

# SCIENTIFIC REPORTS

OPEN

## Hyperinvasiveness of *Salmonella enterica* serovar Choleraesuis linked to hyperexpression of type III secretion systems *in vitro*

Received: 01 June 2016  
Accepted: 01 November 2016  
Published: 25 November 2016

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*Salmonella enterica* serovars Choleraesuis and Typhimurium are among the non-typhoid *Salmonella* serovars that are important zoonotic pathogens. In clinical observation, *S. Typhimurium* typically causes diarrheal diseases; however, *S. Choleraesuis* shows high predilection to cause bacteremia. The mechanism why *S. Choleraesuis* is more invasive to humans remains unknown. In this study, we compared the *S. Typhimurium* LT2 and *S. Choleraesuis* SC-B67 proteomes through stable isotope labeling of amino acid in cell culture (SILAC). In SILAC, the expression of many virulence proteins in two type III secretion systems (T3SSs) were significantly higher in *S. Choleraesuis* than in *S. Typhimurium*. Similar differences were also found at the transcriptional level. Compared to *S. Typhimurium*, *S. Choleraesuis* showed a higher penetration level to Caco-2 (>100-fold) and MDCK (>10-fold) monolayers. In mice after oral challenge, the invasion of spleen and liver was also higher in *S. Choleraesuis* than in *S. Typhimurium*. The transcription of *hilD* in *S. Choleraesuis* was increased in physiological (1 mM) or high (10 mM) concentrations of Mg<sup>2+</sup>, but not in low (8 μM) concentration. We conclude that *S. Choleraesuis* showed hyperinvasiveness in cellular as well as mouse models due to hyperexpression of T3SS genes.

*Salmonella* is one of the most important pathogens to humans. *Salmonella* can be divided into typhoid and non-typhoid *Salmonella* (NTS) serovars according to the respective clinical syndromes. Both *S. Typhimurium* and *S. Choleraesuis* are NTS serovars. *S. Typhimurium* typically causes diarrheal disease in young children, while *S. Choleraesuis* frequently causes diseases in all ages<sup>1,2</sup>. Previous studies have demonstrated the high invasion ability of *S. Choleraesuis* through various *ex vivo* and *in vitro* assays<sup>3,4</sup>. However, the mechanism underlying such high pathogenicity in *S. Choleraesuis* remains unknown.

The most important virulence mechanism in the *Salmonella* is the type III secretion system (T3SS). *Salmonella* contains two sets of T3SS, the genes of which are located in *Salmonella* pathogenicity island-1 and -2 (SPI-1 and SPI-2)<sup>5-7</sup>. SPI-1 T3SS participates in the early stage of infection in *Salmonella* adhesion and invasion<sup>8</sup>, while SPI-2 T3SS is involved in replication of *Salmonella* in both the phagocytic and non-phagocytic cells<sup>9</sup>. SPI-2 T3SS also plays an important role in the maturation of *Salmonella*-containing vacuoles (SCVs)<sup>10</sup>. The regulation of SPI-1 and SPI-2 T3SS is controlled by different mechanisms, in which HilA and HilD are two important proteins that can activate SPI-1 T3SS<sup>11-14</sup>. Environmental signals, like Mg<sup>2+</sup>, also regulate the SPI-1 and SPI-2 T3SS expression through two-component signal transduction systems (TCSs)<sup>15,16</sup>.

To elucidate why *S. Choleraesuis* is more invasive than *S. Typhimurium*, we compared protein expression pattern between the two serovars through stable isotope labeling by amino acids in cell culture (SILAC) analysis. We found that virulence factors linked to invasiveness were highly expressed in *S. Choleraesuis*. *S. Choleraesuis* appeared more pathogenic in cellular and murine models. Under normal and high concentrations of Mg<sup>2+</sup>, the expression of SPI-1 was even higher in *S. Choleraesuis* than in *S. Typhimurium*. It appears that the presence of

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Mg<sup>2+</sup> is associated with the higher expression of SPI-1 virulence factors in *S. Choleraesuis*, thereby promoting the subsequent invasion and penetration of *S. Choleraesuis* through host cells.

## Results

### Comparison of the protein expression patterns between *S. Choleraesuis* and *S. Typhimurium*.

In the differential proteomic study, 1731 proteins were identified in both *S. Typhimurium* LT2 and *S. Choleraesuis* SC-B67. 287 (16.6%) proteins were expressed two times higher in *S. Choleraesuis* SC-B67 than in *S. Typhimurium* LT2; on the other hand, 183 (10.6%) were expressed two times higher in *S. Typhimurium* LT2 than in *S. Choleraesuis* SC-B67 in RPMI cell culture medium (Supplementary Tables S3 and S4). Expression level of major virulence factors involved in *Salmonella* pathogenesis was higher in *S. Choleraesuis* SC-B67 than in *S. Typhimurium* LT2 (35/287, 12.20%; Table 1). The 35 virulence genes were mainly located in the virulence plasmid, SPI-1, SPI-2, SPI-4, SPI-5, and SPI-11 (Table 1). The transcriptional level of *hilA* and *hilD* was 24.0 and 21.1 fold, respectively, higher in *S. Choleraesuis* than in *S. Typhimurium*. Similar results were obtained in both the media RPMI 1640 and DMEM (Supplementary Figure S1). The transcription level of *sipB*, an effector protein gene for SPI-1 T3SS, was also higher in *S. Choleraesuis*, this correlates to the protein expression level (Supplementary Figure 1). In addition to strains used in SILAC, clinical isolates of *S. Choleraesuis* also expressed higher level of *hilD* (Supplementary Figure 2).

