 30 min) (New England BioLabs).

To determine protein interactions by a bacterial two-hybrid assay, SptP (primer pair 22 and 24 and primer pair 23 and 24 for *S. Typhimurium* and *S. Typhi* SptP, respectively) and SicP (primer pair 25 and 26 and primer pair 25 and 27 for *S. Typhimurium* and *S. Typhi* SicP, respectively) were amplified from both *S. Typhimurium* 14028 and *S. Typhi* Ty2 and cloned into the BamHI and KpnI sites of pKT25 and pUT18C, generating SptP and SicP fused to the C terminus of the T25 and T18 adenylate cyclase fragments.

Gene deletions were constructed via bacteriophage  $\lambda$  red recombination as previously described (38). Primers 28 and 30 were used to amplify the kanamycin resistance cassette of pKD4 with regions of homology to *S. Typhimurium sptP*, and primer pair 29 and 30 was used to amplify the kanamycin resistance cassette of pKD4 with regions of homology to *S. Typhi sptP*; primers 30 and 31 were used to create a kanamycin resistance cassette with *sopE* flanking regions; primer pair 32 and 33 was used to create the *sopE2* kanamycin resistance cassette product; and primers 41 and 42 were used to create the kanamycin resistance cassette with regions of *invA* homology. Correct deletion of the gene via insertion of the kanamycin resistance cassette was screened using primers 31 and 32 (*sptP*), 35 and 36 (*sopE*), 39

and 40 (*sopE2*), and 43 and 44 (*invA*). To create the SL1344  $\Delta$ *sopE*  $\Delta$ *sopE2* double mutant strain, the FLP recognition target (FRT)-flanked kanamycin resistance cassette within *sopE* was excised via FLP-mediated recombination (pCP20), generating SL1344 *sopE::FRT*, prior to transformation with the *sopE2* pKD4 PCR product.

**HeLa cell invasion assays.** The invasiveness of the  $\Delta$ *sptP* and  $\Delta$ *sopE* strains relative to that of WT *Salmonella* was determined by gentamicin protection assays. At 24 h prior to infection, HeLa cells were seeded in a 24-well plate at a density of  $7 \times 10^4$  cells/well. To induce expression of the SPI-1 T3SS, *Salmonella* strains were either cultured overnight at 37°C and 200 rpm before they were subcultured 1:33 in LB until late exponential phase (optical density at 600 nm [OD<sub>600</sub>], ~1.8) or cultured overnight at 37°C without shaking. *Salmonella* was added to the cells at a multiplicity of infection (MOI) of 100:1, and the cells and bacteria were incubated for 15 min for *S. Typhimurium* infections or for 30 min for infections involving both *S. Typhimurium* and *S. Typhi*. The cells were then washed three times with phosphate-buffered saline (PBS), before the addition of DMEM supplemented with 100  $\mu$ g/ml gentamicin. The cells were further incubated for 1 h and washed with PBS, and the gentamicin concentration was reduced to 20  $\mu$ g/ml. At the required time points, the cells were again washed three times in PBS before the addition of 0.1% Triton X-100. The cells were incubated for 5 min at room temperature (RT), before serial dilution and plating onto LB agar plates to enumerate the intracellular bacteria.

**Immunofluorescence staining and microscopy.** To compare the ability of the strains to recover the host cytoskeleton postinfection, cells were seeded onto glass coverslips and infected as outlined above. At the required time point, the cells were washed three times in PBS, before fixation with 4% paraformaldehyde for 15 min at room temperature, and were then washed a further three times in PBS. The cells were then quenched in 50 mM ammonium chloride for 10 min at RT, prior to permeabilization in 0.2% Triton X-100 for 4 min. Coverslips were blocked in 0.2% bovine serum albumin (BSA) for 5 min before addition of the primary antibody diluted in 0.2% BSA for 1 h. Intracellular *Salmonella* bacteria were detected using the CSA-1 antibody (1:200; Insight Biotechnology). The cells were again washed and blocked, prior to addition of the appropriate secondary antibody (1:200 donkey anti-goat immunoglobulin-Cy2; Jackson ImmunoResearch), Hoechst (1:1,000; Sigma), and phalloidin-tetramethyl rhodamine isocyanate (1:100; Invitrogen) for 30 min at room temperature. The coverslips were washed in PBS before they were mounted with ProLong Gold antifade reagent (Invitrogen) and visualized using a Zeiss Axio Observer Z1 microscope at  $\times 100$  magnification.

