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2014. Spatial Segregation of Virulence Gene Expression during Acute Enteric Infection with *Salmonella enterica* serovar Typhimurium . mBio 5(1): .
doi:10.1128/mBio.00946-13.

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RESEARCH ARTICLE

Spatial Segregation of Virulence Gene Expression during Acute Enteric Infection with *Salmonella enterica* serovar Typhimurium

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ABSTRACT To establish a replicative niche during its infectious cycle between the intestinal lumen and tissue, the enteric pathogen *Salmonella enterica* serovar Typhimurium requires numerous virulence genes, including genes for two type III secretion systems (T3SS) and their cognate effectors. To better understand the host-pathogen relationship, including early infection dynamics and induction kinetics of the bacterial virulence program in the context of a natural host, we monitored the subcellular localization and temporal expression of T3SS-1 and T3SS-2 using fluorescent single-cell reporters in a bovine, ligated ileal loop model of infection. We observed that the majority of bacteria at 2 h postinfection are flagellated, express T3SS-1 but not T3SS-2, and are associated with the epithelium or with extruding enterocytes. In epithelial cells, *S. Typhimurium* cells were surrounded by intact vacuolar membranes or present within membrane-compromised vacuoles that typically contained numerous vesicular structures. By 8 h postinfection, T3SS-2-expressing bacteria were detected in the lamina propria and in the underlying mucosa, while T3SS-1-expressing bacteria were in the lumen. Our work identifies for the first time the temporal and spatial regulation of T3SS-1 and -2 expression during an enteric infection in a natural host and provides further support for the concept of cytosolic *S. Typhimurium* in extruding epithelium as a mechanism for reseeding the lumen.

IMPORTANCE The pathogenic bacterium *Salmonella enterica* serovar Typhimurium invades and persists within host cells using distinct sets of virulence genes. Genes from *Salmonella* pathogenicity island 1 (SPI-1) are used to initiate contact and facilitate uptake into nonphagocytic host cells, while genes within SPI-2 allow the pathogen to colonize host cells. While many studies have identified bacterial virulence determinants in animal models of infection, very few have focused on virulence gene expression at the single-cell level during an *in vivo* infection. To better understand when and where bacterial virulence factors are expressed during an acute enteric infection of a natural host, we infected bovine jejunal-ileal loops with *S. Typhimurium* cells harboring fluorescent transcriptional reporters for SPI-1 and -2 (*PinvF* and *PssaG*, respectively). After a prescribed time of infection, tissue and luminal fluid were collected and analyzed by microscopy. During early infection (≤ 2 h), bacteria within both intact and compromised membrane-bound vacuoles were observed within the epithelium, with the majority expressing SPI-1. As the infection progressed, *S. Typhimurium* displayed differential expression of the SPI-1 and SPI-2 regulons, with the majority of tissue-associated bacteria expressing SPI-2 and the majority of lumen-associated bacteria expressing SPI-1. This underscores the finding that *Salmonella* virulence gene expression changes as the pathogen transitions from one anatomical location to the next.

Received 5 November 2013 Accepted 23 December 2013 Published 4 February 2014

Citation Laughlin RC, Knodler LA, Barhoumi R, Payne HR, Wu J, Gomez G, Pugh R, Lawhon SD, Bäuml AJ, Steele-Mortimer O, Adams LG. 2014. Spatial segregation of virulence gene expression during acute enteric infection with *Salmonella enterica* serovar Typhimurium. *mBio* 5(1):e00946-13. doi:10.1128/mBio.00946-13.

Invited Editor John Gunn, The Ohio State University **Editor** Stefan Kaufmann, Max Planck Institute for Infection Biology

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The intestinal mucosa is located at an important crossroads of dynamic interactions between the intestinal microbiota, vital absorptive cells, transient as well as resident immune cells, and pathogenic organisms. Intestinal villi extend into the luminal milieu and provide a selective barrier against luminal contents, remove injured or aged epithelial cells via controlled sloughing or extrusion, educate naive immune cells to intestinal symbiotic bacteria, and monitor the local environment for pathogenic threats (1, 2). Appropriate immunologic and cellular responses to the

autochthonous intestinal microbial populations, as well as general luminal conditions, are important for the health of the organism.

Disruption of the autochthonous population plays an important role in the establishment and propagation of infection for several pathogens of the alimentary tract, of which *Salmonella enterica* serovar Typhimurium has received significant attention (3–7). This member of the *Enterobacteriaceae* family is a food-borne pathogen that elicits clinically and pathologically similar disease outcomes in humans and cattle (8–10). Animal models for

this localized gastroenteric infection include neonatal bovines and streptomycin-treated mice (11, 12). In the bovine model, bacterial invasion of intestinal tissue occurs as early as 15 min after exposure and typically affects phagocytic and nonphagocytic cells (13). Ileal Peyer's patch phagocytes, likely tissue-associated dendritic cells and M cells, capture and deliver invading *S. Typhimurium* cells to the local mesenteric lymph node to educate and recruit T cells for return to the site of infection (14, 15). The initial hours of acute *Salmonella* infection in humans and cattle are similarly characterized by polymorphonuclear cell (PMN) infiltration into the lamina propria and then PMN efflux and transit through the intestinal epithelium into the lumen, luminal fluid accumulation, epithelial cell shedding, and villus blunting (16, 17). Similar features of mucosal damage have also been described for *S. Typhimurium* infection of rabbits and rhesus monkeys (18, 19).

S. Typhimurium employs two type III secretion systems (T3SS) to mediate their interactions with host cells (20, 21). T3SS-1 and T3SS-2 are encoded in *Salmonella* pathogenicity islands 1 and 2 (SPI-1 and SPI-2, respectively). Genetic deletion of SPI-1 or SPI-2 can abrogate the virulence and ability of *Salmonella* to invade, colonize, or replicate within host cells (10, 11, 22, 23). The SPI-1 and SPI-2 regulons are induced under different environmental conditions. Expression of the SPI-1 regulon is controlled by numerous proteins, including *invF* and *hilA*, and induced extracellularly (24), consistent with its role in invasion of nonphagocytic host cells, such as enterocytes. Proteins encoded by genes within the SPI-1 regulon include structural components of the T3SS-1 apparatus and several type III effectors that modulate macropinocytosis at the plasma membrane, trafficking of the nascent *Salmonella*-containing vacuole (SCV), and intracellular replication (25). Following invasion into nonphagocytic cells, *S. Typhimurium* down-regulates SPI-1 and induces the SPI-2 regulon (26). SPI-2-encoded T3SS-2 translocates effector proteins that are required for maturation and maintenance of the SCV (25). Although the SCV has been considered the predominant site of intracellular replication for *Salmonella*, recent studies have identified a distinct population of *S. Typhimurium* cells that hyper-replicate in the cytosol of epithelial cells (27, 28). In contrast to the SPI-2-induced vacuolar population, these cytosolic bacteria are induced for SPI-1 and flagellated. In a polarized epithelial cell model, these "invasion-primed" intracellular bacteria are released into the extracellular milieu when the host cell is extruded from the monolayer. In murine gall bladders, extruding epithelial cells were shown to also contain cytosolic invasion-primed *S. Typhimurium* cells (27). In addition, enterocytes containing large numbers of *S. Typhimurium* cells have been observed within rabbit ileal mucosa (18) and chick ileocecal mucosa (29). It has been proposed that this bacterial population may play an important role in cell-to-cell transmission and/or dissemination *in vivo* (27).