**Acid tolerance.** At 10 minutes post inoculation in simulated gastric fluid (SGF, pH 2.0), the number of alive bacteria was 100-fold higher in *S. Choleraesuis* than in *S. Typhimurium* (Supplementary Figure 3). At 10–30 minutes of incubation, *S. Choleraesuis* SC-B67 still outnumbered *S. Typhimurium* SL1344 by approximately 10-fold, although there was no significant difference after 15 minutes. (Supplementary Figure 3), suggesting that *S. Choleraesuis* showed better acid tolerance than *S. Typhimurium*.

**HeLa (non-polar) and MDCK (polar) cell invasion.** To compare the invasion ability between *S. Typhimurium* and *S. Choleraesuis*, we infected the epithelial cells with the two serovars separately. *S. Choleraesuis* and *S. Typhimurium* showed no difference in the level of invasiveness to polar and non-polar cells (Supplementary Figure 4).

**Penetration to epithelial cells.** *S. Choleraesuis* penetrated the MDCK (10-fold) and Caco-2 cell (1000-fold) monolayers at higher efficiency than *S. Typhimurium* (Fig. 1) at 3 hours post-infection. Even at 6 hours after infection, the number of bacteria penetrating the monolayer was still higher in *S. Choleraesuis* than in *S. Typhimurium*, especially in Caco-2 cells (Fig. 1). *S. Choleraesuis* clinical isolates also showed higher penetration ability than *S. Typhimurium* clinical isolates (Supplementary Figure 5).

**Intra-macrophage survival.** We compared the intra-macrophage survival between *S. Typhimurium* and *S. Choleraesuis* by using THP-1 cells. The result showed that *S. Choleraesuis* displayed at least 4-times higher survival than *S. Typhimurium* inside macrophages (Fig. 2). *S. Choleraesuis* clinical isolates also showed better intra-macrophage survival ability at 4 hours and 6 hours post-infection than *S. Typhimurium* clinical isolates (Supplementary Figure 6).

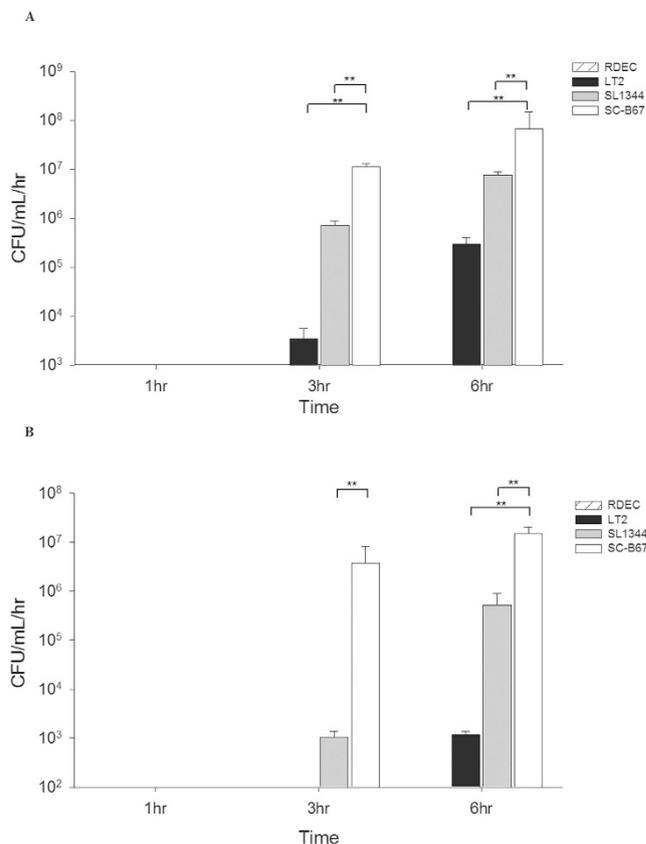
**Mouse experiments.** *S. Choleraesuis* showed higher invasion and intracellular survival compared to *S. Typhimurium* in cellular experiments. We then compared the pathogenesis of the two serovars using a murine model. One day post infection, *S. Choleraesuis* was found in spleen of one mouse (1/10) (Fig. 3A). No bacteria (less than 10 CFU/g) was found in spleen and liver of other mice (Fig. 3A and B). At 3 days post-infection, bacteria was recovered from spleen and liver at least 10 CFU/g in seven mice infected by *S. Choleraesuis*, but only in three mice infected by *S. Typhimurium* (Fig. 3A and B). Both serovars were recovered from the spleen and liver at 5 days post-infection. However, the bacterial numbers recovered were approximately 1000-times higher in *S. Choleraesuis* than in *S. Typhimurium* at 5 days post-infection (Fig. 3A and B). To further confirm the difference, we co-infected the mice with *S. Typhimurium* and *S. Choleraesuis*. After 5 days, *S. Typhimurium* was significantly outnumbered by *S. Choleraesuis* in liver and spleen (Fig. 3C).

