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Salmonella Regulator STM0347 Mediates Flagellar Phase Variation via Hin Invertase

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Abstract: Salmonella enterica is one of the most important food-borne pathogens, whose motility and virulence are highly related to flagella. Flagella alternatively express two kinds of surface antigen flagellin, FliC and FljB, in a phenomenon known as flagellar phase variation. The molecular mechanisms by which the switching orientation of the Hin-composed DNA segment mediates the expression of the fljBA promoter have been thoroughly illustrated. However, the precise regulators that control DNA strand exchange are barely understood. In this study, we found that a putative response regulator, STM0347, contributed to the phase variation of flagellin in S. Typhimurium. With quantitative proteomics and secretome profiling, a lack of STM0347 was confirmed to induce the transformation of flagellin from FliC to FljB. Real-time PCR and in vitro incubation of SMT0347 with the hin DNA segment suggested that STM0347 disturbed Hin-catalyzed DNA reversion via hin degradation, and the overexpression of Hin was sufficient to elicit flagellin variation. Subsequently, the ∆stm0347 strain was outcompeted by its parental strain in HeLa cell invasion. Collectively, our results reveal the crucial role of STM0347 in Salmonella virulence and flagellar phase variation and highlight the complexity of the regulatory network of Hin-modulated flagellum phase variation in Salmonella.

Keywords: Salmonella enterica; STM0347; flagellar phase variation; Hin; motility; virulence

1. Introduction

As one of the most important food-borne pathogens, Salmonella enterica can cause vomiting, diarrhea, gastroenteritis, typhoid fever, systemic infection, and even death [1–3]. It is estimated that there are over 93.8 million cases of gastroenteritis caused by Salmonella infection per year worldwide, leading to 155,000 deaths from gastroenteritis and 84,800 deaths due to diarrhea every year [4,5]. In the process of infecting host cells, Salmonella first invades non-phagocytic intestinal epithelial cells to survive and proliferate intracellularly and then penetrates epithelial cells for deeper infection [6,7]. It is well documented that bacterial motility, chemotaxis, and virulence, including host cell invasion, depend on flagella to a large extent [8,9]. On the one hand, Salmonella flagella can drive bacteria to scan and colonize host cells [10,11]; on the other hand, they can also be recognized by the host innate immune system, triggering immune defenses [12]. Consequently, Salmonella has evolved specific mechanisms to evade host immunity.

Salmonella enterica serovar Typhimurium assembles a sophisticated flagellum structure. There are three basic components to the flagellum, including the basal body, hook, and filament, which alternatively expresses two different antigenic proteins, FliC (H1 antigen) and FljB (H2 antigen) [13,14]. The flagellin type exhibits a regulatory effect on the virulence of Salmonella. Salmonella locked in the FliC-expressing phase is more virulent than the
FljB-expressing mutant, whether by oral or intraperitoneal inoculation, probably due to its superiority at infecting and colonizing the spleen [15]. Additionally, in normal swimming motility observations, there are no marked differences between FliC- and FljB-locked bacteria; however, FljB-expressing bacteria have a greater swimming speed under high-viscosity conditions [16]. An in-depth study revealed a distinct motor pattern between Salmonella with two flagella phases. When flagella were locked in the FliC-expressing phase, Salmonella tended to stop frequently and swim for a prolonged period near the host cell surface, which resulted in excessive gastrointestinal colonization in murine infection models [17].

The alternate expression of these two antigen phases of the flagellum is defined as flagellar phase variation, the study of which dates back to 1922 [18,19]. The regulatory mechanisms that underlie flagellum phase variation have also been widely reported. The gene that promotes fljB in S. Typhimurium is sliced by a reversible DNA segment, which is sandwiched by two repetitive sequences, hixL and hixR [20]. When this DNA segment is in one orientation and the promoter approaches the fljBA operon, the FljB flagellum is expressed, and fliC is suppressed at the transcriptional level by the direct binding of FljA with the fliC promoter [20]. When Hin recombinase controls the flipping of the reversible segments to another orientation, neither fljB nor fljA is transcribed, and the FliC flagellum dominates in the bacteria. Previous research has illustrated the necessity of Fis and HU binding with a recombinational enhancer on the hin gene while assembling the invertasome during DNA inversion [21,22]. However, studies that explore the other regulatory factors of Hin-mediated flagellum phase variation remain insufficient.

A previous study globally compared gene expression across Escherichia coli, Salmonella enterica, and Bacillus subtilis, and a phylogenetic tree analysis revealed a conserved homologous regulator of CsgD (E. coli), STM0347 (S. enterica) [23], which is a putative response regulator whose functions remain to be characterized. In the present study, we found that FliC was downregulated in the stm0347 deletion mutant compared with the wild-type (WT) strain of S. Typhimurium via proteomic profiling. We further confirmed the flagellum phase variation phenomenon in the stm0347 deletion mutant by visualizing secreted proteins. Notably, our data indicated STM0347-dependent Hin-inversion, suggesting a linkage between this transcription factor and Salmonella flagellum phase variation. Importantly, we provide a series of lines of evidence that STM0347 might prevent invertible DNA from flipping orientation by diminishing hin mRNA levels or degrading hin DNA fragments, and the overexpression of Hin was able to induce flagellar phase variation to a certain extent. In addition, a variant lacking stm0347 was outcompeted by WT in terms of host cell invasion.