Although it is clear that SPI-1 and, to a lesser extent, SPI-2 are required for the induction of pathological changes during acute enteric infection (10, 30), the timing and location of bacterial gene expression *in vivo* have received little attention and are poorly understood. Here we have addressed this question using the well-established neonatal bovine ileal loop model. Calves were infected with *S. Typhimurium* harboring transcriptional fusions to representative genes from SPI-1 (*invF*) or SPI-2 (*ssaG*) for various times before tissue was harvested for transmission electron and confocal microscopy. We report the presence of both vacuole membrane-bound and -compromised bacteria within the epithelium and dis-

semination of invasion-primed *S. Typhimurium* by epithelial extrusion, particularly early during infection. In addition, our single-cell expression analysis revealed a distinct temporal and spatial segregation of SPI-1- and SPI-2-positive bacteria during intestinal colonization.

RESULTS AND DISCUSSION

***S. Typhimurium* interactions with the epithelium during early infection.** To observe interactions between *S. Typhimurium* and the host epithelium during early infection, bovine ileal loops were harvested at 2 h postinoculation (p.i.) and processed for transmission electron microscopy (TEM). Analysis of infected tissue revealed numerous examples of *S. Typhimurium* cells colonizing enterocytes, goblet cells, and other cells within the epithelium (Fig. 1; see also Fig. S1 in the supplemental material). In most instances, infected host cells remained part of the epithelium; however, examples of epithelial cells with remnant microvilli, seemingly undergoing extrusion and containing *S. Typhimurium* cells, were also observed (Fig. 1C and E, arrows). We further noted that while many intracellular bacteria were enclosed by a vacuolar membrane (Fig. 1B and S2A and B), others were not (Fig. 1C to F and S2C to E). Instead, the integrity of the vacuole around these bacteria appeared to be compromised (defined as >25% of the bacterial surface not being associated with a vacuolar membrane), affording access to the host cytosol. Approximately 13% of *S. Typhimurium* cells were observed in such compromised vacuoles ($n = 173$ bacteria). By electron microscopy, we were unable to identify a bovine epithelial cell laden with cytosolic *S. Typhimurium* cells, as has been described for chick ileocecal mucosa (31) and polarized human intestinal epithelial monolayers (27). However, our TEM analysis does support previous observations of the presence of *S. Typhimurium* in distinct intracellular compartments in epithelial cells and cellular extrusion acting as a mechanism for bacterial dissemination into the lumen (18, 27). TEM analysis of infected tissue samples from 8 h p.i. revealed *S. Typhimurium* bacteria located in the lamina propria within intact vacuoles, compromised vacuoles, or vacuoles apparently free of any discernible host membrane (Fig. S3).

Further TEM analysis of intracellular *S. Typhimurium* within intact and compromised vacuoles from 2-h-p.i. samples revealed the presence of numerous vesicular bodies within the SCV lumen (Fig. S1). These vesicles ranged in size from <50 nm to >200 nm. Intracellular and extracellular Gram-negative bacteria are known to secrete spherical vesicles, called outer membrane vesicles (OMV), that are 50 to 250 nm in diameter and often have an electron-dense luminal content by electron microscopy, consistent with what we report here (32). Budding or recently formed OMV originating from the pathogen were found associated with all intracellular *S. Typhimurium* cells analyzed (Fig. S1, arrowheads), complementing the findings from a previous *in vivo* study of a human *Salmonella* isolate in chicken ileum (33). OMV were typically found free within the SCV lumen (Fig. S1J) or adjacent to or apparently spanning the SCV membrane (Fig. S1K, arrowhead, and S2C, arrow). Larger, more-electron-lucent membrane structures were also noted within the SCV (Fig. S1J and K and S2E, chevrons), sometimes apparently fusing with or blebbing from the vacuolar membrane (indicated in Fig. S2E, chevron). It is unclear if these larger vesicles originate from the pathogen or the host.

SPI-1 and -2 expression during acute infection. To further our understanding of bacterial virulence gene expression during

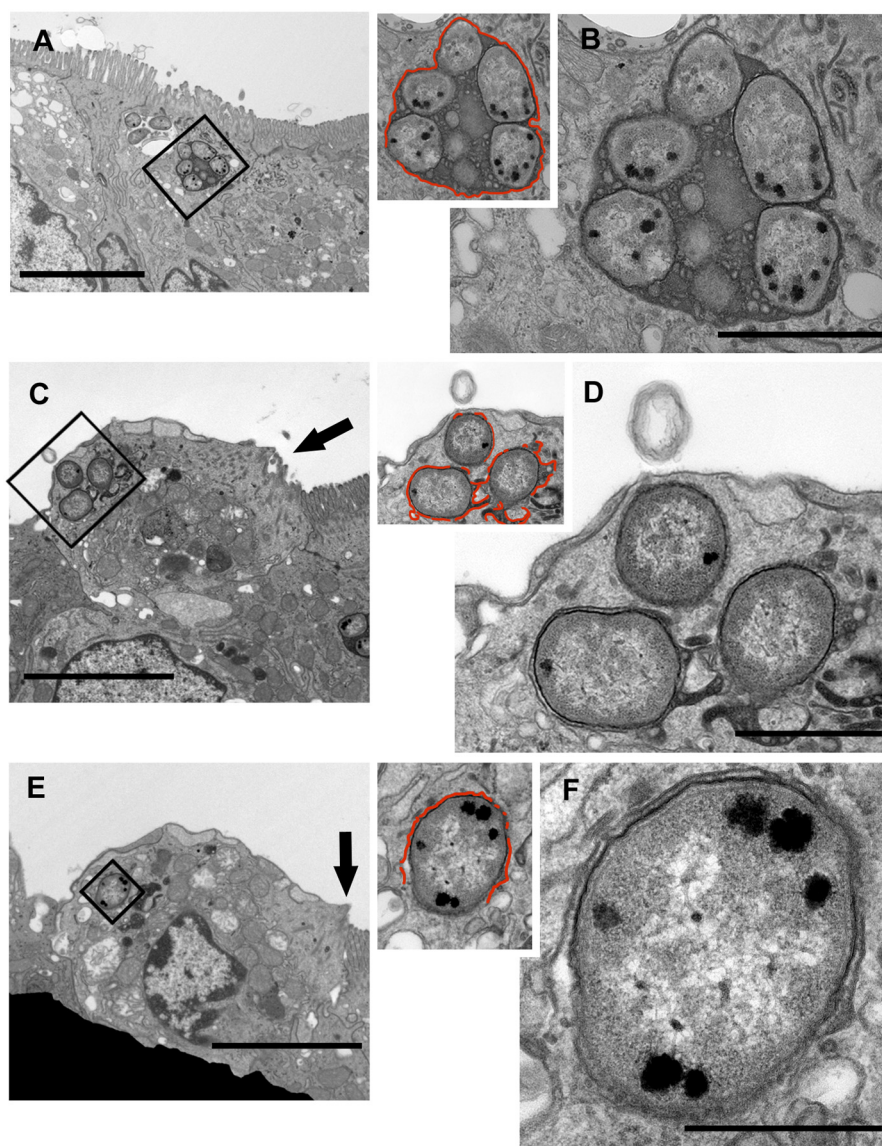


FIG 1 In enterocytes, *S. Typhimurium* cells are surrounded by intact and compromised vacuolar membranes. Representative transmission electron micrograph of infected epithelial cells at 2 h p.i. The vacuolar membrane is highlighted with a red line in the inset. (A and B) Series of images demonstrating *S. Typhimurium* cells enclosed within an intact membrane (C to F). Infected epithelial cells likely undergoing cellular extrusion. At least one bacterium in each cell has a compromised vacuolar membrane (indicated by discontinuous red lines in the insets). (C and E) Arrows indicate remnant microvilli. Bars, 5 μm (A, C, and E), 1 μm (B, D, and G), and 0.5 μm (F).