**Expression of *hilA* and *hilD* in different *hilE* complemented strains.** *HilE* is a negative regulator protein for SPI-1 T3SS virulence factors<sup>17</sup>. The predicted translation start site of *hilE* in *S. Choleraesuis* and *S. Typhimurium* is different due to the presence of an additional nucleotide between the two start codons in *S. Choleraesuis* (Supplementary Figure 7). *HilE* becomes a pseudogene in *S. Choleraesuis* if it uses the first start codon. The same difference was also found in the clinical *S. Choleraesuis* isolates (data not shown). We tried to construct the *hilE* deficient strain in both *S. Typhimurium* and *S. Choleraesuis*. However, all the *S. Choleraesuis* are multiple drug resistant strains, we do not have suitable selection markers to construct mutant strains. To analyze the function of *HilE* in *S. Choleraesuis*, we transformed a plasmid harboring *hilE* sequence from *S. Choleraesuis* into *hilE*-deficient strain in SL1344. The expression of *hilA* and *hilD* was very low in the deletion mutant carrying the *hilE* gene from either *S. Typhimurium* or *S. Choleraesuis* (Fig. 4). In the intra-macrophage survival assay, both *hilE* complemented strains showed poor survival (Fig. 5). The results indicated that the *hilE* gene from either *S. Typhimurium* or *S. Choleraesuis* was functional and thus can complement the *hilE* deletion to suppress the expression of the SPI-1 T3SS.

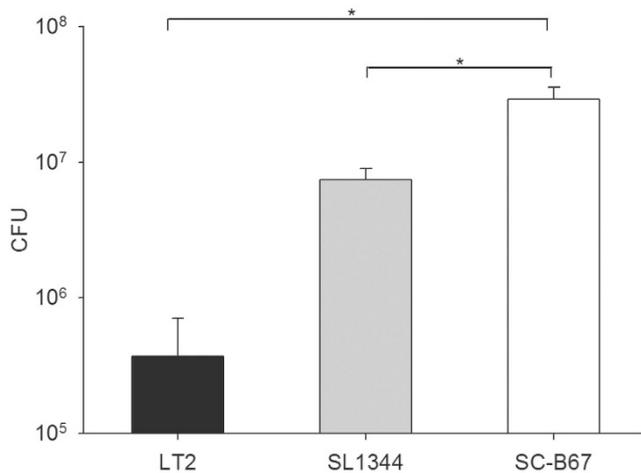
**Hypersensitivity to Mg<sup>2+</sup> in *S. Choleraesuis*.** Environmental signals, like oxygen, Mg<sup>2+</sup> and Fe<sup>3+</sup>, also control the SPI-1 T3SS expression<sup>11,15,18</sup>. We checked whether or not the environmental signals were the cause for differential gene expression between *S. Typhimurium* and *S. Choleraesuis*. Under both high and low oxygen conditions, the level of *hilD* were higher expressed in *S. Choleraesuis* SC-B67, relative to *S. Typhimurium*