2. Results

2.1. Homology Analysis of STM0347 Protein

Salmonella STM0347 is a putative response regulator whose functions remain to be characterized. A genome comparison between E. coli and S. Typhimurium found homology between STM0347 and a LuxR-like protein, CsgD, and proposed that STM0347 might have evolved to be a regulator of pathogenesis-related protein [23]. STM0347 is relatively distinctive among bacterial species, and sequence alignment with its potential homologous protein also indicates that it exerts similarity to a transcriptional regulatory protein, UhpA, which is known to contain a helix-turn-helix (HTH) luxR-type domain (Figure 1A). UhpA belongs to the NarL family, exhibits a regulatory effect on sulfur assimilation pathways in Salmonella, and participates in sugar phosphate transport by directly binding to the uhpT promoter in E. coli [24,25]. In addition, STM0347 is highly conserved among numerous Salmonella strains according to a BLAST search against reference sequences in the NCBI database (some representative STM0347 homologues are shown in Figure 1B). Although the sequence analysis implied a C-terminal winged helix-like DNA-binding domain, the underlying regulon and its potential functions need to be further explored.
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Figure 1. Homology analysis of STM0347 protein. Homologous proteins STM0347 were blasted with the NCBI database (https://blast.ncbi.nlm.nih.gov/Blast.cgi, accessed on 16 October 2021), multiple sequences of STM0347 with its homologs from different bacteria or different Salmonella strains were aligned by the CLUSTALW program (http://www.genome.jp/tools/clustalw/), and the figures were portrayed with ENDScript Web server (https://escript.ibcp.fr/ESPript/cgi-bin/ESPript.cgi, accessed on 16 October 2021) [26].

2.2. Comparative Proteomic Profiling of Wild-Type S. Typhimurium SL1344 and Its Isogenic Strain ΔSTM0347

To elucidate the STM0347 regulon in S. Typhimurium, we performed quantitative proteomic profiling of the stm0347 deletion mutant compared with its parental WT strain. In total, 1802 proteins were identified from three biological replicates. A complete list of all identified proteins is provided in Supplementary Table S1. To globally depict differentially expressed proteins in these two strains, we developed a protein-level volcano plot complying with the criteria described in the “Materials and Methods” section (Figure 2). With the label-free quantitation (LFQ) strategy, 41 proteins in stm0347 deletion mutants were differentially regulated, including 31 downregulated and 10 upregulated proteins, compared with WT (see Supplementary Table S1). The putative outer membrane lipoprotein STM0349 showed an outstanding lead in the upregulation of all proteins tested; however, its precise function still needs to be defined. We then noticed that the H1 antigen flagellin...
(FliC) and the flagella synthesis protein (FlgN) were significantly downregulated when 
stm0347 was lacking. Interestingly, we also found the suppression of HilA, a well-known 
transcriptional regulator of the Salmonella pathogenicity island-1 (SPI-1) type III secretion 
system (T3SS) in Salmonella. These findings strongly suggest that STM0347 might play a 
regulatory role in mediating Salmonella virulence.

Figure 2. A volcano plot of Salmonella WT and Δstm0347 strains detected by LC–MS/MS analysis. 
The logarithmic values of the abundance ratios of Δstm0347 to WT was reported on the x-axis. The 
negative logarithmic p-values processed with t test was presented on the y-axis. Dotted lines denote 
two-fold (vertical) and p < 0.05 cutoff (horizontal). Data was collected from three biological replicates.

2.3. The Deprivation of STM0347 Leads to Flagellum Phase Variation in S. Typhimurium

To test the hypothesis that stm0347 deletion induced the attenuation of Salmonella 
motility and virulence, we enriched the secreted proteins in WT and the Δstm0347 mutant 
and visualized the secreted pattern between these two strains by both Coomassie blue 
staining and immunoblotting. Interestingly, we observed a marked increase in the secretion 
of an approximately 52 kDa protein, as well as a reduction in one near 48 kDa in the 
stm0347 deletion mutant (Figure 3A). We further authenticated the two bands excised from gel by 
LC–MS/MS and identified them as flagellin subunits FljB and FliC (see Supplementary 
Figure S1). Additionally, we double checked the secretion of these two flagellin subunits with immunoblotting (Figure 3B). To confirm that the specification of the phenotype is 
triggered by stm0347 deletion rather than genetic manipulation or other inscrutable factors, 
we constructed several deletion mutants, including two virulence factors (SptP and SseL) and two metabolic regulatory factors (EntC and AraA), and detected the flagella phases. The alternative expression of two flagella phases was not observed in these strains (Figure 3C). The compensation of STM0347 by exogenous expression plasmid or genome 
recombination of the stm0347 coding sequence displayed a consistent flagella phase, FliC, 
with WT Salmonella (Figure 3A), which provided further evidence of the regulatory effect 
of STM3047 on the flagella phase variation.
Figure 3. STM0347 regulates flagellin phase variation in S. Typhimurium. (A) Secreted proteins in the wild-type strain, the Δstm0347 deletion mutant strain, the Δstm0347 complemented with an STM0347-expressing plasmid strain, and the Δstm0347 chromosomally complemented with stm0347 coding sequence strain, were visualized with Coomassie blue strained gel. (B) Protein levels of FliC and FljB in secreted proteins of the WT and Δstm0347 deletion mutant strain, determined with Western blot. (C) Secreted proteins in the WT strain, the Δstm0347 deletion mutant strain, and the sptP, sseL, entC, and araA deletion mutant strain were visualized with Coomassie blue strained gel.