acute infection *in vivo*, we used *S. Typhimurium* harboring reporter plasmids for either the SPI-1 or the SPI-2 regulon in the bovine ileal loop model. The plasmids encode a destabilized green fluorescent protein (GFP) variant, GFP[LVA] (half-time, ~40 min), that is under the control of gene promoters from either SPI-1 (*P_{invF}*) or SPI-2 (*P_{ssaG}*). *InvF* is a transcriptional activator of SPI-1 (34), whereas *SsaG* is a structural component of T3SS-2 (35). In broth cultures, wild-type bacteria harboring pMPMA3 Δ Plac-*P_{invF}*-GFP[LVA] exhibited green fluorescence when grown under SPI-1-inducing conditions (Luria-Bertani [LB]–Miller broth, 3.5 h) but not under SPI-2-inducing conditions (low phosphate magnesium [LPM], pH 5.8, 14 h) (Fig. S4A). Fluorescence was not observed under SPI-1-inducing conditions upon deletion of *hilA*, a master regulator of the SPI-1 regulon (36).

Conversely, bacteria harboring pMPMA3 Δ Plac-*P_{ssaG}*-GFP[LVA] were fluorescent only when grown under SPI-2-inducing conditions, and this was dependent on *ssrB* (Fig. S4), part of a two-component regulatory system that is absolutely required for expression of the SPI-2 regulon (37). Upon examination of infected ileal loop tissues, we found that the intrinsic fluorescence of GFP[LVA] was often too weak to detect by confocal microscopy. To circumvent this, we used rabbit polyclonal anti-GFP antibodies to amplify the fluorescence signal associated with individual bacteria. Mammalian tissue culture invasion assays were used to assess the expression kinetics and frequencies of GFP-positive bacteria and to determine that intrinsic fluorescence and antibody amplification of the GFP[LVA] signal were comparable to those of methods of reporter activity quantification

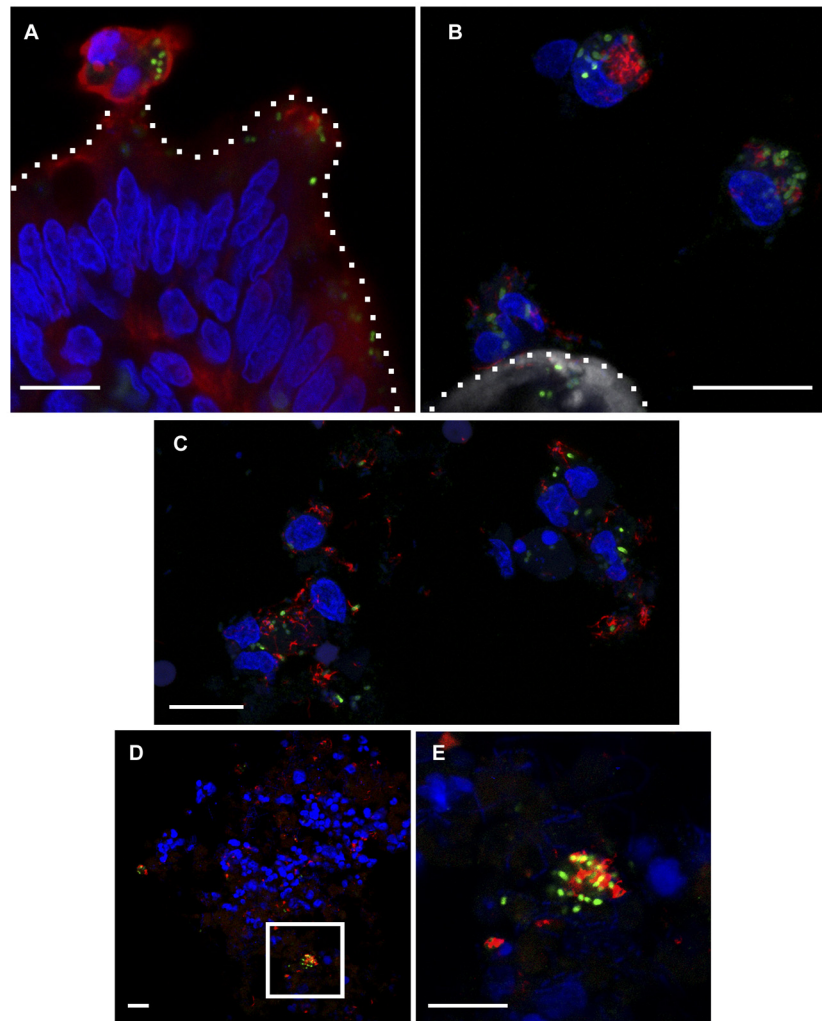


FIG 2 *PinvF*-positive, flagellated *S. Typhimurium* bacteria are associated with host cells proximal to villus tips and in the lumen. Ligated jejunal-ileal loops were infected with *S. Typhimurium* cells harboring *PinvF*-GFP[LVA] for 2 h. (A) Confocal image showing an extruded epithelial cell (cytokeratin 8 positive) containing numerous *PinvF*-GFP-positive bacteria. Green, anti-GFP; red, anti-cytokeratin 8; blue, DNA. (B) Confocal image showing several bovine cells adjacent to a villus tip containing a significant burden of flagellin-positive and *PinvF*-GFP positive *Salmonella* cells. Green, anti-GFP; red, anti-FliC; shades of gray, phalloidin for actin; blue, DNA. For A and B, dotted lines indicate the apical surface of the villus tip. (C) Confocal image of cells adjacent to villus tips revealing flagellin-positive and *PinvF*-GFP-positive bacteria intimately associated with bovine cells sloughed from the tissue. Green, anti-GFP; red, anti-FliC; blue, DNA. (D) Confocal image of luminal fluid content showing *PinvF*-GFP-positive, flagellated *Salmonella* bacteria found in clusters within a heterogeneous milieu of eukaryotic and prokaryotic cells. Green, anti-GFP; red, anti-FliC; blue, DNA. (E) Enlarged inset from panel D. Bars, 10 μm .

(Fig. S4B). Furthermore, at both 2 h and 8 h p.i. in the *in vivo* model, luminal fluid accumulation and bacterial burdens in tissue, mucus, and fluid samples from loops infected with *S. Typhimurium* harboring the GFP reporter plasmids were consistent with those of wild type (WT)-infected loops (Fig. S5). Additionally, plasmids were retained throughout the duration of an 8-h infection (Fig. S5). Collectively, these experiments validated the use of pMPMA3 Δ Plac-*PinvF*-GFP[LVA] and pMPMA3 Δ Plac-*PssaG*-GFP[LVA] as accurate transcriptional reporters of the SPI-1 and SPI-2 regulons, respectively, *in vitro*.

