

Salmonella Pathogenicity Island 2 Is Expressed Prior to Penetrating the Intestine

Nat F. Brown¹✉, Bruce A. Vallance²✉, Brian K. Coombes¹, Yanet Valdez¹, Bryan A. Coburn¹, B. Brett Finlay^{1*}

1 Michael Smith Laboratories, University of British Columbia, Vancouver, British Columbia, Canada, **2** Division of Gastroenterology, BC Children's Hospital, Vancouver, British Columbia, Canada

***Salmonella enterica* serovar Typhimurium is a facultative intracellular pathogen that causes disease in mice that resembles human typhoid. Typhoid pathogenesis consists of distinct phases in the intestine and a subsequent systemic phase in which bacteria replicate in macrophages of the liver and spleen. The type III secretion system encoded by *Salmonella* pathogenicity island 2 (SPI-2) is a major virulence factor contributing to the systemic phase of typhoid pathogenesis. Understanding how pathogens regulate virulence mechanisms in response to the environment, including different host tissues, is key to our understanding of pathogenesis. A recombinase-based in vivo expression technology system was developed to assess SPI-2 expression during murine typhoid. SPI-2 expression was detectable at very early times in bacteria that were resident in the lumen of the ileum and was independent of active bacterial invasion of the epithelium. We also provide direct evidence for the regulation of SPI-2 by the *Salmonella* transcription factors *ompR* and *ssrB* in vivo. Together these results demonstrate that SPI-2 expression precedes penetration of the intestinal epithelium. This induction of expression precedes any documented SPI-2-dependent phases of typhoid and may be involved in preparing *Salmonella* to successfully resist the antimicrobial environment encountered within macrophages.**

Citation: Brown NF, Vallance BA, Coombes BK, Valdez Y, Coburn BA, et al. (2005) *Salmonella* pathogenicity island 2 is expressed prior to penetrating the intestine. PLoS Pathog 1(3): e32.

Introduction

Salmonella is a Gram-negative bacterial pathogen that causes substantial morbidity and mortality worldwide. Human-adapted serovars cause typhoid, a systemic and life-threatening infection, while non-human-adapted serovars commonly cause enteritis. Following ingestion of contaminated food or water, the pathogenesis of both typhoid and *Salmonella* enteritis begins with an intestinal phase, while only typhoid progresses to a systemic phase. The intestinal phase of typhoid involves colonization of the intestine and penetration of the intestinal epithelium through two separate mechanisms. The first involves active bacterial invasion [1], and the second involves passive uptake of *Salmonella* during dendritic cell (DC) sampling of luminal microflora [2,3]. Once *Salmonella* has penetrated the intestinal epithelium, the systemic phase of typhoid begins by dissemination from the intestine via the lymphatics followed by colonization of macrophages of the liver and spleen [4,5]. The niche occupied by *Salmonella* within these cells is a membrane-bound compartment termed the *Salmonella*-containing vacuole. Much of our understanding of typhoid pathogenesis has come from mice infected with *S. enterica* serovar Typhimurium, which models human typhoid in several respects.

The current understanding of typhoid pathogenesis suggests that distinct virulence systems operate during the intestinal and systemic phases of infection and that these virulence systems display little overlap in their spatiotemporal activation. These virulence factors are type III secretion systems (T3SSs) that translocate numerous *Salmonella* virulence proteins (termed effectors) directly into the host cell,

where they alter various host cell processes [6]. The T3SS encoded by *Salmonella* pathogenicity island 1 (SPI-1) allows *Salmonella* to invade non-phagocytic cells and penetrate the intestinal epithelium, and is the major factor involved during the intestinal phase of typhoid [7–9]. SPI-1 mutant *Salmonella* Typhimurium is fully virulent when inoculated intraperitoneally into mice, indicating that the role played by SPI-1 is limited to the intestinal phase of *Salmonella* infection [9]. In contrast, *Salmonella* pathogenicity island 2 (SPI-2) mediates *Salmonella* replication within macrophages at systemic sites, and SPI-2 mutant *Salmonella* is avirulent when inoculated intraperitoneally [10–12]. Studies of *Salmonella*-induced enteritis have found that the role of SPI-2 in the intestine is subtle when compared to the role of SPI-1 [13–16]. However, it should be noted that *Salmonella*-induced enteritis and typhoid are distinct diseases involving different intestinal pathologies, and as such the pathogenesis of the diseases is not directly comparable. In particular, typhoid does not typically involve

Received June 15, 2005; Accepted October 17, 2005; Published November 18, 2005
DOI: 10.1371/journal.ppat.0010032

Copyright: © 2005 Brown et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Abbreviations: DC, dendritic cell; LB, Luria-Bertani; RIVET, recombinase-based in vivo expression technology; SPI-1, *Salmonella* pathogenicity island 1; SPI-2, *Salmonella* pathogenicity island 2; T3SS, type III secretion system

Editor: Ralph Isberg, Tufts University School of Medicine, United States of America

* To whom correspondence should be addressed. E-mail: bfinlay@unixg.ubc.ca

✉ These authors contributed equally to this work.

Synopsis

Typhoid fever is a disease caused by specific *Salmonella* strains and is a significant cause of mortality in many regions of the developing world. Following a person's ingestion of *Salmonella*, the bacteria initially colonize the intestine, which they subsequently breach to reside in immune cells of the liver and spleen. The ability to survive inside immune cells directly contributes to the ability of *Salmonella* to cause typhoid, and is conferred upon *Salmonella* by the so-called *Salmonella* pathogenicity island 2 (SPI-2) type III secretion system. Previous work has shown that while SPI-2 is specifically turned on inside host cells, it is not active when grown in typical laboratory medium. Owing to these facts, it has been hypothesized that *Salmonella* specifically turn on SPI-2 inside host cells after breaching the host intestine. The researchers developed a sensitive system in *Salmonella* to test this hypothesis using a mouse model of typhoid. Interestingly, SPI-2 was specifically turned on before *Salmonella* breached the intestine, suggesting that SPI-2, which is integral to virulence, is active in a preemptive fashion to allow *Salmonella* to survive within the immune system.

significant intestinal inflammation [16]. This has led to models in which *Salmonella* employs a two-tiered expression of virulence systems that correspond to the biphasic pathogenesis of typhoid, with SPI-1 mediating intestinal pathogenesis and SPI-2 mediating systemic pathogenesis.