2.4. ΔSTM0347-Induced Flagella Phase Variation Depends on the Modulation of Invertible Promoter of fljB

We subsequently explored whether the expression levels of fljB and fliC were altered in the Δstm0347 deletion mutant at the transcriptional level. As is shown in Figure 4A, the deprivation of Δstm0347 dramatically reduced the mRNA level of fliC and accompanied a surge in the fljB level. Correspondingly, complementation with the STM0347-expressing plasmid rescued the fljC and fljB transcripts back to WT levels (Figure 4A), suggesting a potential transcriptional regulatory effect of STM0347 on fljB. Given the putative role of the response regulator with a winged helix-like DNA-binding domain (see Figure 1), we next sought to test whether STM0347 exhibits a direct transcriptional regulatory role in fljB via an electrophoretic mobility shift assay (EMSA). DNA fragments containing the fljB promoter were incubated with purified 3×FLAG-tagged STM0347 protein; however, no retardation in electrophoretic mobility was shown at varying concentrations of STM0347 (Figure 4B).

We next focused our attention on other possible mechanisms. Since the Hin-involved DNA segment inversion participates in the flip-orientation of the fljBA promoter, and ultimately controls the alternative expression of fliC and fljB [20], the proportion of the two different orientations of the invertible segment were measured with real-time PCR and PCR amplification in the WT strain, in the Δstm0347 deletion mutant, and under complement conditions. The primers were designed according to the principles of previous studies [27,28]. For real-time PCR, one primer was located outside the invertible region, and the other two primers were located within the invertible region, which made the PCR products of hin-on-orientation (FljB-expressed) and hin-off-orientation (FliC-expressed)
both nearly 500 bp. Compared with the WT strain, the hin-on to hin-off-orientation ratio was significantly elevated in the \( \Delta stm0347 \) deletion mutant, while the ratio was restored by the heterogeneous expression of the STM0347 protein on the plasmid (Figure 5A). For the PCR amplification strategy, two recombination sites, \( hixL \) and \( hixR \), were produced from two pairs of primers. In the \( \Delta stm0347 \) strain, the hin-off-orientation fragment could barely amplify, while the products of the hin-on-orientation fragment were markedly enhanced (Figure 5B). In addition, chromosomally complemented \( stm0347 \) was adequate to resume the flip sequence switching back to WT levels. Interestingly, flagella phase variation did not occur when we locked the invertible sequence to one orientation or knocked out the coding sequence of hin at a specific orientation (Figure 5C,D). Together, these findings reveal that STM0347-triggered flagella phase variation is related to the switch orientation of the Hin-containing DNA segment.

Figure 4. Transcriptional regulatory effect of fliC and fljB by STM0347. (A) Relative mRNA levels of fliC and fljB in WT, \( \Delta stm0347 \), and \( \Delta stm0347 \) complemented with an STM0347-expressing plasmid induced with 0.001% arabinose. Expression levels are calibrated with 16 s rRNA. Results are presented as mean ± SD. Asterisks indicate significant differences (**p < 0.001). (B) EMSA experiments denied the direct regulation of STM0347 on fljB promoter. Different amounts of purified STM0347 (ranging from 0 to 5 \( \mu \)g) were incubated with fljB promoter prior to electrophoretic separation. ‘Free’ indicates free DNA.
WT level (Figure 6B). On the other hand, we hypothesized that STM0347 might disturb Fis-initiated invertasome assembly via competitive binding to a Hin enhancer; thus, an STM0347-mediated flagella phase variation. (A) A real-time PCR strategy determined the ratio of fliC- and fljB-expressing DNA orientation in WT, Δstmo347 and Δstmo347 complemented with a STM0347-expressing plasmid induced with 0.001% arabinose. Results are presented as mean ± SD. Asterisks indicate significant differences (* p < 0.05). (B) Two hix sites representing the off orientation (top) or the on orientation (bottom) were PCR amplified from the genomic DNA of the WT, Δstmo347, Δstmo347 chromosomally complemented with stmo347 coding sequence strains, to identify the orientation of the fliC- and fljB-expressed DNA orientation. The sizes of PCR products were listed as follows: hixL of hin-off-orientation (550 bp), hixR of hin-off-orientation (1700 bp), hixL of hin-on-orientation (1137 bp), hixR of hin-on-orientation (1113 bp). (C) Secreted proteins in WT, the stmo347 deletion mutant strain and four phase lock strains, was visualized with Coomassie blue strained gel. (D) Secreted proteins in WT, the stmo347 deletion mutant strain and four hin deletion mutant strains, were visualized with Coomassie blue strained gel.