Virulence locus	ID	Gene	Ratio <sup>a</sup>	
SPI-1	SC2794	<i>sitA</i>	0.99	
	SC2802	<i>orgA</i>	4.66	
	SC2806	<i>PrgH</i>	12.38	
	SC2807	<i>hilD</i>	24.04	
	SC2808	<i>hilA</i>	21.60	
	SC2814	<i>sipA</i>	22.47	
	SC2815	<i>sipD</i>	5.54	
	SC2816	<i>sipC</i>	9.65	
	SC2817	<i>sipB</i>	13.04	
	SC2824	<i>invJ</i>	6.38	
SPI-2	SC2828	<i>invA</i>	17.86	
	SC1412	<i>ssrB</i>	9.77	
	SC1415	<i>ssaC</i>	24.04	
SPI-4	SC1421	<i>sseC</i>	5.75	
	SC4140	<i>ssiE</i>	22.88	
SPI-5	SC4141	<i>ssiF</i>	5.04	
	SC1040	<i>pipB</i>	0.74	
SPI-11	SC1043	<i>sopB</i>	18.73	
SPI-13	SC1256	<i>pagC</i>	3.63	
CS54	SC3061	<i>stmR</i>	1.45	
	SC2510	<i>xseA</i>	0.84	
Gifsy-1	SC2513	<i>sinI</i>	2.26	
	SC2579	<i>gogB</i>	0.72	
Gifsy-2	SC0997	<i>sodC</i>	1.45	
	SC1459	<i>sodC</i>	4.69	
pSLT	SCH_v26	<i>pefD</i>	0.57	
	SCH-V05	<i>spvC</i>	129.87	
	SCH_V04	<i>spvB</i>	45.25	
	SCH_V48	<i>mig-5</i>	0.81	
Regulators	SC1462	<i>slyA</i>	2.63	
	SC3434	<i>ompR</i>	2.13	
	SC4120	<i>zur</i>	1.76	
	SC2896	<i>relA</i>	1.89	
	SC1181	<i>phoQ</i>	1.26	
	SC1182	<i>phoP</i>	1.17	
	SC0713	<i>fur</i>	1.12	
	SC1746	<i>hns</i>	1.26	
	SC1748	<i>hnr</i>	1.51	
	SC2645	<i>rpoE</i>	1.18	
	SC2856	<i>rpoS</i>	1.85	
	SC4014	<i>oxyR</i>	0.71	
	SC3453	<i>rtcR</i>	1.27	
	SC1950	<i>sirA</i>	1.01	
	Other effectors	SC2712	<i>virK</i>	5.60
		SC0798	<i>slrP</i>	7.42
		SC1852	<i>sopE2</i>	8.32
SC0926		<i>sopD2</i>	15.38	
SC1174		<i>sifA</i>	4.66	
SC1626		<i>sseJ</i>	22.08	
SC1252		<i>msgA</i>	2.91	
SC2710		<i>pipB2</i>	6.75	
SC2713		<i>mig</i>	3.05	
SC4040		<i>sseK1</i>	6.76	
SC3130		<i>tolC</i>	1.10	
SC2708		<i>iroN</i>	0.41	
SC2290		<i>sseL</i>	6.38	
SC1691		<i>steC</i>	18.15	

**Table 1.** The expression level of virulence factors between *S. Choleraesuis* SC-B67 and *S. Typhimurium* LT2. <sup>a</sup>SC-B67/LT2.

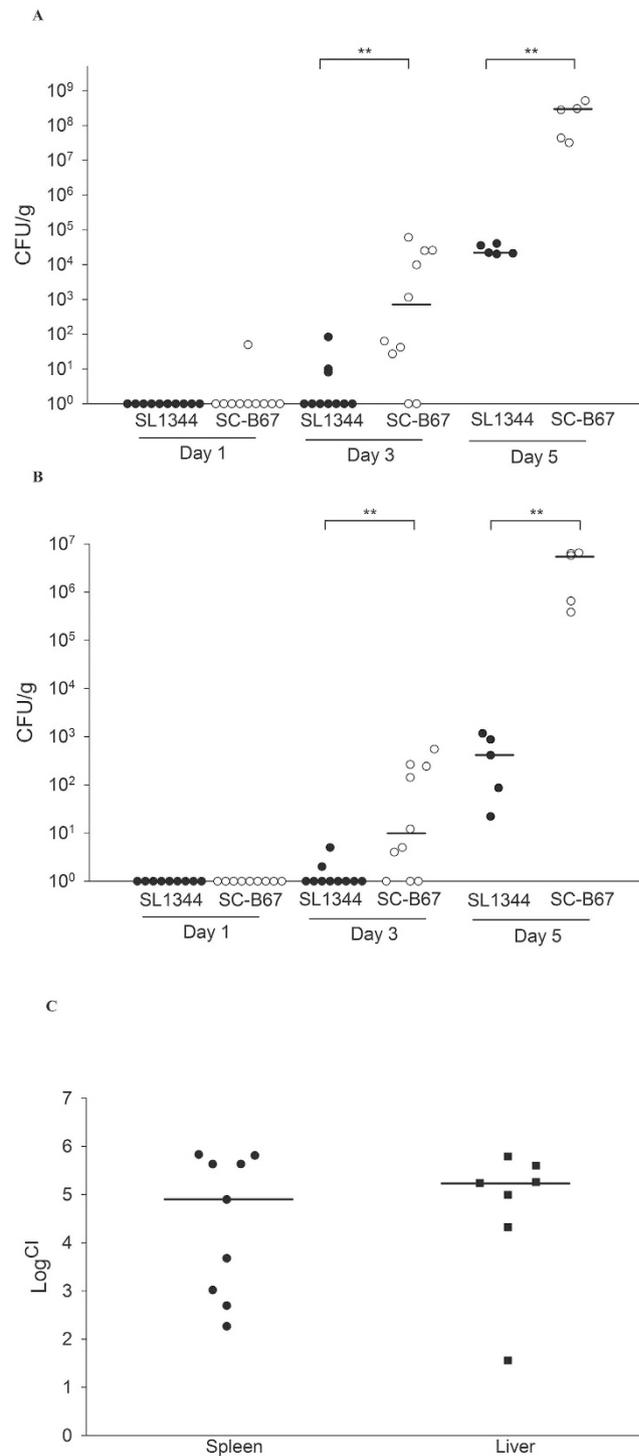


**Figure 1. Penetration assay of *S. Typhimurium* SL1344 and *S. Choleraesuis* SC-B67 through cellular monolayer.** The cells used were MDCK (A) and Caco-2 (B) cells. At appropriate time points, the number of *Salmonella* was calculated by plating on LB agar. The experiment was repeated 3 times. \*\* $p < 0.01$ .