While it is clear that SPI-2 is a major virulence factor leading to mortality in murine typhoid, our current understanding of its molecular function is limited. Its primary role appears to be subversion of host cell membrane traffic, allowing replication of intracellular *Salmonella* [17]. Consistent with its role during intracellular growth, in vitro studies have shown that expression of SPI-2 is induced inside host cells and in culture medium that mimics the environment of the *Salmonella*-containing vacuole [18–20]. In vitro, SPI-2 expression is regulated by the SPI-2-encoded regulatory system SsrA/B [18,19], and efficient transcription of *ssrAB* requires the regulator OmpR [21]. Recent work by Hensel and colleagues has shown that the SPI-2 T3SS is typically expressed as one apparatus per bacterium, during growth in host cells or during inducing in vitro culture [22].

Countless studies have shown that bacteria regulate their expression profile by sensing the extracellular environment and responding accordingly. As an environment in which to study bacterial gene expression, the mammalian host poses unique challenges because of its complexity and dynamic nature. Consequently, analysis of the spatiotemporal expression pattern of virulence genes in vivo has rarely been undertaken. Of particular interest to the field of *Salmonella* research is the expression pattern of SPI-2 during typhoid because it is the central virulence mechanism and is considered to play a role specific to a particular phase of pathogenesis. Based on this, we hypothesized that SPI-2 expression would be confined to the peripheral lymphoid tissues, spleen, and liver. To address this, we examined the spatiotemporal expression pattern of three SPI-2 promoters during experimental murine typhoid. In contrast to the current model of SPI-2-mediated pathogenesis that argues for exclusive intracellular expression, we found that expression of SPI-2 occurs during initial stages of pathogenesis in the lumen of the intestine.

Results

SPI-2 Gene Expression In Vitro Assessed Using RIVET

Recombinase-based in vivo expression technology (RIVET) is an exquisitely sensitive reporter of gene expression. This system involves the construction of a transcriptional fusion to a site-specific recombinase, which mediates the loss of a selectable genetic marker (a process called resolution) [23]. This approach has been applied to elucidate the spatiotemporal expression patterns of the toxin-coregulated pilus and cholera toxin during *Vibrio cholerae* infection of mice [24]. We designed and constructed a *Salmonella* Typhimurium strain possessing all the genetic requirements for RIVET and *lacZ* fusion analysis of P_{sseA} (see Materials and Methods). The ability of this strain to function as a RIVET reporter of P_{sseA} activity was initially assessed during in vitro growth in media that have previously been shown to induce (LPM medium) or not induce (Luria-Bertani [LB] medium) expression from SPI-2 promoters (Figure 1). P_{sseA} activity was assessed using RIVET and β -galactosidase activity as well as by monitoring cytoplasmic levels of SseB, a protein expressed from P_{sseA} . A low level of P_{sseA} activity was detected for each output in the non-inducing LB medium whereas high levels were detected from bacteria cultured in the inducing LPM medium, indicating that RIVET correlates well with standard methods. Additionally, RIVET was considerably more sensitive at detecting P_{sseA} activity than *lacZ* transcriptional fusion and immunoblotting for native SseB levels. This was most obvious when the number of bacteria in the culture was below the sensitivity of conventional reporter systems. Furthermore, the high degree of sensitivity of RIVET was deemed important for the study of the monocopy expression level of the SPI-2 T3SS. Using RIVET, we also confirmed the role of SsrB and OmpR in regulating SPI-2 gene expression (Figure 1D).

Salmonella infection of cultured mammalian cells is a model that is frequently used to approximate conditions encountered during infection in vivo. We determined the expression from P_{sseA} at 1 h following uptake into various mammalian cells using the RIVET system (Figure 2). In HeLa cells (epithelial), where bacteria actively induce their own uptake using the SPI-1 T3SS, substantial induction of expression from P_{sseA} occurred after 1 h in an SsrB- and an OmpR-dependent fashion. This demonstrates that SPI-2 is activated rapidly following SPI-1-mediated *Salmonella* invasion of an epithelial cell. The role of SPI-2 in mediating replication within macrophages has been thoroughly documented [17], and DCs are also a potentially important cell type encountered by *Salmonella* in vivo [2,3]. We therefore determined the kinetics of P_{sseA} induction in RAW264.7 murine macrophages and murine DCs. A significant induction of SsrB- and OmpR-dependent P_{sseA} expression could be detected within 1 h of *Salmonella* being added to either cell type (Figure 2).

SPI-2 Gene Expression In Vivo Assessed Using RIVET

Once it was established that the RIVET system was sensitive and specific, the RIVET system was used to determine the spatiotemporal expression pattern of P_{sseA} during infection of mice. To facilitate the synchronized arrival of a bacterial load in the distal ileum, the inoculum was delivered into a single loop constructed in the ileum of each mouse tested (see Materials and Methods). At times ranging from 15 min to 4 h