2.5. ΔSTMO347-Induced Flagella Phase Variation Is Probably Involved in the Degradation of Hin

It has been confirmed that Fis and HU binding to Hin is an essential step to assemble the ‘invertasome’ during DNA strand exchange [29]. Thus, we first suspected that the STM0347 might suppress the expression of fis, hin, or the two subunits of HU, hupA, and hupB. We did not find any difference in the transcriptional levels of fis, hupA, or hupB among the WT, the stmo347 deletion mutant, or Δstmo347 complemented with an STM0347-expressing plasmid (Figure 6A). Notably, stmo347 deletion led to a statistically significant upregulation of the mRNA levels of hin. Due to the low endogenous expression level of STM0347, knockout of stmo347 did not induce a dramatic change in hin. In addition, when we tried to induce the exogenous STM0347-expressing plasmid close to the endogenous expression level with 0.001% (vol/vol) arabinose, hin expression was attenuated to the WT level (Figure 6B). On the other hand, we hypothesized that STM0347 might disturb Fis-initiated invertasome assembly via competitive binding to a Hin enhancer; thus, an EMSA was conducted to test whether STM0347 directly binds to the hin DNA segment. Interestingly, we observed a decrease in the free DNA band without a corresponding shift in band binding with STM0347 proteins (Figure 6C). Combined with the qPCR results,
we suspected that sufficient amounts of STM0347 protein could degrade the hin gene cassette, and consequently restrain the DNA inversion. To explore this possibility, we further introduced excessive Hin proteins by exogenous plasmids (0.2% arabinose) into the WT strain, and we inspected whether flagella phase variation occurred. As shown in Figure 6D, overexpression of Hin could elicit a portion of flagella transformation from FliC to FljB. Taken together, these data indicated that STM0347 might inhibit DNA inversion by degrading hin DNA segments, ultimately preventing flagella phase variation to a certain extent.

Figure 6. STM0347 inhibits Hin-catalyzed DNA reversion through diminishing hin level. (A) Relative mRNA levels of fis, hupA, and hupB in WT, Δstm0347, and Δstm0347 complemented with a STM0347-expressing plasmid induced with 0.001% arabinose. Expression levels are calibrated with 16s rRNA. Results are presented as mean ± SD. (B) Relative mRNA levels of hin in WT, Δstm0347, and Δstm0347 complemented with a STM0347-expressing plasmid induced with 0.001% arabinose. Expression levels are calibrated with 16s rRNA. Results are presented as mean ± SD. Asterisks indicate significant differences (* p < 0.05). (C) In vitro incubation of STM0347 and hin DNA segment. Different amounts of purified STM0347 (ranging from 0 to 5 µg) were incubated with hin DNA segment prior to electrophoretic separation. ‘Free’ indicates undegraded DNA bands. Gels were stained with Goldview (up) or Coomassie blue (down). (D) Secreted proteins in WT, and WT complemented with a Hin-expressing plasmid induced with 0.2% arabinose, were visualized with Coomassie blue.
2.6. ΔSTM0347 Deletion Disadvantages S. Typhimurium for Cell Invasion

Based on our observation that STM0347 regulated flagellin expression at both the transcriptional and translational levels, we sought to investigate whether STM0347 exerts a regulatory role in *Salmonella* motility. Swimming and swarming motility were assessed with 0.3% and 0.7% soft-agar plates. Not unexpectedly, there were barely any differences in terms of motility between the *stm0347* deletion mutant and its parental WT strain (Figure 7A,B), probably because STM0347 mainly mediates flagellin alteration rather than suppressing overall flagellar expression. The differences in the structure and dynamics of these two flagellar filaments might impact *Salmonella* motility under specific circumstances [16]. It was shown that FljB-expressed *Salmonella* (Δ*stm0347*) had a slightly greater swimming motility under high viscosity conditions compared with FliC-expressed bacteria (WT) (Supplementary Figure S2). Given that FliC and FljB represent two different antigen flagellins [14], flagellar phase variation might be involved in immune evasion during host infection. We exploited a HeLa-cell competitive invasion assay using a 1:1 ratio mixture of WT *Salmonella* and the *stm0347* deletion mutant, and colony forming units (CFUs) were calculated 2 h post-infection. Meanwhile, to rule out the impact of growth rate differences on invasion efficiency, we monitored the growth situation of those two strains over a 10 h period, and Δ*stm0347* appeared to be indistinguishable from WT *Salmonella* (Figure 7C). The results indicated that the WT strain remarkably outcompeted the *stm0347* deletion mutant (Figure 7C). Together, these data suggest that the absence of *stm0347* does not affect *S. Typhimurium* motility; however, it is disadvantageous for host cell invasion.