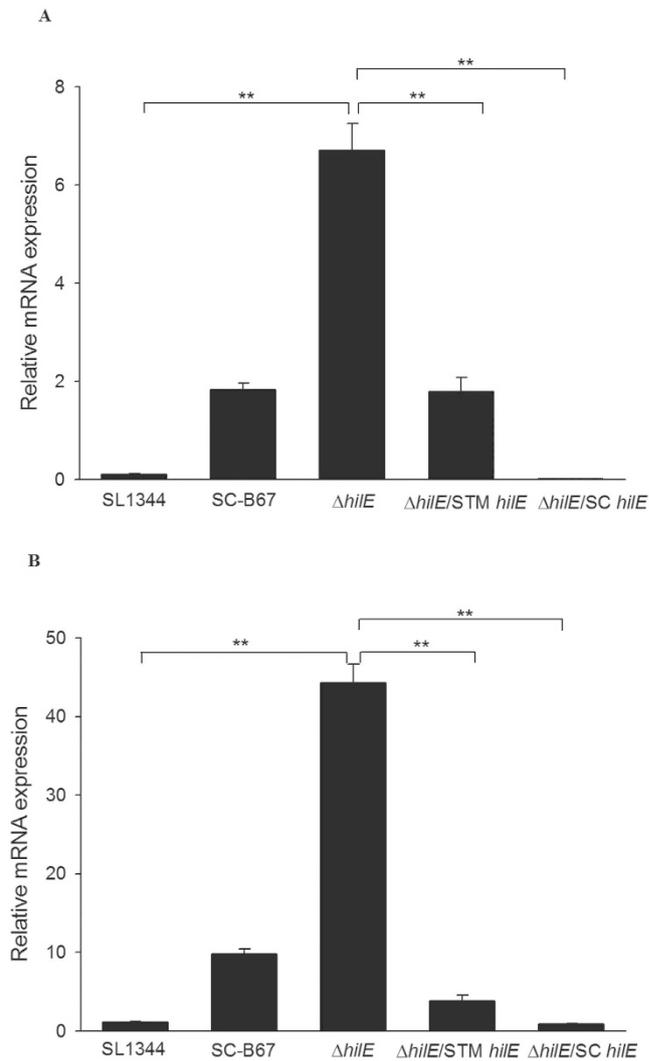
**Figure 2. Intra-macrophage survival assay.** THP-1 cells were infected by *S. Choleraesuis* SC-B67 and *S. Typhimurium* LT2 and SL1344. After 1 hour post-infection, cells were treated with gentamicin for 30 minutes. After 4 hours post-infection, cells were lysed and plated on LB agar. The experiment was repeated 3 times. \* $p < 0.05$ .

(Supplementary Figure 8).  $Mg^{2+}$  is a ligand bound to PhoPQ TCS. Under low  $Mg^{2+}$  condition, PhoPQ TCS is activated to suppress *hilA* and *hilD* expression; it is contrary under high  $Mg^{2+}$  condition<sup>15</sup>. In low ( $8 \mu M$ )  $Mg^{2+}$  condition, the expression of *hilD* was low in both serovars (Fig. 6A). Interestingly, *hilD* expression was increased in *S. Choleraesuis* SC-B67 under normal (1 mM) and high (10 mM)  $Mg^{2+}$  condition (Fig. 6A). PhoPQ TCS in *S. Choleraesuis* SC-B67 appeared hypersensitive to environmental  $Mg^{2+}$ , leading to higher expression of *HilD*.

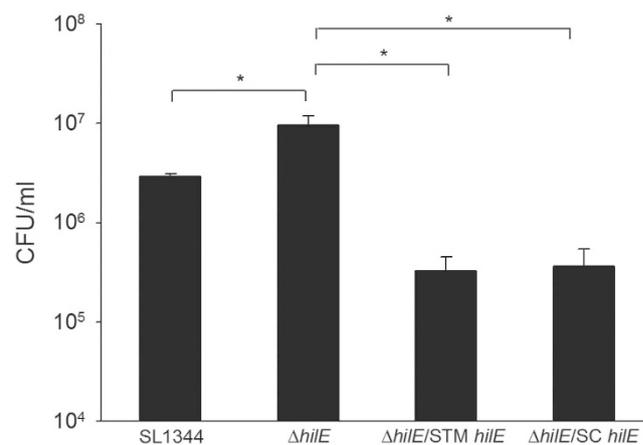


**Figure 3. Isolation of *Salmonella* from the organs.** BALB/c mice were infected by different *Salmonella* serovars through oral infection. At 1, 3, 5, and 6 day(s) post-infection, mice were sacrificed. Spleen (A) and liver (B) were grounded and plated on the SS agar plates. Mice were co-infected with SL1344 and SC-B67 (C). At 5 days post infection, both serovars were recovered from the spleen and liver, and analyzed by CI as described in the text. \*\* $p < 0.01$ .