Confocal microscopy analysis of villus tips and cells immediately adjacent to tissue revealed numerous instances of extruding or sloughed cells positive for staining by cytokeratin 8, an intermediate filament protein found in epithelial cells containing *PinvF*-positive *S. Typhimurium* cells at 2 h (Fig. 2A). In bovine

cells found adjacent to villus tips, many of the bacteria also immunostained for flagellin (FliC) (Fig. 2B). *Salmonella* has previously been associated with epithelial cell extrusion in polarized monolayers, rabbit ileal loops, and mouse gall bladders (18, 27). Such extrusion has been postulated as a dissemination mechanism, since bacteria within these dying cells are induced for SPI-1 and flagellated (18). Our observations provide further evidence for the presence SPI-1-induced, flagellated *S. Typhimurium* bacteria associating with host cells sloughing from villus tips *in vivo*. Infected cells adjacent to the villus often contained multiple regions positively stained for DNA, suggesting either nuclear degradation of a single cell or multiple cells clumping together (Fig. 2A and B). Extruding epithelial cells or near-tissue *PinvF*-positive bacteria were rarely observed at 8 h p.i., likely due to the significant villus blunting and immune responses seen at this later time point.

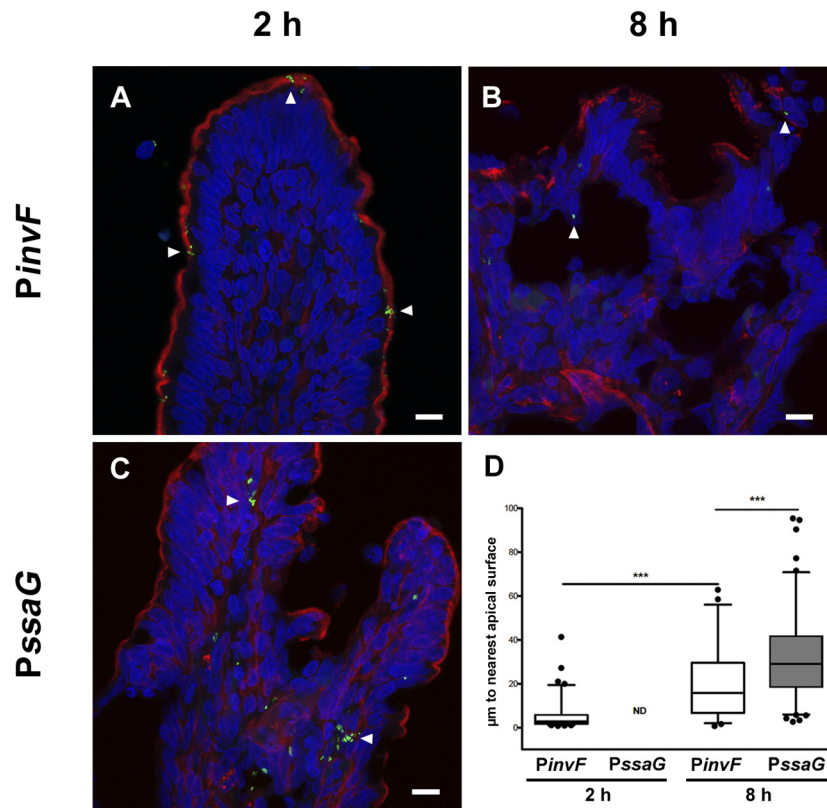


FIG 3 Temporal and spatial distribution of *PinvF*- and *PssaG*-positive *S. Typhimurium* cells. Ligated jejunal-ileal loops were infected with *S. Typhimurium* cells harboring either *PinvF*-GFP[LVA] or *PssaG*-GFP[LVA]. (A to C) Representative confocal images of tissue stained for GFP (green), phalloidin (for actin, red), and DNA (blue), with GFP-positive bacteria indicated by white arrowheads. (A) *PinvF*-GFP[LVA] infection at 2 h p.i.; (B) *PinvF*-GFP[LVA] infection at 8 h p.i.; (C) *PssaG*-GFP[LVA] infection at 8 h p.i.; (D) box-and-whisker plot of tissue-associated *PinvF*- or *PssaG*-GFP[LVA]-positive *S. Typhimurium* cells and their distance from the nearest apical surface (assessed from phalloidin staining). Whiskers represent the 5-to-95% confidence interval. ● symbols represent outlier data points. ***, $P < 0.001$. ND, not determined. Numbers of GFP-positive bacteria assessed for distance to the luminal surface were as follows: 82 bacteria at 2 h for *PinvF*-GFP[LVA]-infected cells, 50 bacteria at 8 h for *PinvF*-GFP[LVA]-infected cells, and 199 bacteria at 8 h for *PssaG*-GFP[LVA]-infected cells (values are from samples from at least three animals).

Immunostaining of the luminal fluid samples for GFP and FliC and fluorescent staining of DNA with DAPI (4',6-diamidino-2-phenylindole) revealed a heterogeneous population of *PinvF*-positive and -negative *S. Typhimurium* cells, other bacterial species, and host cells (Fig. 2D and E). Interestingly, *PinvF*-positive *S. Typhimurium* appeared to aggregate in this environment (Fig. 2E) and also immunostained for flagellin (Fig. 2B and C). Stecher et al. used a reporter plasmid expressing GFP $mut2$ under the control of the *fliC* promoter in a streptomycin-treated mouse model of colitis and showed there a gradation of *fliC* expression by *S. Typhimurium* in the gut at 1 day p.i. (38). Approximately half of the bacteria in the cecal lumen were GFP positive, whereas almost all were GFP positive (92%) when in close proximity to the cecal epithelium. In an independent study of flagellin expression during systemic salmonellosis, Cummings et al. used a *fliC::gfp mut3* reporter to show that ~60% of *S. Typhimurium* cells in the Peyer's patches transcribed *fliC* at 7 days p.i. but that none in the mesenteric lymph nodes or spleen were GFP positive (39). Our finding that luminal *S. Typhimurium* cells are flagellated supports the concept of an anatomical restriction of FliC expression to the gut.

We did not detect any *PssaG*-positive bacteria in luminal fluid samples at either 2 h or 8 h p.i. (Fig. S6). This is in contrast to the findings of a recent report that identified a very small population

(1.34% of the total) of SPI-2-induced *S. Typhimurium* cells in the lumen of the cecum during a mouse model of colitis. Approximately two-thirds of these bacteria were extracellular; the remaining cells were within luminal CD18⁺ neutrophils (40). Using recombinase-based *in vivo* expression technology (RIVET), another group has also reported that SPI-2 is expressed prior to bacterial penetration of the epithelial layer in a murine model of salmonellosis (41). The considerable differences in pathology and host response between the bovine and mouse models of salmonellosis, in addition to infection parameters and detection methods, may account for the discrepancy between these observations.