**SPI-1 secretion assays and Western blotting.** To determine the secretion of SptP, *Salmonella* strains were grown aerobically to late exponential phase in 50 ml LB. The OD<sub>600</sub> was recorded, and 1 ml bacteria were pelleted and resuspended in 2 $\times$  SDS loading buffer (1 M Tris, pH 6.8, 2% SDS, 20% glycerol, 5%  $\beta$ -mercaptoethanol, bromophenol blue) in proportion to the OD<sub>600</sub> reached to normalize bacterial numbers (10  $\mu$ l of loading buffer per 0.1 OD<sub>600</sub> unit). The remaining culture was centrifuged for 20 min at 3,300  $\times$  g before syringe filtering (pore size, 0.2  $\mu$ m). Five milliliters of the cleared supernatant was precipitated overnight at 4°C following the addition of 10% trichloroacetic acid (TCA). Precipitated proteins were recovered by centrifugation at 20,000  $\times$  g, followed by two acetone washes. The pellets were left to air dry before the addition of 2 $\times$  SDS loading buffer in the same volume used for the pellet fractions. To determine SptP stability or intracellular protein levels, the strains were grown aerobically to late exponential phase in 5 ml LB, before the OD<sub>600</sub> was recorded and 1 ml was pelleted and resuspended in 2 $\times$  SDS loading buffer. Samples were then heated at 95°C for 10 min. Proteins were separated on 12% acrylamide SDS-polyacrylamide gels, followed by semidry or wet transfer onto a polyvinylidene difluoride membrane (GE Healthcare). The membranes were blocked in 5% milk in PBS–0.05% Tween 20 and blotted with either anti-DnaK 8E2/2 (1:10,000; Enzo Life Sciences), anti-HA (1:1,000; Sigma), or anti-SipD (1:5,000) primary antibodies, followed by the addition of horseradish peroxidase-conjugated secondary antibody (1:10,000; Jackson ImmunoResearch). Following the addition of the EZ-ECL reagent (Geneflow), chemiluminescence was detected using a LAS-3000 imager (Fuji).

**Bacterial two-hybrid and  $\beta$ -galactosidase assays.** *E. coli* DHM1 was cotransformed with pKT25 and pUT18C constructs in various combinations, and transformants were selected on LB agar plates supplemented with kanamycin and ampicillin. Three colonies from each transformation were subsequently grown overnight with antibiotic selection at 37°C. Ten microliters of these cultures was spotted onto LB agar plates supplemented with kanamycin, ampicillin, 1 mM isopropyl- $\beta$ -D-thiogalactopyranoside (IPTG), and 0.1 mg/ml of 5-bromo-4-chloro-3-indolyl- $\beta$ -D-galactopyranoside (X-Gal) and incubated at 30°C for 24 h. Colonies with reconstituted adenylate cyclase activity mediated by interaction of the fused proteins were distinguished by their ability to metabolize X-Gal and subsequent blue coloration (33).

For quantitative analysis, colonies were resuspended in 1.2 ml LB. The OD<sub>600</sub> was recorded, and the bacteria were pelleted and resuspended in Z buffer (0.06 M Na<sub>2</sub>HPO<sub>4</sub>, 0.04 M NaH<sub>2</sub>PO<sub>4</sub>, 0.01 M KCl, 0.001 M MgSO<sub>4</sub>, 0.05 M  $\beta$ -mercaptoethanol, pH 7).  $\beta$ -Galactosidase assays were then performed as previously described (39).

**Statistical analysis.** Statistical tests were performed using GraphPad Prism (version 7.00) software for Windows (GraphPad Software, San Diego, CA, USA). All data are expressed as the mean  $\pm$  standard deviation (SD). Significance ( $P < 0.05$ ) was determined either by unpaired *t* test or by one-way analysis of variance (ANOVA) followed by Tukey's *post hoc* test.

## SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at <https://doi.org/10.1128/JB.00647-16>.

**TEXT S1**, PDF file, 0.2 MB.



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