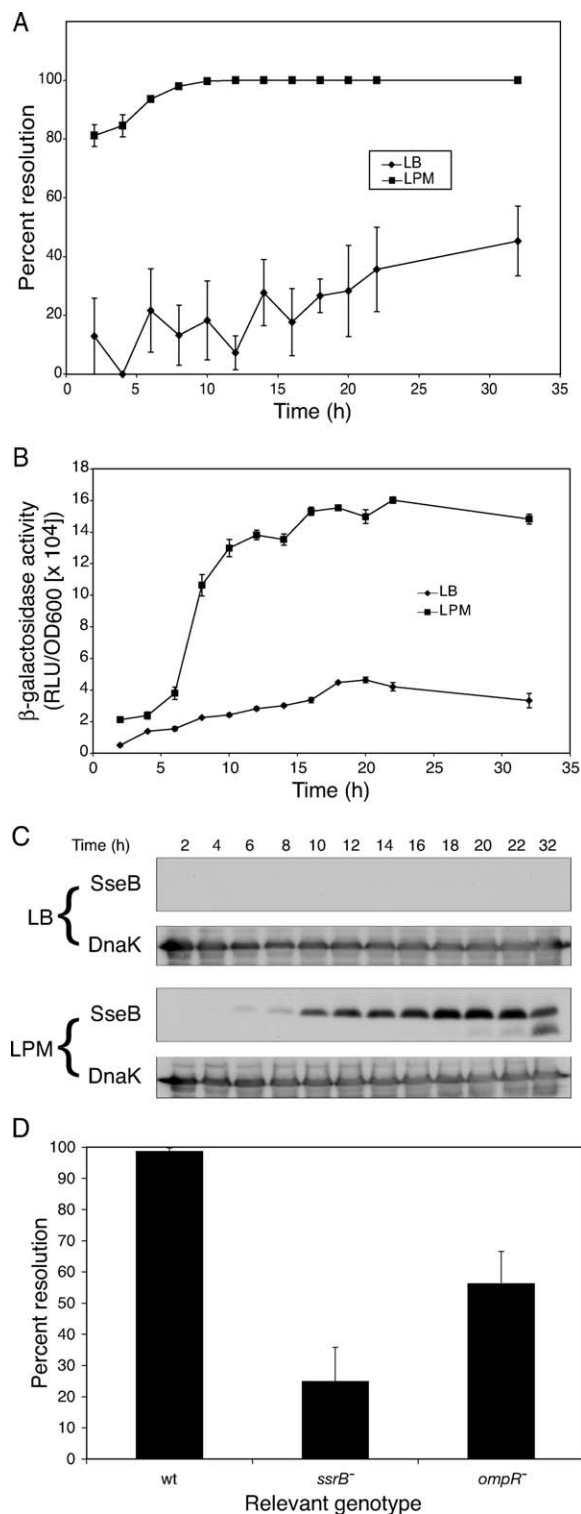


Figure 1. Expression from P_{sseA} during Growth of *Salmonella* in Inducing and Non-Inducing Culture Media

(A–C) The wild-type *Salmonella* RIVET strain (NB25) was inoculated into LPM (inducing) and LB (non-inducing) media at 1/100 of the culture volume, and cultures were incubated with shaking at 37 °C. At the indicated time points, samples were taken for measurement of culture OD₆₀₀ (unpublished data), colony-forming units per millilitre (unpublished data), β -galactosidase activity, percent resolution, and native SseB levels. The results shown for percent resolution (A) and β -galactosidase activity (B) are the mean \pm standard error of the mean from three independent experiments. The levels of SseB associated with bacteria were detected by immunoblotting (C). Protein loading was normalized

to culture OD₆₀₀ and was confirmed by immunoblotting for the abundantly expressed cytoplasmic protein DnaK.

(D) Wild-type (NB25), *ssrB*⁻ (NB7), and *ompR*⁻ (NB15) were grown in LPM medium and percent resolution was determined from the cultures at 10 h. DOI: 10.1371/journal.ppat.0010032.g001

following inoculation, the liver, spleen, mesenteric lymph nodes, and ileal loop were dissected, homogenized, and plated to determine the degree of P_{sseA} induction using RIVET (Figure 3A). These results clearly demonstrated that expression from P_{sseA} was induced within 15 min following inoculation and that this induction occurred within the ileum. RIVET reports gene expression in an irreversible fashion, and data on expression at systemic sites and later time points was not considered reliable as the majority of bacteria had undergone resolution within 15 min. We also assessed *ssrB*⁻ and *ompR*⁻ strains for their ability to induce expression from P_{sseA} within the ileum 15 min following inoculation (Figure 3B). These data showed that both regulators play a significant role in the observed induction in the ileum and directly showed the involvement of these regulators in the expression of SPI-2 in vivo.

The experiments described above showed that P_{sseA} , the promoter driving expression of the SPI-2 T3SS translocon and effectors, is active within 15 min of inoculation into ileal loops. To see if the promoters driving expression of the remaining components of the SPI-2 T3SS were active on a similar timescale, we constructed strains for RIVET analysis of transcription from P_{spiC} and P_{ssaG} . These strains were confirmed to have the typical SPI-2 expression characteristics during in vitro culture, including being dependent on the SPI-2-encoded transcription factor SsrB (data not shown). When these strains were inoculated into mouse ileal loops, expression after 15 min from both P_{spiC} and P_{ssaG} could be observed (Figure 3C), similar to that of previous experiments with P_{sseA} . These results showed that all SPI-2 T3SS components are expressed in a large proportion of *Salmonella* within 15 min of arriving in the ileum.

The short time frame in which SPI-2 expression was induced in vivo prompted us to investigate the location of

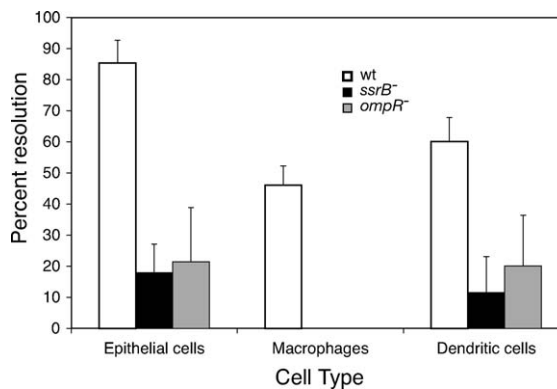


Figure 2. Expression from P_{sseA} during Infection of Mammalian Cells In Vitro

The human epithelial cell line HeLa, murine macrophage cell line RAW264.7, and bone-marrow-derived DCs were infected with wild-type (NB25), *ssrB*⁻ (NB7), and *ompR*⁻ (NB15) RIVET strains as described in Materials and Methods. At 1 h post-infection intracellular bacteria were recovered and the percent resolution was determined. DOI: 10.1371/journal.ppat.0010032.g002