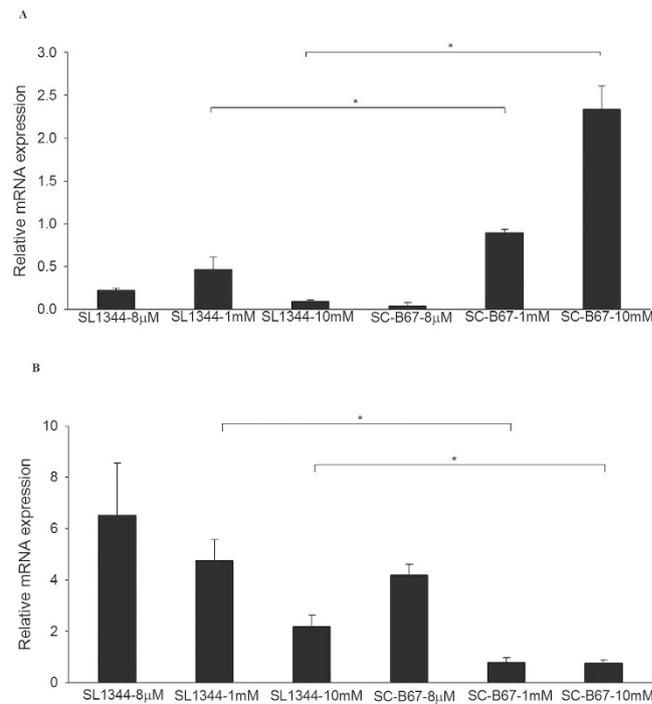
To support this, we tested expression patterns of the PhoP-activated genes (Pags) under different  $Mg^{2+}$  concentrations in the two serovars. Under low  $Mg^{2+}$  condition, the expression of *pagC* was similar between both serovars, but it was 5.4-fold lower in the *S. Choleraesuis* than in *S. Typhimurium* under normal  $Mg^{2+}$  condition, suggesting *S. Choleraesuis* was hypersensitive to  $Mg^{2+}$  to cause HilD activation (Fig. 6B).



**Figure 4. Expression of *hilA* and *hilD* in different *Salmonella* strains.** All the bacterial strains were cultured in the DMEM for 6 hours. Total RNA were isolated and analyzed by qPCR. The expression level of *hilA* (A) and *hilD* (B) are shown relative to the 16S rDNA. \*\* $p < 0.01$ .



**Figure 5. Intra-macrophage survival of *S. Typhimurium* SL1344 and its derivative strains.** THP-1 cells were infected by SL1344, *HilE* mutant strains, and its complemented strains. Bacteria were recovered at 4 hours post-infection. \* $p < 0.05$ .



**Figure 6. Gene expression of SL1344 and SC-B67 under different Mg<sup>2+</sup> concentrations.** Both serovars were cultured in N-Salt medium with 8 μM, 1 mM, and 10 mM Mg<sup>2+</sup> concentrations. Then the total RNA were isolated and analyzed by qPCR. The expression level of *hilD* (A), and *pagC* (B) in both serovars was examined in different Mg<sup>2+</sup> concentrations. All the expression levels are relative to the 16S rDNA. \**p* < 0.05.

## Discussion

Systemic approaches, like genomic, transcriptomic, and proteomic methods, have been used to compare the difference among different serovars of *S. enterica*<sup>19–21</sup>. In genomic studies, *S. Typhimurium* and *S. Choleraesuis* all contained well-defined virulence genes in SPI-1 and SPI-2<sup>19</sup>. It is hard to use sequence comparison to explain why *S. Choleraesuis* caused more bacteremia than other serovars in humans. Previous studies compared the protein expression pattern between *S. Typhimurium* and *S. Choleraesuis* through 2-dimensional SDS-PAGE analysis<sup>21</sup>, indicating that only one enzyme, GabD, showed different expression level between the two serovars<sup>21</sup>. GabD is succinate semialdehyde dehydrogenase I and the expression level of GabD appeared not directly related to invasiveness of *S. Choleraesuis*. In this study we used proteomic approach, SILAC, to analyze the differential protein expression between *S. Typhimurium* and *S. Choleraesuis*. The two serovars shared 4021 proteins in common. A total of 1731 proteins were indentified, and more than 70% of the identified proteins did not express at different level between the two. Many flagella synthesis and chemotaxis response proteins were expressed higher in *S. Typhimurium*, suggesting that motility of *S. Typhimurium* may be better than *S. Choleraesuis*, as previously described<sup>3</sup>. Some proteins from predicted pseudogenes in *S. Choleraesuis* were also detected in our analysis. This may be because *Salmonella* pseudogenes could undertake a recoding mechanism<sup>22</sup>. Most of the genes expressed higher in *S. Choleraesuis* SC-B67 are metabolism-related genes. Interestingly many virulence factors, like those in SPI-1 and SPI-2, were higher expressed in *S. Choleraesuis*. The result might explain why *S. Choleraesuis* is more invasive than *S. Typhimurium*.