![Figure 7](image-url). The absence of *stm0347* does not affect *S. Typhimurium* motility, however disadvantageous for host cell invasion. (A) Swimming motility of WT and Δ*stm0347* strains were investigated on the 0.3% agar LB plate, and the diameters of the swimming zones were monitored at 1, 3, and 8 h. Data
were presented as mean ± SD. *n = 12. (B) Swarming motility of WT and ∆stm0347 strains were detected on the 0.7% agar LB plate, and the diameters of the swarming zones were measured at 48 h. Data were presented as mean ± SD. *n = 12. (C) Growth curves of WT and ∆stm0347 deletion mutants in LB broth. Data were presented as mean ± SD. *n = 6. (D) The competitive invasion between WT and ∆stm0347 strains. HeLa cells were infected with a 1:1 mixture of streptomycin resistance WT strain and kanamycin resistance ∆stm0347 strain at a multiplicity of infection (MOI) of 10, and CFU was determined 2 h post infection. (n = 9). Asterisks indicate significant differences (*** p < 0.001).

3. Discussion
The goal of our current study was to define the functions and regulatory mechanisms of STM0347 in *Salmonella* virulence. Our proteomics data revealed remarkable inhibition levels of HilA and FliC in the ∆stm0347 deletion mutant compared with the parental WT strain. The pathogenesis of *Salmonella* mainly depends on effector proteins, which are encoded by *Salmonella* pathogenicity islands (SPIs) on its genome, and transported into the host cell cytoplasm by the T3SS [30]. HilA has been widely reported to have a direct activation effect on two promoters of SPI-1, which plays a critical role in *Salmonella* invasion during the early stage of infection [31]. In a further investigation of the ∆stm0347 deletion mutant secretion protein patter, we observed that ∆stm0347 absence induced flagellin expression in *S. Typhimurium* to alter from FliC to FljB, which relied on the invertible expression of the fljB promoter flanked by a reversible DNA segment.

There are two flagellins, FliC and FljB, expressed in *Salmonella*, and they switch at a frequency rate of 10^{-3} to 10^{-5} per cell per generation [32]. Only FljB flagellin is produced upon fljBA operon transcription, and it further inhibits fliC expression by directly binding to its promoter region; meanwhile, only FliC flagellin is present when the operon inverts to the other orientation [33]. In this study, ∆stm0347 deprivation drastically decreased the expression of FliC and increased the expression of FljB at the transcriptional and translational levels. Meanwhile, our data showed that ∆stm0347 induced flagellar phase variation via a Hin-participating DNA segment flipping reaction. Interestingly, in addition to STM0347, IacP and QseG have also been reported to regulate flagellar phase variation by Hin inversion in *Salmonella* [34,35]; however, an in-depth understanding of the underlying regulatory mechanisms requires further investigation.

Three critical steps are essential for Hin-catalyzed DNA exchange [29,36,37]. First, the Fis-bound enhancer initiates the assembly of the invertasome, which is also known as a γδ resolvase topological structure. Afterwards, the Hin dimer binds to the synapsed hix site invertase subunits and covalently joins each 5′ end of the broken strands via an ester linkage at serine 10. Ultimately, DNA strand exchange occurs under catalytic conditions. During invertasome assembly, the presence of either of two HU subunits, HupA or HupB, is required to bind with a nonspecific sequence between the enhancer and recombination site [22]. In this case, the mRNA levels of fis, hupA, and hupB were unchanged after ∆stm0347 deletion. Remarkably, the hin transcription level was significantly increased after ∆stm0347 knockout (approximately twofold) and was attenuated by complementing the exogenous STM0347 protein. The growth of BL21 was inhibited when GST-tagged STM0347 was expressed on pGEX-6p-1 plasmid, and inclusion body proteins formed when 6×His-tagged STM0347 was expressed on pET-28a plasmid (data not shown). Thus, we suspected that STM0347 could only be expressed at a relatively low level in *Salmonella*, so endogenous knockout of ∆stm0347 simply caused a limited fold change in the hin expression level. However, despite the modest regulatory levels, ∆stm0347 deletion was still adequate for the alteration of flagellin expression with a biological amplification cascade. Combined with our in vitro incubation results, flagellar phase variation was probably arrested by the degradation of Hin when STM0347 was present in WT *Salmonella*. We further discovered that overexpression can also induce the alternative expression from FliC to FljB.