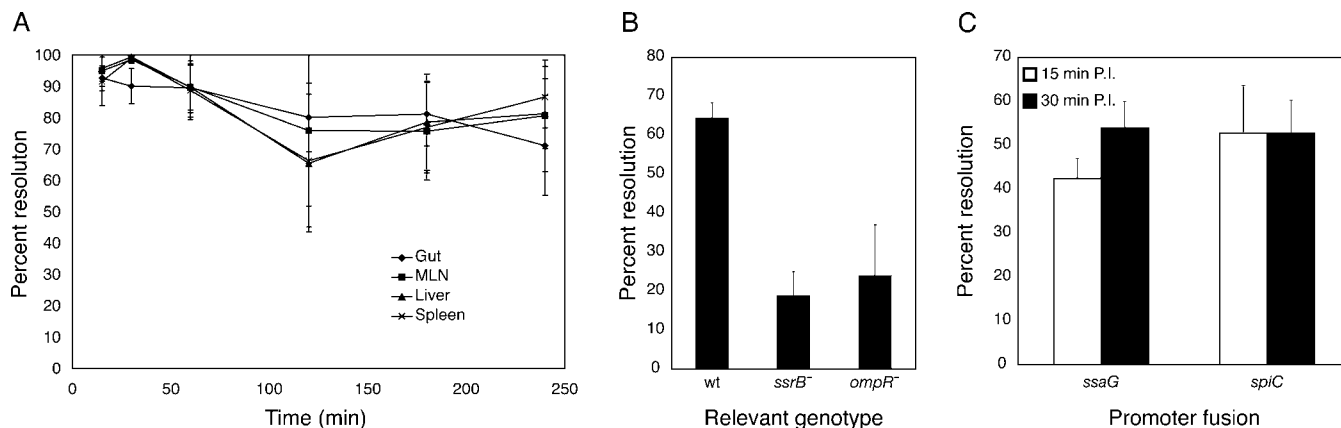


Figure 3. Kinetics of SPI-2 Expression during Murine Typhoid

(A) Mice were infected with the wild-type P_{sseA} RIVET strain (NB25) as described in Materials and Methods. At the indicated times following inoculation, the ileal loops, mesenteric lymph nodes, spleen, and liver were taken and assessed for expression from P_{sseA} by determining the percent resolution. The results shown are the mean \pm standard error of the mean from three independent experiments.

(B) Mice were infected with the wild-type (NB25), $ssrB^-$ (NB7), and $ompR^-$ (NB15) P_{sseA} RIVET strains, and at 15 min following inoculation, the ileal loop was removed and homogenized, and the percent resolution of the infecting *Salmonella* was determined.

(C) Mice were infected with the wild-type P_{spiC} (NB33) and P_{ssaG} (NB31) RIVET strains for 15 and 30 min, the ileum was removed and homogenized, and the percentage resolution determined. The results shown are the mean \pm standard error of the mean for three independent experiments.

DOI: 10.1371/journal.ppat.0010032.g003

the bacteria that had induced SPI-2 expression. Specifically, we wanted to know whether *Salmonella* was located within host cells when SPI-2 expression was induced. As SPI-1 mediates the major route of epithelial penetration [2,8], we tested expression from P_{sseA} in a SPI-1 mutant ($invA^-$) in vivo. At 15 min following inoculation, the induction of expression from P_{sseA} was independent of SPI-1 (Figure 4A), strongly suggesting that the SPI-2 expression we were observing was occurring in the lumen of the intestine. To further test this, we directly determined the localization of wild-type *Salmonella* 15 min following inoculation into ileal loops using confocal microscopy. As expected, the vast majority of bacteria were not located within host cells but rather were associated with the apical surface of the epithelium (Figure 4B). These data

are inconsistent with the observed induction of SPI-2 occurring exclusively in an intracellular compartment in vivo.

To test whether *Salmonella* association with the cytoplasmic membrane of a host cell initiates expression from P_{sseA} , we infected HeLa cells in vitro with noninvasive $invA^-$ *Salmonella*. Extracellular, cell-associated $invA^-$ *Salmonella* at 1 h post-infection had not induced significant expression from P_{sseA} ($3.54\% \pm 3.01\%$ resolution). In contrast, internalized wild-type *Salmonella* controls showed a high level of expression ($75.10\% \pm 5.75\%$ resolution). This is consistent with cell culture medium providing a non-inducing environment for SPI-2 expression (data not shown) and shows that the host cell plasmalemma does not induce SPI-2 expression. This suggested that a stimulus for expression from P_{sseA} exists in the luminal contents of the small intestine. We therefore conducted experiments to address this hypothesis. Wild-type, $ssrB^-$, and $ompR^-$ RIVET strains were incubated for up to 30 min at 37 °C with contents collected from mouse small intestine and then resolution was measured. No resolution was observed for any strain, suggesting that the stimulus inducing expression of SPI-2 in the intestine is not capable of being extracted with naive luminal contents of the small intestine.

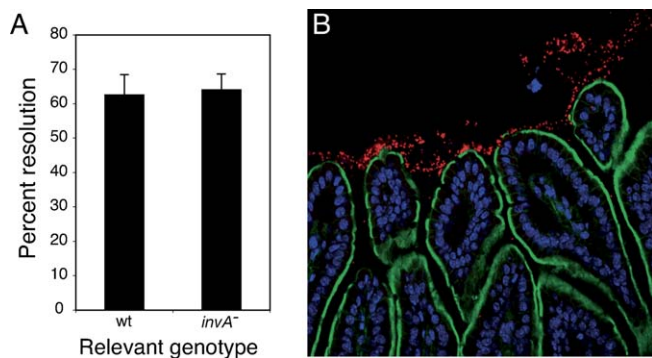


Figure 4. *Salmonella* Induces Expression of SPI-2 in the Lumen of the Ileum

(A) Mouse ileal loops were infected with either wild-type or $invA^-$ P_{sseA} RIVET strains for 15 min before the loops were removed and homogenized, and the percent resolution was determined.

(B) Mouse ileal loops were infected with wild-type *Salmonella* for 15 min before being removed, fixed, stained, and analyzed by confocal microscopy with an oil immersion 40 \times 1.3 N.A. objective. A representative image is displayed showing host cell nuclei in blue, actin in green, and *Salmonella* in red.