*Salmonella* encounter different acid stress environment from gastric acid (pH 2.0~3.0) to phagosome (pH 5.5~6.0) during its infection route<sup>23</sup>. Acid tolerance response is important for their virulence<sup>24</sup>. *S. Choleraesuis* was more tolerant to the SGF than *S. Typhimurium*. When *S. Choleraesuis* achieves the intestine lumen, they have to pass through the intestinal mucosal barrier before they establish systemic infection. In terms of the non-polar or polar epithelial cell invasion, *S. Typhimurium* and *S. Choleraesuis* were almost the same. However, all the *S. Choleraesuis* strains showed better penetration than the *S. Typhimurium*. Epithelial cell invasion indicates whether bacteria can enter the cells by their own ability. Monolayer cell penetration is more complicated than epithelial cell invasion. Besides entry into epithelial cells, bacteria have to have transcytose or disrupt tight junction to pass through the cellular barrier<sup>25,26</sup>. SPI-1- and SPI-4-encoded genes have been proved to play a vital role in invasion of epithelial cells<sup>27–29</sup>. These effectors also can disrupt the structure and function of tight junction, thereby helping *Salmonella* to penetrate intestine for further dissemination<sup>29,30</sup>. In our study, the SPI-1 and SPI-4 genes were shown highly expressed in *S. Choleraesuis* in SILAC data. High expression of these proteins might contribute to *S. Choleraesuis* to breach the intestinal barrier to further cause systemic infection.

After penetrating the intestine barrier, survival in the macrophage is an important issue for *Salmonella* to cause systemic infection. *S. Choleraesuis* showed better intra-macrophage survival than *S. Typhimurium*. SPI-2 T3SS and *spv* genes play a critical role in intracellular survival and systemic infection<sup>31–33</sup>. Our results showed that

the level of proteins encoded in SPI-2 and *spv* was expressed at least 5-fold higher in *S. Choleraesuis*. This may explain why *S. Choleraesuis* had better intra-macrophage survival than *S. Typhimurium*.

HilA is a key positive regulator that directly binds the promoter of the structure and effector genes in SPI-1 T3SS<sup>34</sup>. HilA is negatively regulated by HilE which binds to HilD to inhibit the transcription of *hilA*<sup>17</sup>. In the genomic analysis of *S. Choleraesuis*, *hile* was predicted as a pseudogene. We found in this study that the plasmid-harboring *S. Typhimurium* or *S. Choleraesuis hile* ORF could restore the *hile* function in *S. Typhimurium hile* deletion strain, meaning that the HilE in *S. Choleraesuis* could still be functional. Recently Nuccio and Baumler analyzed different *Salmonella* serovars with their genome sequences available in NCBI<sup>35</sup>. They redefined some pseudogenes to normal genes, including *hile* in *S. Choleraesuis*<sup>35</sup>. Combining the genomic and functional assays, the hyperexpression of SPI-1 T3SS in *S. Choleraesuis* found in this study appears not related to *hile*, which previously was thought to be a pseudogene.

Environmental signals, such as oxygen, Mg<sup>2+</sup>, and osmolarity, also regulate the SPI-1 T3SS expression. High oxygen concentration is a negative regulator for *hilA* expression. High oxygen inhibits the expression of *hilD* which down regulates *hilA*<sup>11</sup>. We tested the sensitivity to oxygen of the two serovars. The expression of *hilA* and *hilD* in *S. Choleraesuis* was higher in high and low oxygen conditions (data not shown). Thus, oxygen appears not the factor that causes SPI-1 hyperexpression in *S. Choleraesuis*. On the other hand, PhoPQ TCS negatively regulates SPI-1 T3SS after sensing Mg<sup>2+</sup> in the environment. When *Salmonella* invades into cells, it encounters a low pH and Mg<sup>2+</sup> environment. Low Mg<sup>2+</sup> activates the PhoPQ TCS to suppress SPI-1 T3SS<sup>36</sup>. However, under high Mg<sup>2+</sup> condition, Mg<sup>2+</sup> binds to PhoP, which in turn activates its phosphatase activity to remove the phosphate from autophosphorylated PhoP<sup>37</sup>. The expression of *hilA* and *hilD* showed significant difference in the two serovars under low Mg<sup>2+</sup> concentration. Interestingly, *hilD* in *S. Choleraesuis* was expressed higher at normal and high Mg<sup>2+</sup> concentrations. Moreover, the downstream gene, *pagC*, was less expressed in *S. Choleraesuis*. Taken together, *S. Choleraesuis* seemed to be more sensitive to Mg<sup>2+</sup>. High concentration of Mg<sup>2+</sup> might cause more PhoP dephosphorylation, which subsequently inhibits the PhoPQ to suppress the *hilA* and *hilD*, or to activate the *pagC* in *S. Choleraesuis*. In conclusion, this study provides sufficient *in vitro* evidence to support that *S. Choleraesuis* is more invasive than *S. Typhimurium* by hyperexpressing T3SS virulence genes. These findings are consistent with the clinical observation that *S. Choleraesuis* is among the non-typhoid *Salmonella* serovars more invasive to the host.