Virulence gene expression within gut tissue. During enteric infections, many different cell types are targets for *S. Typhimurium*, including enterocytes, goblet cells, macrophages, and dendritic cells (42, 43). However, it is unclear when these particular host cell-bacterium interactions occur during the course of acute infection. To monitor the physical location of *PinvF*- and *PssaG*-induced *S. Typhimurium* cells, GFP-positive bacteria were quantified in two ways by confocal microscopy: (i) by determining the distance of tissue-associated bacteria from the nearest apical surface, denoted by phalloidin staining (Fig. 3), and (ii) by categorizing the subtissue localization of GFP-positive bacteria (Fig. 4).

At 2 h p.i., *PinvF*-positive bacteria were largely associated with

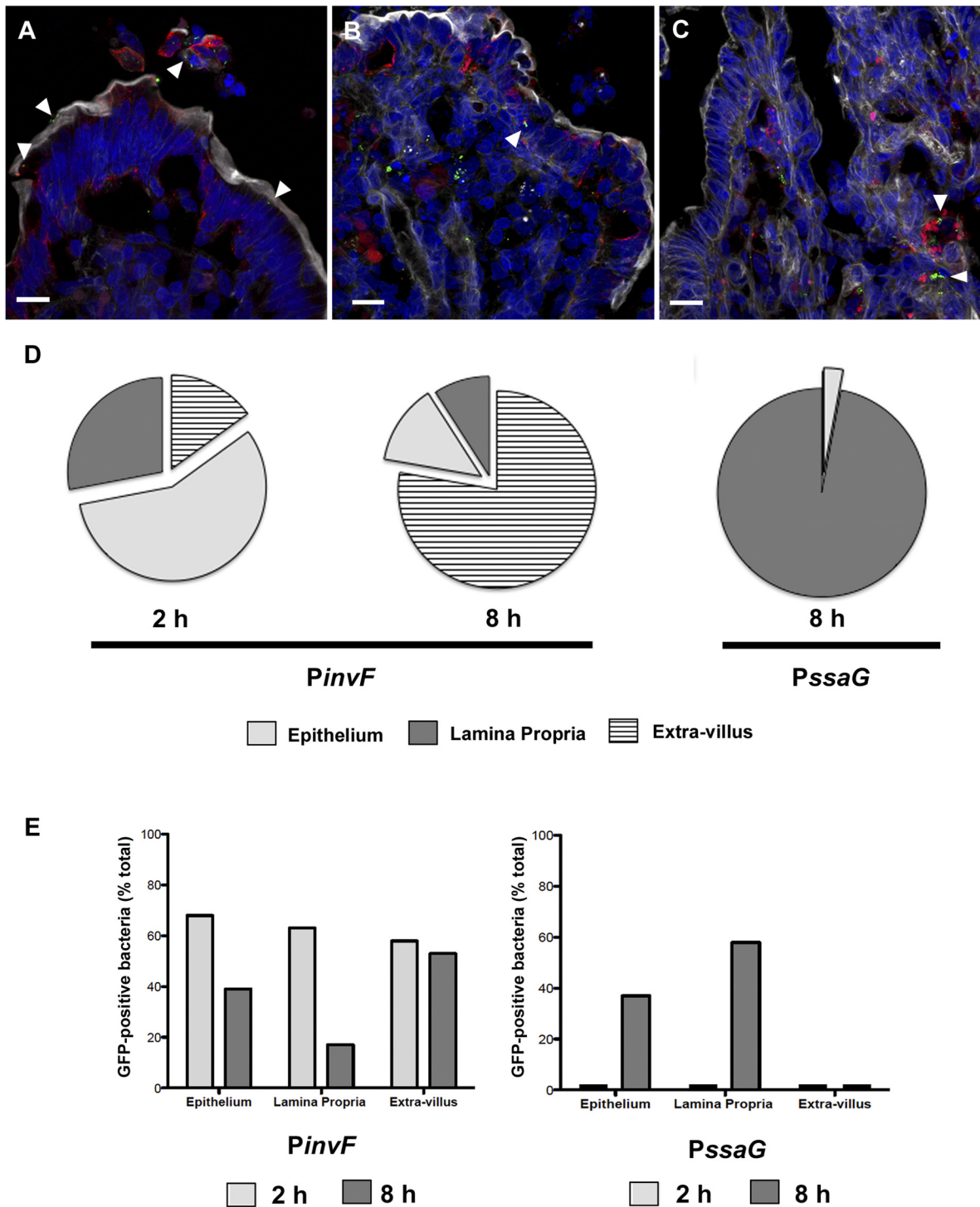


FIG 4 *PinvF*- and *PssaG*-positive *S. Typhimurium* bacteria predominantly associate with different host cell types. Ligated jejunal-ileal loops were infected with *S. Typhimurium* cells harboring either *PinvF*-GFP[LVA] or *PssaG*-GFP[LVA] and processed for confocal microscopy. (A) Representative image of *PinvF*-GFP[LVA] infection at 2 h p.i. (B and C) *PssaG*-GFP[LVA] infection at 8 h p.i. (A and B) Green, anti-GFP; red, anti-cytokeratin 8; shades of gray, phalloidin for actin; blue, DNA. (C) Green, anti-GFP; red, anti-HLA-DR α ; shades of gray, phalloidin for actin; blue, DNA. Bars = 10 μ m. Arrowheads indicate GFP-positive *S. Typhimurium* bacteria within epithelial cells (A and B) or phagocytes (C). (D) The locations of *PinvF*- and *PssaG*-positive bacteria were scored as in the epithelium (light gray), extravillus (hatched), or lamina propria (dark gray). Percentages of bacteria in the extravillus/epithelium/lamina propria were as follows: 15/57/28% for *PinvF*-GFP[LVA]-infected specimens at 2 h ($n = 439$), 77/13/9% for *PinvF*-GFP[LVA]-infected specimens at 8 h ($n = 298$), and 0/3/97% for *PssaG*-GFP[LVA]-infected specimens at 8 h ($n = 308$). (E) Proportions of total bacteria positive for either *PinvF*- or *PssaG*-GFP in the epithelium, lamina propria, or extravillous space at 2 h (light gray) or 8 h p.i. (dark gray). Percentages of *PinvF*-GFP-positive bacteria at 2 h/8 h were as follows: 68/39% in the epithelium, 63/17% in the lamina propria, and 58/53% in the extravillous space. Percentages of *PssaG*-GFP-positive bacteria at 2 h/8 h were as follows: 0/37% in the epithelium, 0/58% in the lamina propria, and 0/0% in the extravillous space.

the epithelium (57%) (Fig. 4A and D) and typically situated <10 μm from the nearest apical surface (Fig. 3A and D). The remaining *PinvF*-positive bacteria were distributed between the extravillous milieu (15%) and the lamina propria (28%), implying that the SPI-1 regulon is expressed after the initial interaction of bacteria with enterocytes. *PinvF*-positive bacteria were still detected at 8 h p.i., (Fig. 3B), although at a reduced frequency compared to that at 2 h p.i., indicating that at a relatively late stage in acute enteritis, SPI-1 is down-regulated but continues to be expressed. The vast majority of *PinvF*-positive bacteria were outside the villus (77%) at 8 h p.i., typically within sloughed-off cells in the mucosal layer immediately adjacent to the tissue (Fig. 4D). Of the tissue-associated *PinvF*-positive bacteria at 8 h p.i., there was more variation in their distances from the apical surface than at 2 h p.i. Some were within 10 μm of the apical surface, but others were found much deeper within the tissue (Fig. 3D). In concordance, tissue-associated *PinvF*-positive bacteria were similarly distributed between the epithelium and lamina propria at 8 h p.i. (Fig. 4D).