DOI: 10.1371/journal.ppat.0010032.g004

Discussion

By using a sensitive methodology and testing earlier time points than have been presented in previous studies of SPI-2 function, we have established that SPI-2 genes are expressed very early in the intestine. The data presented above are inconsistent with the currently held view that the initial induction of SPI-2 expression occurs in response to an intracellular environment. Our attempts to confirm the RIVET expression data using β -galactosidase reporters to measure SPI-2 expression in vivo failed to detect sufficient signal to reliably report SPI-2 expression (data not shown). However, extensive controls were performed in vitro, where sufficient numbers of bacteria can be analyzed to detect β -

galactosidase reporters as well as SPI-2 proteins by western blotting. These control experiments confirmed that RIVET was a sensitive and specific reporter of SPI-2 expression. The failure of β -galactosidase to accurately report on SPI-2 expression in vivo is most likely a reflection of low bacterial numbers in the examined tissues. This highlights a major advantage of RIVET in its capacity to report on gene expression from small populations of bacteria.

Previous work has focused on mimicking the intracellular environment encountered by *Salmonella* to define the cues for inducing expression of SPI-2. Our results show that SPI-2 is induced in the gut lumen, prior to encountering an intracellular environment. We have investigated signals present in the small intestine as potential cues for SPI-2 expression and have determined that intestinal SPI-2 induction is not a result of association with host cell plasmalemma or luminal contents from the small intestine of uninfected mice. Additionally, we have been able to rule out low oxygen tension as a cue for SPI-2 induction in the intestine (data not shown). Our data are compatible with a rapid host response to *Salmonella* acting as a cue for the induction of SPI-2 expression. One such response would be the secretion of antimicrobial products by paneth cells, specialized epithelial cells that reside at the base of the crypts and respond to antigenic stimuli including *Salmonella* [25]. However, experiments with *mmp7*^{-/-} mice, which are deficient in paneth cell products, have indicated that these products are not a stimulus for SPI-2 expression in vivo (data not shown). Other potential host responses acting as a stimulus for SPI-2 expression will be a topic of future investigation.

Previous studies on SPI-2 and its translocated effectors have almost exclusively focused on the role played during the systemic phase of typhoid, and consequently little is known about the potential actions of SPI-2 during the intestinal phase of typhoid. Others have shown that SPI-2 mutant *Salmonella* colonizes the caecum and Peyer's patches to a lesser extent than wild-type *Salmonella* during typhoid [19]. This, together with our data on expression, firmly establishes a role for SPI-2 prior to colonization of systemic sites during typhoid. It is important to consider the localization of intestinal *Salmonella* in order to speculate on the role SPI-2 could be playing at this location. Although the vast majority of bacteria are present in the luminal space of the intestine, it is well established that *Salmonella* transits through epithelial cells and can reside within sub-epithelial phagocytic cells. Given that type III secretion mediates the direct delivery of effector proteins into the host cell cytosol [6] and that the system encoded by SPI-2 allows *Salmonella* to replicate within macrophages [17], this suggests that the location where SPI-2 is actually functioning is within phagocytic cells of the intestine. It is likely that SPI-2 expression initiated in the intestinal lumen primes *Salmonella* for residency in intestinal phagocytic cells prior to the bacteria reaching this niche. We do not conclude that SPI-2 enables replication of luminal *Salmonella*.

The rapid induction of SPI-2 in the lumen of the ileum suggests that for *Salmonella* to initiate a successful infection of sub-epithelial and systemic sites, it must be prepared for the harsh environment of the macrophage endosomal system prior to the encounter. This finding may establish a paradigm for all pathogens where preemptive synthesis of important virulence factors occurs prior to the transition from

colonization of a mucosal surface to colonization of systemic sites. We also speculate that functions mediated by SPI-2 during early stages of infection may not be limited to the establishment of a *Salmonella*-containing vacuole that supports bacterial replication. Such possibilities include enhancing dissemination to systemic sites. By identifying additional SPI-2-dependent functions at these early intestinal stages of typhoid pathogenesis, we hope to elucidate key mechanisms that facilitate the successful parasitic lifestyle of *Salmonella*.

Materials and Methods

Bacterial strains and plasmids. Bacterial strains and plasmids are shown in Table 1. All *Salmonella* strains used in this study were derived from the wild-type strain SL1344. All bacteria were routinely cultured in LB medium. LPM medium (pH 5.8) has been described previously [26]. Kanamycin was used at 50 $\mu\text{g ml}^{-1}$, chloramphenicol at 10 $\mu\text{g ml}^{-1}$, and ampicillin at 100 $\mu\text{g ml}^{-1}$. *Salmonella* RIVET strains were always grown in LB medium containing ampicillin and chloramphenicol prior to conducting experiments.

For RIVET studies, it is desirable to insert the resolvable cassette (*res*-marker-*res*) into a neutral site within the genome. *ushA* is a silent gene in *Salmonella* Typhimurium because of an inactivating S139Y substitution in the expressed protein product [27], and as such this was considered to be a good site to insert the resolvable cassette for RIVET. A region of approximately 1.8 kb spanning *ushA* was amplified by PCR using the oligonucleotides *ushako-f* (5' GCAAC TAGTGGATGTTGGAGATAGTAGGATGTG 3') and *ushako-r* (5' GATGTCCACCCTACCATTGGCGTAAACG 3') and was cloned into pBLUESCRIPT KS+ using SpeI and SalI to create pUshA. A resolvable cassette mediating resistance to chloramphenicol was constructed by subcloning the *res-npt-sacB-res*-containing SphI fragment into pUC19 to create pUC-RES. The *npt-sacB* portion was replaced with *cat* to give pUC-RESCm. The *res-cat-res* cassette was then inserted into the middle of *ushA* in pUshA, and then the *ushA::res-cat-res* allele was cloned into pCVD442 using SacI and KpnI to give pUshA::RES KO. Using this plasmid, the *ushA::res-cat-res* allele was introduced into SL1344 and SL1344 Δ *ompR* using standard allelic exchange methodologies [28] to give strains NB24 and NB13, respectively. NB24 was shown to have no detectable defect in virulence when inoculated intraperitoneally into mice by either a time-to-death assay (N. F. B. and B. A. V., unpublished data) or competitive index (B. K. C. and M. Wickham, unpublished data). *Salmonella* strains NB6 and NB16 were created by P22 generalized transduction of *ssrB::Km* and *invA::Km* alleles into NB24, respectively.