Although with highly homologous sequences and identical helical parameters, the distinguished structure and dynamics of the D3 domains mediates different motility functions in FliC and FljB flagellin-expressing *Salmonella* [16]. Similar to previous studies,
no difference in motility was detected between the \textit{stm0347} deletion mutant and the WT strain \cite{17,34,35}. Since we only observed the phase variation of flagella, rather than the expression of total amount of flagella, this phenotype was not entirely unexpected. Furthermore, flagellin is a major surface antigen for many bacterial species, and both purified recominant FliC and FljB protein can comparably activate the NF-κB pathway and induce interleukin 1β (IL-1β) secretion from host cells \cite{38,39}. In this study, we found that the invasion of the \textit{stm0347} deletion mutant was significantly outcompeted by the WT stain during the competitive invasion of HeLa cells. This was most likely because the WT strain, which expressed mainly FliC flagellin, had a distinct advantage in scanning the target cell surfaces and near-surface motility behavior as an immune evasion strategy \cite{17}. HilA has been defined as a master regulator for the SPI-1 gene cluster, which plays critical roles in \textit{Salmonella} invasion and virulence \cite{40–42}. A previous study also showed that knockout of \textit{hilA} in \textit{S}. Enteritidis resulted in the repression of the cell invasion ability \cite{43}. On the other hand, in clinically isolated antibiotic resistant \textit{Salmonella} strains, \textit{invA} (100%) was the most spread gene, followed by \textit{hilA} (88.24%), \textit{stn} (58.82%), and \textit{fliC} (52.94%) \cite{44}. Consistent with these theories, our data showed that the decrease in HilA protein levels was at least partially responsible for the impairment of invasion ability in the Δ\textit{stm0347} strain.

In conclusion, the present work took advantage of high-resolution mass spectrometry to quantitatively profile the protein expression of a \textit{Salmonella} mutant lacking \textit{stm0347} compared to its parental strain, which depicted the biological functions of \textit{STM0347}. Notably, we proposed that \textit{STM0347} diminished the Hin level, which inhibited the inversion of the Hin-mediated DNA segment, and further prevented flagellin expression from alternating from FliC to FljB (Figure 8). Importantly, the loss of \textit{STM0347} might induce, or at least partially contribute to, \textit{Salmonella} virulence attenuation during host cell invasion. Our study compensates for the lack of research on the biological functions of \textit{STM0347} and highlights the complexity of \textit{STM0347}'s regulation of Hin-mediated flagellum phase variation in \textit{Salmonella}.

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure8.png}
\caption{A proposed regulatory pathway of \textit{STM0347} mediated flagellin phase variation. \textit{STM0347} disrupted the Hin-catalyzed DNA strand exchange through attenuation of hin levels, and further inhibited the expression of flagellin transforming from FliC to FljB.}
\end{figure}
4. Methods and Materials

4.1. Bacterial Strains, Mutant Construction and Molecular Cloning

The S. Typhimurium strain SL1344 was used in this study, and all other strains are detailed in Supplementary Material Table S2. The bacteria were routinely cultivated at 37 °C in Luria–Bertani (LB) plates with 2% agar and 30 µg/mL streptomycin, unless otherwise specified. The *stm0347* deletion mutant (Δ*stm0347*) was constructed by the λ red recombinase system based on the protocol of a previous report [45]. Briefly, a kanamycin resistance gene cassette with 56 bp homologous fragments of *stm0347* at each end was amplified from pKD4 plasmids and electroporated into SL1344 carrying pKD46. The resulting *stm0347* deletion mutant with kanamycin resistance was preserved for the following competitive infection assay. To further generate chromosomally compensated *stm0347*, the phase lock mutants, and *hin* deletion mutants with the λ red system, the kanamycin resistance gene cassette in Δ*stm0347* was eliminated with the pCP20 plasmid. Successfully deleted target genes or mutated DNA segments were verified by both PCR and sequencing analysis. To construct an STM0347 complementation plasmid, the *stm0347* fragment was amplified, restriction digested, and ligated into the pSB3313 plasmid with an arabinose-inducible promoter and a C-terminal 3×FLAG tag. All primers used in this study are listed in Supplementary Material Table S2.

4.2. Proteomic Sample Preparation and LC–MS/MS Detection

To explore the underlying STM0347-regulated targets, we conducted proteomic analyses of Δ*stm0347* along with the WT strain. In brief, a single colony of each strain was inoculated in LB broth with 30 µg/mL streptomycin, and then the overnight culture was passaged 1:20 into fresh LB broth and harvested when the optical density of 600 nm reached 0.9. After centrifugation at 8000 × g for 5 min and washing with ice-cold PBS 3 times, the resulting bacterial pellets were resuspended in SDS–PAGE loading buffer and denatured at 100 °C for 5 min. Protein samples wereprefractionated using 10% SDS–PAGE and separated into 6 fractions, and in-gel trypsin digestion was performed according to a previous description [46].