To better understand the prevalence of SPI-1 gene expression within the subtissue compartments over time, we determined the proportion of GFP-positive *S. Typhimurium* cells in the epithelium, lamina propria, and extravillous space. In the epithelium and lamina propria, we observed a marked decrease in the proportion of *PinvF*-GFP[LVA] expression from 2 h p.i. to 8 h p.i.: 68% ($n = 298$) to 39% ($n = 85$) in the epithelium and 63% ($n = 171$) to 17% ($n = 162$) in the lamina propria (Fig. 4E). A previous report has observed a similar decrease in expression of another gene within SPI-1, *sicA*, in the murine intestine after oral infection (44). In contrast, the proportion of *PinvF*-GFP[LVA]-positive bacteria remained constant from 2 h to 8 h in the extravillous space, with 58% ($n = 93$) positive at the early time point and 53% ($n = 389$) positive later in the infection (Fig. 4E). Historically, a compelling role for SPI-1 in the colonization of nonphagocytic cells, such as epithelial cells, has been well established (45–47). Much less is known about this pathogenicity island and phagocytic cells, but SPI-1 does contribute to cell death in macrophages and dendritic cells and to nitric oxide production in macrophages (48, 49). Our tissue localization data indicate that both enterocytes and cells within the lamina propria are colonized by SPI-1-induced *S. Typhimurium* during acute enteritis.

For the SPI-2 regulon, no tissue-associated *PssaG*-positive bacteria were observed at 2 h (Fig. S6B and see above). In contrast, *PssaG*-positive bacteria were prevalent at 8 h p.i. and strongly associated with subepithelial tissue (97%, $n = 308$), especially cells positive for HLA-DR α (Fig. 4C), which is a cell surface receptor found on dendritic cells, B cells, and monocytes/macrophages. These immune-originating cell types are often found near the central lacteal. This preferential localization of *PssaG*-positive *S. Typhimurium* cells to the lamina propria was also reflected in the distance of the bacteria from the apical surface (Fig. 3C and D); GFP-positive bacteria were significantly further from the apical surface than *PinvF*-positive *S. Typhimurium* cells at the same time point. Despite this strong association with the lamina propria, in some instances, *PssaG*-positive bacteria were found within cytokeratin 8-positive epithelial cells (Fig. 4B). Proportionally, 58% ($n = 518$) of bacteria in the lamina propria were *PssaG* positive at 8 h p.i., while only 37% ($n = 27$) were positive in the epithelium (Fig. 4E). Intracellular induction of SPI-2 is known to occur in

both nonphagocytic and phagocytic cells (50, 51), in agreement with the tissue localization that we report here.

In this work, we have utilized a natural-host model of infection, i.e., bovine, ligated jejunal-ileal loops, to better understand host-pathogen interactions during acute enteritis. One striking observation was that of extruding or sloughing enterocytes harboring *S. Typhimurium* during early stages of infection. Bacteria within these extruded cells expressed SPI-1 and flagella. We have previously reported a similar phenotype in polarized epithelial monolayers (27). Epithelial cell turnover is a normal process in healthy, uninfected tissue that must be exquisitely regulated; an imbalance can lead to states of acute or chronic pathological disturbances in the gut. Interestingly, gastrointestinal infections are often associated with altered rates of epithelial cell extrusion in the gut. For example, after oral inoculation of calves, enteropathogenic *Escherichia coli* (EPEC) induces enterocyte exfoliation in the terminal rectum (52). Likewise, extensive epithelial shedding into the lumen of the small intestine is observed upon *Vibrio parahaemolyticus* infection of infant rabbits (53). Perhaps increased epithelial cell shedding is a common host defense mechanism against enteric pathogens.

In order to establish a successful infection, pathogens must precisely regulate virulence gene expression, both temporally and spatially. By documenting bacterial gene expression at the single-cell level *in vivo*, we have demonstrated for the first time a distinct segregation of SPI-1- and SPI-2-induced *S. Typhimurium* cells over a time course of acute intestinal infection. *PinvF*-positive bacteria were found predominantly within enterocytes early during infection but were mostly limited to the extravillous space as the infection progressed, likely due to immunological responses (i.e., PMNs, professional phagocytes, etc.) and pathological responses (i.e., villus blunting, epithelial sloughing) by the host. Extruded epithelial cells contained SPI-1-induced, flagellated bacteria, suggestive of reseeding of the lumen with invasive bacteria (26). In contrast, SPI-2-induced bacteria were almost exclusively found within the lamina propria and only at 8 h p.i. Notably, less than half of the tissue-associated *Salmonella* cells were induced for either SPI-1 or SPI-2 at 2 h or 8 h p.i., although it is likely that at earlier time points, a larger proportion of the bacterial population is induced for SPI-1, given the central role that this pathogenicity island plays in enterocyte entry and colonization (45, 54). Variable gene expression in genetically identical populations of *Salmonella* is well recognized and described for broth culture conditions (55–57) and infection of tissue culture cells (26–28, 58) but has only lately received attention *in vivo* (59, 60). Heterogeneity is most certainly dictated by both host and bacterial factors (61), and the complex nature of the gut suggests a high likelihood of population heterogeneity during colonization. For example, movement from the anaerobic lumen of the intestine to the zone of relative oxygenation at the epithelial surface is considered a trigger for T3SS activation on the surface of *Shigella flexneri*, the causative agent of dysentery (62). Heterogeneity in virulence gene expression *in vivo* has also recently been reported for *Vibrio cholerae*, the etiological agent of cholera. During infection of rabbit ileal loops, only bacteria that are closest to the epithelial surface, not those in the luminal fluid, express *tcpA*, a structural subunit of the toxin-coregulated pilus (63). The existence of these subpopulations of bacteria *in vivo* underscores the importance of single-cell studies for analyzing gene expression. Future studies directed at deciphering the distinct microenvironments within the gut and how bac-

teria respond to each of these environments is key to our understanding of the complexities of pathogen-host interactions during gastroenteritis.

MATERIALS AND METHODS

Bacterial strains and culture. *Salmonella enterica* serotype Typhimurium derivative IR715 was transformed by electroporation with the plasmid pMPMA3ΔPlac, containing a destabilized version of green fluorescent protein (GFP[LVA]) under the control of either the *invF* or the *ssaG* promoter (26, 64). A promoterless GFP[LVA] plasmid (EMPTY-GFP) served as a vector control (26). Due to the reduced stability of GFP[LVA] (~40 min in *S. Typhimurium*), these plasmids are valid reporters for transient gene expression (26). Bacterial cultures were grown either in shaking Luria-Bertani (LB) broth or on LB agar plates containing nalidixic acid (50 mg/liter) with carbenicillin (100 mg/liter) when appropriate. Bacterial inocula for the ligated ileal loop surgeries were prepared as described previously (22). Briefly, IR715, pMPMA3ΔPlac-*PinvF*-GFP[LVA] (*PinvF*), and pMPMA3ΔPlac-*PssaG*-GFP[LVA] (*PssaG*) were grown in LB broth with appropriate antibiotics for 14 h at 37°C at 220 rpm in a shaking incubator (model 24; New Brunswick Scientific). Cultures were then diluted 1:100 in LB broth containing carbenicillin (100 mg/liter), where appropriate, and incubated as described above for 4 h. Bacteria in the exponential phase of growth were quantified using a Genesys 10S visible-light spectrophotometer (Thermo Scientific) and diluted to 10⁹ CFU in 3 ml of LB broth. Bacterial densities were confirmed by plating on LB agar plates with appropriate antibiotics.