Transcriptional fusions were generated as follows. An approximately 1.5-kb region upstream of *sseA* was amplified by PCR using oligonucleotides *pssea-f* (5' ATACTCGAGCGTATTCTTCATTTT CATCGGTG 3') and *pssea-r* (5' ATACAATTGCCCTTTCAG CAAGCTGTTGAC 3') and cloned into pIVET5nMut135 using XhoI and MfeI. Other reporter strains were generated using the same strategy using oligonucleotides *psagfu-f* (5' ATGCTCGAGAAT TACCTCCGTTAGCCTGAC 3') and *psagfu-r* (5' ATG CAATTGCTGCCTGGTGCGCCATGTG 3') for *P*_{*ssaG*} and oligonucleotides *psab-f* (5' ATCTCGAGTCGGCTGCAATT GAAGG 3') and *psab-r* (5' ATACAATTGCCAGCAT GAATCCCTCCTCAGAC 3') for *P*_{*spiC*}. The resulting reporter plasmids were conjugated into the desired *Salmonella* strains from *Escherichia coli* SM10 λ *pir*. The resulting strains had the reporter plasmid integrated into the chromosome by homologous recombination, resulting in a merodiploid genotype for the promoter region being studied.

Gene expression reporter assays. Resolution assays were performed by plating serial dilutions of sample material onto LB ampicillin plates and incubating these overnight at 37°C. The following day, plates with between 50 and 200 colonies were replica-plated onto LB ampicillin plates and LB ampicillin chloramphenicol plates, which were incubated overnight at 37°C. Colonies that grew on the ampicillin only plates, but not on the ampicillin chloramphenicol plates were considered to have undergone the resolution event. β -galactosidase assays were performed as previously described [29] using Galacto-Star chemiluminescent substrate (Applied Biosystems, Foster City, California, United States) and were read using a Spectrafluor Plus (Tecan, Mannedorf, Switzerland) in luminescence mode.

Antibodies and reagents. Antibodies to SseB have been described previously [30] and were used at a concentration of 1:1,000, and a

Table 1. Bacterial Strains and Plasmids Used in This Study

Category	Strain/Plasmid	Genotype/Description	Reference/Source
<i>E. coli</i>	DH5 α	<i>supE44 Δlacu169 (ϕ80 <i>lacZ</i>ΔM15) <i>hsdR17 recA1 endA1 gyrA96 thi-1 relA1</i></i>	[31]
	SM10 λ pir	<i>thi recA thr leu tonA lacY supE RP4-2-Tc::Mu λpir</i>	[32]
<i>Salmonella</i>	SL1344	Wild-type strain	[33]
	SB103	<i>invA::Km</i>	[9]
	SL1344 Δ ompR	Δ ompR	B. B. F. (unpublished data)
	SL1344 <i>ssrB::Kan</i>	<i>ssrB::Km</i>	[28]
	NB6	<i>ssrB::Km ushA::res-cat-res</i>	This study
	NB7	<i>ssrB::Km ushA::res-cat-res sseA::pPseAMut135</i>	This study
	NB13	Δ ompR <i>ushA::res-cat-res</i>	This study
	NB15	Δ ompR <i>ushA::res-cat-res sseA::pPseAMut135</i>	This study
	NB16	<i>invA::Km ushA::res-cat-res</i>	This study
	NB23	<i>invA::Km ushA::res-cat-res sseA::pPseAMut135</i>	This study
	NB24	<i>ushA::res-cat-res</i>	This study
	NB25	<i>ushA::res-cat-res sseA::pPseAMut135</i>	This study
	NB31	<i>ushA::res-cat-res ssaG::pPssaGMut135</i>	This study
	NB32	<i>ssrB::Km ushA::res-cat-res ssaG::pPssaGMut135</i>	This study
	NB33	<i>ushA::res-cat-res spiC::pPsiCMut135</i>	This study
NB34	<i>ssrB::Km ushA::res-cat-res spiC::pPsiCMut135</i>	This study	
Plasmid	pRES	<i>lacZ::res-npt-sacB-res bla oriT oriR6K</i>	[34]
	pIVET5nMut135	<i>tnpR::lacZ bla oriT oriR6K</i>	[23]
	p345-Cm2	<i>cat</i>	[35]
	pCVD442	<i>bla oriT oriR6K sacB</i>	[26]
	pBLUESCRIPT KS+	<i>oriColE1 oriF1(+)</i> <i>bla lacZα</i>	Stratagene (La Jolla, California, United States)
	pUC19	<i>oriColE1 bla lacZα</i>	[36]
	pUshA	<i>ushA</i> . pBLUESCRIPT KS+ derivative	This study
	pUC-RES	<i>res-npt-sacB-res</i> . pUC19 derivative	This study
	pUC-RESCm	<i>res-cat-res</i> . pUC-RES derivative	This study
	pUshA::RES KO	<i>ushA::res-cat-res</i> . pCVD442 derivative	This study
	pPseAMut135	<i>P_{sseA}::tnpR::lacZ</i> . pIVET5nMut135 derivative	This study
	pPssaGMut135	<i>P_{ssaG}::tnpR::lacZ</i> . pIVET5nMut135 derivative	This study
	pPsiCMut135	<i>P_{spiC}::tnpR::lacZ</i> . pIVET5nMut135 derivative	This study

DOI: 10.1371/journal.ppat.0010032.t001

monoclonal antibody to DnaK (Stressgen Biotechnologies, Victoria, British Columbia, Canada) was used at a concentration of 1:2,000. Anti-*Salmonella* Typhimurium LPS antiserum (Difco, Becton-Dickinson, Franklin Lakes, New Jersey, United States) was used at a concentration of 1:200. HRP-labeled anti-mouse and anti-rabbit antibodies were used at a concentration of 1:5,000 and were purchased from Jackson Immunoresearch Laboratories (West Grove, Pennsylvania, United States). Alexafluor 568-labeled anti-rabbit antibodies (Molecular Probes, Eugene, Oregon, United States) were used at a concentration of 1:200. Alexafluor 488-labeled phalloidin and DAPI were purchased from Molecular Probes.