The resulting peptides were dissolved in HPLC-grade water prior to LC–MS/MS analyses. For label-free proteomics analyses, a nanoflow reversed-phase LC (nLC, Thermo Fisher Scientific, Waltham, MA, USA) coupled with a hybrid ion trap orbitrap mass spectrometer (LTQ Orbitrap Velos, Thermo Scientific, Waltham, MA, USA) was applied for peptide analyses. Homemade C18 analytical columns (75 µm × 150 mm) were equipped with 4 µm of 100 Å Magic C18AQ silica-based particles (Michrom BioResources Inc., Auburn, CA, USA). A 47-min gradient was employed at the start with 5% solvent B (100% ACN, 0.1% FA), which gradually rose to 45% within 40 min. Then, the concentration of solvent B ascended to 95% in 5 min, which was followed by elution with 95% solvent B for the last 2 min. Eluted peptides were electrospayed directly into the mass spectrometer for MS and MS/MS analyses in data-dependent acquisition mode. The full MS scan was (m/z 350–1200) exploited, and the 10 most intense ions were chosen for subsequent MS/MS analyses.

4.3. Proteomic Data Processing

The raw files were processed with MaxQuant (http://maxquant.org/, version 1.5.3.8, accessed on 18 November 2021) based on the combination 5. Typhimurium protein database of LT2 (Taxon identifier: 99287) and SL1344 (Taxon identifier: 216597) downloaded from the UniProt website (www.uniprot.org, accessed on 25 June 2020) [47]. The precursor mass tolerance and the fragment mass tolerance were set at 20 ppm and 0.8 Da, respectively. Meanwhile, methionine oxidation was set as a variable modification. Trypsin was set as a digestion enzyme with a maximum of two missed cleavages. The false discovery rates (FDRs) of peptides and proteins were restricted to under 1%. We then filtered the proteins that were only identified by site, matched the reverse database, and contained potential contaminants with Perseus, and we also replaced the missing values with random
Two-group comparisons were performed with paired Student’s t-tests. Proteins with $p < 0.05$ and average fold change $> 2.0$ were considered significantly regulated.

### 4.4. Western Blot for Secreted Proteins

The supernatants of WT and $\Delta stm0347$ were collected after centrifugation at $7000 \times g$ for 20 min and filtered with a 0.22 µm membrane to further remove the remaining bacteria. Sodium deoxycholate was added to a final concentration of 0.2% (wt/vol) before the secreted proteins were precipitated with 20% (wt/vol) trichloroacetic acid (TCA) overnight at 4 °C. Finally, the protein pellets were collected, washed with ice-cold acetone 3 times, and denatured in SDS–PAGE loading buffer with 100 mM Tris-HCl (pH 8.0) at 95 °C for 10 min. Secreted protein samples were loaded for SDS–PAGE electrophoresis and were either transferred onto polyvinylidene difluoride membranes (Millipore, Billerica, MA, USA) or stained with Coomassie blue. For immunoblotting, membranes were incubated with antibodies and visualized with ECL Western Blotting Substrate (Tanon, Shanghai, China). Antibodies and dilutions were as follows: FLAG (Zen-bio, Chengdu, China, #T201126-3A6, 1:5000) and FliC (Abcam, Cambridge, UK, #ab93713, 1:5000).

### 4.5. RNA Extraction and Quantitative Real-Time PCR

Bacterial cells, including WT, $stm0347$ deletion mutant, and $\Delta stm0347$ complemented with an STM0347-expressing plasmid, were collected at OD$_{600}$ = 0.9 and ground in liquid nitrogen. Total RNA was extracted with TRIzol™ Reagent (Invitrogen™, Grand Island, NY, USA) and reverse transcribed to cDNA using the PrimeScript™ RT–PCR Kit (Takara, Dalian, China). Real-time PCR was performed on a 7500 Fast Real-Time PCR system (Applied Biosystems™, Foster City, CA, USA), which was collocated with a SYBR Premix EX Taq II Kit (Takara, Dalian, China) to quantify $fljB$, $fliC$, $fis$, $hupA$, $hupB$, and $hin$ at the transcriptional level. We used 16S ribosomal RNA as a housekeeping gene. The primers used for real-time PCR are listed in Supplementary Material Table S2.

### 4.6. DNA Extraction and Hin-Mediated Invertible Segment Orientation Determination

To explore the induction of flagellar phase variation, we picked 4 primers based on a previous depiction [28] to determine the orientation of the Hin-mediated invertible segment and the $fljBA$ promoter region. WT, $\Delta stm0347$, and $\Delta stm0347::stm0347$ bacteria were harvested at OD$_{600}$ = 0.9, and total DNA was extracted using an Ezup Column Bacteria Genomic DNA Purification Kit (Sangon, Shanghai, China) according to the manufacturer’s instructions. DNA concentrations were measured with a NanoDrop™ spectrophotometer (Thermo Scientific, Waltham, MA, USA). Equal amounts of DNA were loaded for PCR amplification, and the primers and PCR product sizes were as follows: $hixL$ of hin-off-orientation ($hix-1/hix-2$, 550 bp), $hixR$ of hin-off-orientation ($hix-3/hix-4$, 1700 bp), $hixL$ of hin-on-orientation ($hix-1/hix-3$, 1137 bp), and $hixR$ of hin-on-orientation ($hix-2/hix-4$, 1113 bp). To quantify the proportion of the on or off orientation of the invertible segment in each bacterial strain, real-time PCR was performed as described above.