Animals and surgeries on bovine, ligated jejunal-ileal loops. Surgeries on ligated jejunal-ileal loops were performed as described previously (22, 65). Brangus calves 4 weeks of age and 45 to 55 kg were used in accordance with the Texas A&M University International Animal Care and Use Committee (IACUC) animal use policies and approved under Animal Use Protocol 2011-077. Calves were obtained from the Texas A&M University Veterinary Medical Park and received colostrum prior to isolation. Animals were fed antibiotic-free milk replacer twice daily and water *ad libitum*. Prior to surgery, calves were twice tested for *Salmonella* spp. in fecal excretions. Rectal swabs were collected immediately after isolation and again 1 week prior to surgery. Swabs were placed in tetrathionate broth (BBL) overnight and subsequently streaked onto XLT-4 agar plates (BBL). All calves were negative for *Salmonella* species colonies on XLT-4 plates after 48 h of incubation. Loops from seven calves were utilized for this study, with at least three independent loops for each bacterial strain.

For the surgical procedure, calves were fasted for 12 h prior to surgery. Anesthesia was induced with propofol (Abbot Laboratories), followed by intubation and maintenance with isoflurane (Isoflor; Abbot Laboratories) for the duration of the experiment. After laparotomy, the distal jejunum and ileum were externalized and 20 to 30 loops, each ~6 cm in length, were formed with a 1-cm spacer loop in between. Bacterial cultures of 3 ml containing 1 × 10⁹ total CFU were prepared as described above and loaded into a 5-ml syringe with a 26-gauge needle and kept on ice until inoculation into the loop via intraluminal injection. Cultures were inoculated into separate loops as WT (*S. Typhimurium* IR715), LB (LB broth), *PinvF* (IR715 harboring pMPMA3ΔPlac-*PinvF*-GFP[LVA]), *PssaG* (IR715 pMPMA3ΔPlac-*PssaG*-GFP[LVA]), or Empty (IR715 pMPMA3ΔPlac-null-GFP[LVA]). Following inoculation, the loops were returned to the body cavity (the surgical incision was temporarily secured) and maintained at approximately 37°C. Infections were allowed to continue for 2 h or 8 h before excision and processing for bacteriology and confocal and electron microscopy.

Pathology, bacteriology, and plasmid retention. To assess the level of tissue-associated *S. Typhimurium* cells, two 6-mm biopsy punches (0.1 g) from the Peyer's patch portion of the loop (antimesenteric side of the intestinal mucosa) were collected from each loop. Extracellular bacteria were removed by washing the samples three times in sterile phosphate-buffered saline (PBS), followed by incubation for 1 h in 10 μg/ml genta-

micin in PBS. The biopsy specimens were then homogenized and diluted in PBS (10-fold) before being plated on selective LB agar plates. For samples containing WT *S. Typhimurium*, LB agar plates contained 50 mg/liter nalidixic acid (LB-NAL). *PinvF* or *PssaG* *S. Typhimurium* samples were plated on LB-NAL plates, and LB-NAL was supplemented with 100 mg/liter carbenicillin (LB-NAL/CARB). LB control samples were plated on LB-NAL plates. Plates were incubated overnight at 37°C, and colonies were then counted. Data are reported as log numbers of CFU per mg of tissue. To quantify *S. Typhimurium* cells in the mucus, ileal loops were opened and tissue surfaces were scraped to collect mucus. Scrapings were placed in a preweighed container, reweighed, serially diluted, and spread on LB-NAL or LB-NAL/CARB selective plates, and colonies were counted the next day. Data are reported as log numbers of CFU per mg of mucus. For quantification of *S. Typhimurium* cells in the luminal fluid, luminal contents were collected from intact loops and quantified for weight and volume, 10-fold serially diluted in sterile PBS, and plated on either LB-NAL or LB-NAL/CARB plates as described above; the plates were incubated overnight at 37°C and colonies counted the next day. Data are reported as log numbers of CFU per volume of fluid.

Tissue and fluid fixation for microscopy. For each loop, 6-mm biopsy specimens were taken from the Peyer's patch tissue and placed into 10% buffered formalin for confocal microscopy or 2.5% glutaraldehyde, 2.5% formaldehyde in 0.1 M sodium cacodylate buffer for transmission electron microscopy (TEM) for 24 h. Samples for confocal microscopy were then floated into 20% sucrose with 0.05% sodium azide and stored at 4°C until use. Samples for TEM were placed into 0.1 M sodium cacodylate buffer and kept at 4°C until processed further.

Fluid samples were removed from loops as described above, and 100 μl of a sample was placed in 500 μl of 10% buffered formalin overnight. Following fixation, samples were centrifuged at 10,000 × *g* for 10 min, the supernatant was removed, and the pellet was gently resuspended in 20% sucrose with 0.05% sodium azide. Samples were stored at 4°C until processed further.

Confocal microscopy. Once infused with 20% sucrose, tissue samples were enrobed with optimum-cutting-temperature compound (Sakura Finetek USA, Inc.) and snap-frozen in liquid nitrogen. Frozen samples were sectioned at 10 μm on an OTF/AS 5000 Cryostat (Vibratome). For immunostaining, sections were rehydrated with PBS for 5 min, followed by permeabilization and blocking with 2% normal donkey serum, 1% bovine serum albumin (BSA), 0.1% Triton X-100, and 0.05% Tween 20 in PBS for 45 min at room temperature (RT). Primary antibodies were diluted in 1% BSA, 0.1% Triton X-100, and 0.05% Tween 20 in PBS overnight at 4°C, followed by 3 washes for 5 min each in 0.05% Tween 20 in PBS (PBST). Secondary antibodies were diluted in 0.1% Triton X-100 in 0.05% Tween 20 in PBS and incubated for 1 h at RT. Sections were washed 3 times for 5 min each in PBST, coated in SlowFade gold antifade reagent with 4',6-diamidino-2-phenylindole (DAPI; Life Technologies), and covered with a coverslip. Samples were cured overnight at RT. Slides were viewed at appropriate fluorescent wavelengths on a Carl Zeiss LSM 510 META NLO Multiphoton confocal microscope (Texas A&M University) or an LSM 710 confocal laser scanning microscope (Rocky Mountain Laboratories, National Institutes of Health). Images were processed and rendered with ImageJ (W. S. Rasband, National Institutes of Health, Bethesda, MD) and assembled using Adobe Photoshop CS3 or Elements 9 (ACD Systems).