Cell culture and infection of cultured cells. HeLa and RAW264.7 cells were cultured using DMEM containing 10% FCS. Bone-marrow-derived DCs were derived as previously described [31]. Briefly, 2.5×10^6 bone marrow cells were cultured in 10 ml of Iscove's Modified Dulbecco's Medium supplemented with 10% FCS, GM-CSF (20 ng/ml), and IL-4 (10 ng/ml). The cells were harvested after 6 d of culture, and DCs were purified using anti-CD11c conjugated MACS microbeads and magnetic separation columns (Miltenyi Biotec, Bergisch Gladbach, Germany). The purity of DCs was assessed by FACS analysis and ranged from 70% to 90% CD11c⁺. Cells were incubated in an atmosphere containing 5% CO₂ at 37 °C.

HeLa cells were infected with invasive *Salmonella* prepared according to previous studies [32]. RAW264.7 and DCs were infected with stationary phase *Salmonella* opsonized in 20% normal human serum for 30 min. An incubation of 10 min at 37 °C in 5% CO₂ was performed once bacteria were applied to cell cultures to allow for internalization. Following this, the infecting medium was aspirated, cells were washed four times with PBS, fresh cell culture medium containing gentamicin (100 μ g ml⁻¹) was added, and incubation at 37 °C in 5% CO₂ was continued. At 1 h post-infection, cell culture medium was removed and the cells were again washed four times with PBS. The cells were then lysed in a solution of Triton X-100 (1% v/v) and sodium dodecyl sulfate (0.1% w/v) to release intracellular bacteria. Samples were diluted and assayed for resolution.

Mouse infections. Female C57BL/6 mice (6–10 wk of age) were purchased from Jackson Laboratory (Bar Harbor, Maine, United States), and housed in the animal facility at the University of British Columbia in direct accordance with guidelines drafted by the University of British Columbia's Animal Care Committee and the Canadian Council on the Use of Laboratory Animals. For ileal loop experiments, bacterial inocula of approximately 10^7 colony-forming units were prepared in 100 μ l, and the resolution status of the strain was confirmed directly before inoculation. Ileal loop experiments were modified from those previously described [1]. In brief, mice were anaesthetized by intraperitoneal injection of ketamine and xylazine. Following a midline abdominal incision, the small bowel was exposed and the bowel was ligated twice, close to the cecum, to create a loop approximately 4 cm in length into which the inoculum was injected. The bowel was then returned to the abdominal cavity and the incision closed with discontinuous sutures. At given time points, the mice were euthanized and tissues collected for bacterial enumeration and RIVET. The intestinal lumen was rinsed with PBS to remove non-adherent bacteria. Tissues were homogenized in PBS using a Polytron homogenizer (Kinematica, Lucerne, Switzerland).

Immunohistochemistry. Tissues were fixed for 3 h in 3% paraformaldehyde, and washed three times with PBS prior to cryoembedding and sectioning. Sections were fixed in acetone at -20 °C for 10 min and then air dried at room temperature. Tissue sections were outlined with a wax pen and blocked 30 min in 10% goat serum in PBS-BSA at room temperature. Sections were then washed three times in PBS-BSA prior to incubation for 30 min at room temperature with anti-*Salmonella* LPS antiserum. Sections were then washed three times in PBS-BSA prior to incubation with appropriate Alexafluor 568-conjugated anti-rabbit antibodies. Sections were then washed three times in PBS-BSA, incubated with Alexafluor 488-labeled phalloidin, washed a further three times in PBS-BSA, and then incubated in DAPI. Imaging was performed on a Nikon (Tokyo, Japan) TE2000 inverted microscope equipped with a Bio-Rad (Hercules, California, United States) Radiance 2000 confocal

scan head and a two-photon laser source using a plan apochromat 40× 1.3 N.A. objective.

Acknowledgments

We thank members of the Finlay and Vallance laboratories for advice with experiments and critical reading of this manuscript. We also thank Andrew Camilli for providing reagents and advice on the use of RIVET and Ifor Beacham for advice on silent genes in *Salmonella*. NFB is a Michael Smith Foundation for Health Research (MSFHR) fellow, and BKC is a MSFHR and Canadian Institutes of Health Research (CIHR) fellow. BAC is supported by CIHR and MSFHR studentships.