### 4.7. Recombinant Protein Expression and Purification

FLAG-tagged STM0347 was expressed in the S. Typhimurium WT + pSTM0347 strain. Briefly, 400 mL bacterial suspensions were cultured to OD$_{600}$ = 0.6, and then 0.4% arabinose was added and induced for 3 h. Bacterial cells were lysed in 30 mL of ice-cold PBS buffer via sonication, and cell lysates were clarified by centrifugation at 7000 × g for 20 min at 4 °C and filtered through a 0.45 µm membrane. The resulting supernatants were incubated with anti-FLAG M2 agarose beads (50% slurry, Sigma-Aldrich, St. Louis, MO, USA) for 4 h at 4 °C to capture FLAG-tagged STM0347. Afterwards, the beads were washed 3 times with a washing buffer containing 25 mM Tris-HCl (pH 7.5) and 150 mM NaCl. FLAG peptides were used to elute the STM0347 protein and were removed by ultrafiltration.
4.8. Electrophoretic Mobility Shift Assays (EMSAs)

DNA fragments of the putative promoter region of the \( \textit{fljBA} \) operon and the coding sequence of \( \textit{hin} \) were amplified by PCR, purified with a Gel Extraction Kit (Axygen\textsuperscript{®}, Corning, CA, USA), and dissolved in water. Then, purified STM0347 proteins were incubated with 30 ng DNA fragments in 20 \( \mu \)L of binding buffer (10 mm Tris-HCl (pH 8.3), 50 mM KCl, 1 mM MgCl\(_2\), 0.5 mM DTT, 0.5 mM EDTA, 50 \( \mu \)g/mL BSA and 0.2% vol/vol glycerol). The contents of STM0347 were set at 0, 0.5, 2, and 5 \( \mu \)g. The reaction mixtures were incubated at room temperature for 20 min and then loaded onto 6% native polyacrylamide gels for electrophoresis in 1 \( \times \) TAE buffer (40 mM Tris, 20 mM acetic acid and 1 mM EDTA) in an ice bath. The gel was then stained with Goldview Nucleic Acid Gel Stain (Yeasen, Shanghai, China) or Coomassie blue and photographed using the ChemiDoc\textsuperscript{TM} imaging system (Bio-Rad, Woodinville, WA, USA).

4.9. Competitive Invasion Assay

The HeLa cells were cultured in Dulbecco’s modified Eagle medium (DMEM, HyClone, Logan, UT, USA) supplemented with 10% fetal bovine (TransGen, Beijing, China) at 37 °C under 5% \( \text{CO}_2 \) conditions. Cells were ready for the invasion assay when the density approximated 80–90% confluence. The overnight cultured WT \( \textit{Salmonella} \) and \( \textit{stm0347} \) deletion mutant with kanamycin resistance were diluted 1:20 until the OD\(_{600}\) reached 0.9, and were then mixed at a 1:1 ratio. Cells were pre-equilibrated with Hanks’ buffered salt solution (HBSS) and infected with a mixed bacterial suspension at a multiplicity of infection (MOI) of 10 for 1 h. Then, the cells were washed 3 times with HBSS and treated with 100 \( \mu \)g/mL gentamicin to kill extracellular bacteria. Finally, the cells were lysed and plated on streptomycin- or kanamycin-supplemented LB plates to calculate colony forming units (CFUs).

4.10. Growth and Motility Study

To obtain the growth curve of WT \( \textit{Salmonella} \) and the \( \textit{stm0347} \) deletion mutant, overnight cultured bacteria were diluted 1:50 in LB broth, and the optical density was monitored at 600 nm hourly over a 10 h period.

Swimming motility was detected as previously described [49] with slight modification. Briefly, bacteria were diluted 1:20 when the OD\(_{600}\) reached 0.9, and 2 \( \mu \)L was pipetted onto the middle of an LB plate with 0.3% agar. The plates were incubated at 37 °C, and the swimming zone was measured at 1, 3, and 8 h. To test the swimming motility under various viscosities, Ficoll PM400 (Sigma-Aldrich, St. Louis, MO, USA) was added to the swimming plates at a final concentration of 5% or 10% according to a previous description [16]. Swarming motility was analyzed on LB plates with 0.7% agar [50], which were cultured in closed containers with 100% humidity for 48 h.

4.11. Statistical Analysis

Data were collected from at least three independent tests, and the results are expressed as the mean ± SD. Statistical analyses were processed with SPSS 19.0 software (IBM, Armonk, NY, USA). Two-group comparisons were implemented with Student’s \( t \) test, while multiple-group comparisons were conducted with one-way ANOVA, which was followed by LSD \( t \) tests for between-group comparisons. Values of \( p < 0.05 \) were defined as statistically significant.

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