Antibodies and reagents. Primary antibodies for indirect immunofluorescence staining were mouse monoclonal anti-FliC (1:100; BioLegend), rabbit polyclonal anti-*Salmonella* O-antigen group B factors 1, 4, 5, and 12 (1:200; Difco), rabbit polyclonal anti-GFP (1:1,000; Life Technologies), mouse monoclonal anti-major histocompatibility complex class II DR alpha DA6.147 (HLA-DRα) (1:100; Santa Cruz), and mouse monoclonal anti-human cytokeratin 8 CK3-3E4 (1:200; Miltenyi Biotec). Fluorescent secondary antibodies used included Alexa Fluor 488-, 568-, or 647-conjugated secondary antibodies (1:1,000; Life Technologies) and

DyLight 488-conjugated secondary antibodies (1:500; KPL). Alexa Fluor 647 phalloidin was used at a 1:50 dilution (Life Technologies).

Tissue localization of SPI-1- and SPI-2-induced bacteria. Tissue sections from 2-h or 8-h infections with *PinvF* or *PssaG* strains from multiple calves were immunostained with anti-cytokeratin 8 and anti-GFP antibodies. The locations of GFP-positive bacteria were designated (i) “extravillous” if bacteria were in the lumen, (ii) “epithelium” if they were in cytokeratin-positive cells, or (iii) “lamina propria” if bacteria were in the tissue beneath the epithelium. At least 20 villi were scored for each time point and infection. Distances of tissue-associated *PinvF*- or *PssaG*-positive bacteria to the nearest apical surface were quantified with the measurement tool in ImageJ and analyzed with Prism 5 (GraphPad Software Inc.). Data were displayed in box-and-whisker plots, with the whiskers representing the 5th-to-95th percentiles. Data were analyzed statistically by a one-way analysis of variance followed by Tukey’s multiple-comparison test, with a *P* of <0.01 considered significant.

Electron microscopy. Tissue samples fixed for TEM were postfixed for 1.3 h in 1% OsO₄ reduced with 0.35% K₄[Fe(CN)₆] and buffered with 0.1 M sodium cacodylate. The samples were dehydrated in an ascending ethanol gradient and embedded in epoxy resin. Thin sections (60 to 90 nm) were prepared with a Leica EM UC6 ultramicrotome and post-stained with uranyl acetate and lead citrate. The sections were viewed and imaged with a Morgagni 268 transmission electron microscope (FEI). Images were cropped, and exposure was optimized and sharpened in Photoshop Elements 9 (Adobe).

Analysis of GFP reporter expression. Bacteria were grown under SPI-1- or SPI-2 inducing conditions as follows. The SL1344 wild type (66), *Δhila* mutant (67), or *ssrB::kan* mutant (68) harboring pMPMA3ΔPlac-*PinvF*-GFP[LVA] or pMPMA3ΔPlac-*PssaG*-GFP[LVA] was grown overnight with shaking in LB-Miller broth at 37°C and then subcultured 1:33 in LB-Miller broth for 3.5 h with shaking at 37°C to induce SPI-1. For SPI-2 induction, bacteria were grown in LB-Miller broth with shaking for 4 to 5 h at 37°C. Bacteria (0.5 ml) were centrifuged, washed once in low-phosphate, low-magnesium (LPM) medium, pH 5.8 (69), and resuspended in an equal volume of LPM medium. Bacteria were then diluted 1:50 in LPM medium and grown for 14 h with shaking at 37°C.

Cultures were centrifuged at 8,000 × *g* for 2 min, and bacterial pellets were washed once in PBS and then fixed in 1% paraformaldehyde (PFA) for 10 min at room temperature. Bacteria were washed in PBS and then stained with mouse monoclonal anti-*S.* Typhimurium group B lipopolysaccharide (LPS; 1:200 dilution, clone 1E6; Meridian Life Science), followed by Alexa Fluor 568-conjugated goat anti-mouse IgG (1:400 dilution). After being washed once in PBS, bacteria were mounted on glass slides using Mowiol 4-88 (Calbiochem) and viewed on a Leica DM4000 upright fluorescence microscope.

HeLa epithelial cells (ATCC CCL-2) and RAW264.7 macrophage-like cells (ATCC TIB-71) were obtained from the American Type Culture Collection and used within 15 passages of receipt. HeLa cells were grown in Eagle’s minimum essential medium (EMEM; Corning Cellgro) containing 10% heat-inactivated fetal calf serum (FCS; Invitrogen). RAW264.7 cells were grown in Dulbecco’s modified Eagle’s medium (DMEM; Corning Cellgro) containing 10% heat-inactivated FCS. Cells were seeded on acid-washed glass coverslips in 24-well plates at 6 × 10⁴ cells/well (HeLa) or 2 × 10⁵ cells/well (RAW264.7). SPI-1-induced bacteria were grown to late log phase as described above, centrifuged at 8,000 × *g* for 2 min, and then resuspended in Hanks’ buffered saline solution (HBSS; Corning Cellgro). Bacteria were added to epithelial and macrophage-like cells at multiplicities of infection of 50 and 10, respectively, for 10 min. Noninternalized bacteria were removed by three washes with HBSS and cells incubated in growth medium until 30 min p.i. Then, growth medium containing 50 μg/ml gentamicin was added for 1 h, followed by growth medium containing 10 μg/ml gentamicin for the remainder of the experiment. Infected monolayers were fixed in 2.5% PFA for 10 min at 37°C and then permeabilized and blocked in 10% normal goat serum–0.2% Triton X-100–PBS for 20 min. Primary anti-

bodies were rabbit anti-GFP (1:1,000 dilution; Molecular Probes) and mouse monoclonal anti-*S.* Typhimurium group B LPS (1:2,000 dilution, clone 1E6; Meridian Life Science). Secondary antibodies were Alexa Fluor 488-conjugated goat anti-rabbit IgG and Alexa Fluor 568-conjugated goat anti-mouse IgG (1:800 dilution; Molecular Probes). Coverslips were mounted on glass slides in Mowiol, and the percentages of GFP-positive bacteria were scored by fluorescence microscopy.

SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at <http://mbo.asm.org/lookup/suppl/doi:10.1128/mBio.00946-13/-/DCSupplemental>.

Figure S1, TIF file, 5.3 MB.

Figure S2, TIF file, 4.2 MB.

Figure S3, TIF file, 2.8 MB.

Figure S4, TIF file, 0.8 MB.

Figure S5, TIF file, 0.8 MB.

Figure S6, TIF file, 1.7 MB.

ACKNOWLEDGMENTS

This work was funded by grants to A.J.B. and L.G.A. from the National Institute of Allergy and Infectious Diseases (A144170 and A1076246). L.A.K. is supported by startup funds from the Paul G. Allen School of Global Animal Health. O.S.-M. is supported by the Intramural Research Program of the National Institutes of Allergy and Infectious Diseases. Confocal and electron microscopy performed at the Image Analysis Laboratory, College of Veterinary Medicine, Texas A&M University, were funded by an NIH NCRR Shared Instrumentation grant (1S10RR22532-01). R.C.L. was partially supported by a College of Veterinary Medicine, Texas A&M University, Postdoctoral Trainee Research award. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

We thank Clay Ashley and Destiny Taylor at the Veterinary Medical Park, College of Veterinary Medicine, Texas A&M University, for their assistance during surgery. We thank Robert Alaniz for helpful conversations and critical review of the manuscript.

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