References

- Jones BD, Ghori N, Falkow S (1994) *Salmonella typhimurium* initiates murine infection by penetrating and destroying the specialized epithelial M cells of the Peyer's patches. *J Exp Med* 180: 15–23.
- Vazquez-Torres A, Jones-Carson J, Baumler AJ, Falkow S, Valdivia R, et al. (1999) Extraintestinal dissemination of *Salmonella* by CD18-expressing phagocytes. *Nature* 401: 804–808.
- Rescigno M, Urbano M, Valzasina B, Francolini M, Rotta G, et al. (2001) Dendritic cells express tight junction proteins and penetrate gut epithelial monolayers to sample bacteria. *Nat Immunol* 2: 361–367.
- Richter-Dahlfors A, Buchan AM, Finlay BB (1997) Murine salmonellosis studied by confocal microscopy: *Salmonella typhimurium* resides intracellularly inside macrophages and exerts a cytotoxic effect on phagocytes in vivo. *J Exp Med* 186: 569–580.
- Salcedo SP, Noursadeghi M, Cohen J, Holden DW (2001) Intracellular replication of *Salmonella typhimurium* strains in specific subsets of splenic macrophages in vivo. *Cell Microbiol* 3: 587–597.
- Hueck CJ (1998) Type III protein secretion systems in bacterial pathogens of animals and plants. *Microbiol Mol Biol Rev* 62: 379–433.
- Wallis TS, Galvov EE (2000) Molecular basis of *Salmonella*-induced enteritis. *Mol Microbiol* 36: 997–1005.
- Penheiter KL, Mathur N, Giles D, Fahlen T, Jones BD (1997) Non-invasive *Salmonella typhimurium* mutants are avirulent because of an inability to enter and destroy M cells of ileal Peyer's patches. *Mol Microbiol* 24: 697–709.
- Galan JE, Curtiss R III (1989) Cloning and molecular characterization of genes whose products allow *Salmonella typhimurium* to penetrate tissue culture cells. *Proc Natl Acad Sci U S A* 86: 6383–6387.
- Hensel M, Shea JE, Gleeson C, Jones MD, Dalton E, et al. (1995) Simultaneous identification of bacterial virulence genes by negative selection. *Science* 269: 400–403.
- Shea JE, Hensel M, Gleeson C, Holden DW (1996) Identification of a virulence locus encoding a second type III secretion system in *Salmonella typhimurium*. *Proc Natl Acad Sci U S A* 93: 2593–2597.
- Ochman H, Soncini FC, Solomon F, Groisman EA (1996) Identification of a pathogenicity island required for *Salmonella* survival in host cells. *Proc Natl Acad Sci U S A* 93: 7800–7804.
- Bispham J, Tripathi BN, Watson PR, Wallis TS (2001) *Salmonella* pathogenicity island 2 influences both systemic salmonellosis and *Salmonella*-induced enteritis in calves. *Infect Immun* 69: 367–377.
- Tsolis RM, Adams LG, Ficht TA, Baumler AJ (1999) Contribution of *Salmonella typhimurium* virulence factors to diarrheal disease in calves. *Infect Immun* 67: 4879–4885.
- Hapfelmeier S, Stecher B, Barthel M, Kremer M, Muller AJ, et al. (2005) The *Salmonella* pathogenicity island (SPI)-2 and SPI-1 type III secretion systems allow *Salmonella* serovar *typhimurium* to trigger colitis via MyD88-dependent and MyD88-independent mechanisms. *J Immunol* 174: 1675–1685.
- Barthel M, Hapfelmeier S, Quintanilla-Martinez L, Kremer M, Rohde M, et al. (2003) Pretreatment of mice with streptomycin provides a *Salmonella enterica* serovar Typhimurium colitis model that allows analysis of both pathogen and host. *Infect Immun* 71: 2839–2858.
- Waterman SR, Holden DW (2003) Functions and effectors of the *Salmonella* pathogenicity island 2 type III secretion system. *Cell Microbiol* 5: 501–511.
- Valdivia RH, Falkow S (1997) Fluorescence-based isolation of bacterial genes expressed within host cells. *Science* 277: 2007–2011.
- Cirillo DM, Valdivia RH, Monack DM, Falkow S (1998) Macrophage-dependent induction of the *Salmonella* pathogenicity island 2 type III secretion system and its role in intracellular survival. *Mol Microbiol* 30: 175–188.
- Deiwick J, Nikolaus T, Erdogan S, Hensel M (1999) Environmental regulation of *Salmonella* pathogenicity island 2 gene expression. *Mol Microbiol* 31: 1759–1773.
- Lee AK, Detweiler CS, Falkow S (2000) OmpR regulates the two-component system SsrA-SsrB in *Salmonella* pathogenicity island 2. *J Bacteriol* 182: 771–781.
- Chakravorty D, Rohde M, Jager L, Deiwick J, Hensel M (2005) Formation of a novel surface structure encoded by *Salmonella* Pathogenicity Island 2. *EMBO J* 24: 2043–2052.
- Camilli A, Beattie DT, Mekalanos JJ (1994) Use of genetic recombination as a reporter of gene expression. *Proc Natl Acad Sci U S A* 91: 2634–2638.
- Lee SH, Hava DL, Waldor MK, Camilli A (1999) Regulation and temporal expression patterns of *Vibrio cholerae* virulence genes during infection. *Cell* 99: 625–634.
- Ayabe T, Satchell DP, Wilson CL, Parks WC, Selsted ME, et al. (2000) Secretion of microbicidal alpha-defensins by intestinal Paneth cells in response to bacteria. *Nat Immunol* 1: 113–118.
- Coombes BK, Brown NF, Valdez Y, Brummell JH, Finlay BB (2004) Expression and secretion of *Salmonella* pathogenicity island-2 virulence genes in response to acidification exhibit differential requirements of a functional type III secretion apparatus and SsaL. *J Biol Chem* 279: 49804–49815.
- Innes D, Beacham IR, Beven CA, Douglas M, Laird MW, et al. (2001) The cryptic *ushA* gene (*ushA(c)*) in natural isolates of *Salmonella enterica* (serotype Typhimurium) has been inactivated by a single missense mutation. *Microbiology* 147: 1887–1896.
- Donnenberg MS, Kaper JB (1991) Construction of an *eae* deletion mutant of enteropathogenic *Escherichia coli* by using a positive-selection suicide vector. *Infect Immun* 59: 4310–4317.
- Kehres DG, Zaharik ML, Finlay BB, Maguire ME (2000) The NRAMP proteins of *Salmonella typhimurium* and *Escherichia coli* are selective manganese transporters involved in the response to reactive oxygen. *Mol Microbiol* 36: 1085–1100.
- Knodler LA, Celli J, Hardt WD, Vallance BA, Yip C, et al. (2002) *Salmonella* effectors within a single pathogenicity island are differentially expressed and translocated by separate type III secretion systems. *Mol Microbiol* 43: 1089–1103.
- Lutz MB, Kukutsch N, Ogilvie AL, Rossner S, Koch F, et al. (1999) An advanced culture method for generating large quantities of highly pure dendritic cells from mouse bone marrow. *J Immunol Methods* 223: 77–92.
- Steele-Mortimer O, Meresse S, Gorvel JP, Toh BH, Finlay BB (1999) Biogenesis of *Salmonella typhimurium*-containing vacuoles in epithelial cells involves interactions with the early endocytic pathway. *Cell Microbiol* 1: 33–49.
- Wray C, Sojka WJ (1978) Experimental *Salmonella typhimurium* infection in calves. *Res Vet Sci* 25: 139–143.
- Osorio CG, Crawford JA, Michalski J, Martinez-Wilson H, Kaper JB, et al. (2005) Second-generation recombination-based in vivo expression technology for large-scale screening for *Vibrio cholerae* genes induced during infection of the mouse small intestine. *Infect Immun* 73: 972–980.
- Dennis JJ, Zylstra GJ (1998) Improved antibiotic-resistance cassettes through restriction site elimination using Pfu DNA polymerase PCR. *Biotechniques* 25: 772–774–776.
- Yanisch-Perron C, Vieira J, Messing J (1985) Improved M13 phage cloning vectors and host strains: Nucleotide sequences of the M13mp18 and pUC19 vectors. *Gene* 33: 103–119.