## Materials and Methods

**Bacterial strains, plasmids, and mutagenesis.** All *Salmonella* strains and plasmids used in this study are listed in Table S1. Bacteria were cultured on the Luria-Bartani (LB) agar at 37 °C, with added ampicillin (100 µg/ml), kanamycin (50 µg/ml), and tetracycline (12.5 µg/ml) when appropriate. A λ-red recombinase mutagenesis method was used for  $\Delta$ *hile* mutant construction, as previously described<sup>38</sup>. The primers used to amplify pKD4 kanamycin resistance gene are listed in Table S2. To construct the *hile* complement strain, primers to amplify the *hile*-contained the putative promoter region from the *S. Typhimurium* and *S. Choleraesuis*, respectively. PCR products were digested by *Bam*HI and *Eco*RI at 37 °C for 1 hour. Digested fragments were ligated into pBR322. The plasmids harbored the *hile* from two serovars were then transferred to the  $\Delta$ *hile* mutant strains by electroporation.

**SILAC proteome analysis.** To compare the protein expression pattern between *S. Typhimurium* LT2 and *S. Choleraesuis* SC-B67, we used the SILAC method as previously described<sup>22</sup>. Briefly, *S. Typhimurium* and *S. Choleraesuis* were cultured to late log phase in the SILAC medium (Mg<sup>2+</sup>, 0.4 mM) supplement with different isotope form of amino acids. Equal numbers of the two serovars were mixed together for further protein extraction by sonication. Protein concentration was measured by BCA method (Thermo). 40 µg of protein sample were separated by 2D-SCX/RPLC system (Dionex). Separated samples were analyzed by a LTQ-Orbitrap hybrid mass spectrometer (Thermo). Raw peptide sequences collected from the LTQ-Orbitrap were analyzed by Mascot v2.3 and MaxQuant v1.2 (Matrix Science). The sequences of *S. Typhimurium* (NC\_003197) and *S. Choleraesuis* (NC\_004631) were downloaded from NCBI and used as references for comparison. All experiments of SILAC analysis were repeated twice.

**qPCR validation.** To test that the mRNA expression was consistent with SILAC results, *Salmonella* strains were cultured in SILAC media to late log-phase. Wild type,  $\Delta$ *hile*, and *hile* complement strain of SL1344 were cultured in DMEM (Mg<sup>2+</sup>, 0.8 mM) or RPMI (Mg<sup>2+</sup>, 0.4 mM) media to late log-phase for analyzing the expression of *hilA* and *hilD*. To analyze the *hilD* expression under different oxygen level, SL1344 and SC-B67 were cultured in DMEM with (high oxygen) or without (low oxygen) shaking to late log phase. To compare the *hilD* and *pagC* expression level in different Mg<sup>2+</sup> concentration, SL1344 and SC-B67 were inoculated in the N-salt media with 8 µM (low), 1 mM (normal), and 10 mM (high) of Mg<sup>2+</sup> as previously described<sup>39</sup>. Bacteria RNA were isolated by TRIzol reagent (Invitrogen) according to the phenol-base method. RNA were treated with DNase I (Fermentas) 20 min at 37 °C to remove genomic DNA and then purified using a RNA clean up kit. Before converting RNA to cDNA, 1 µg of total RNA were used in PCR reaction to make sure that DNA has been removed. 1 µg of total RNA were reverse transcribed to cDNA by ToolsQuant II Fast RT kit (Tools Biotechnology Co., Ltd.). qPCR experiments were performed with SybrGreen Supermix (Bio-Rad) in an iCycler iQ5 (Bio-Rad) instrument. Expression of each gene was normalized to that of 16S rDNA. All the primers used in the experiments were listed in Table S2. All the qPCR experiments were done in triplicate in each independent experiment.

**Acid tolerance assay.** *S. Choleraesuis* SC-B67 and *S. Typhimurium* SL1344 were precultured in LB broth and then 1 ml of the culture was transferred to the tube containing 4 ml of simulated gastric fluid (SGF, pH 2.0).

Suspensions were cultured in the 37 °C incubator without shaking and survival was monitored every 5 mins. SGF contained 8.3 g/L proteose-peptone (Sigma), 3.5 g/L D-glucose (Sigma), 2.05 g/L NaCl (Sigma), 0.6 g/L KH<sub>2</sub>PO<sub>4</sub> (Sigma), 0.11 g/L CaCl<sub>2</sub> (Sigma), 0.37 g/L KCl (Sigma), 0.1 g/L lysozyme (Sigma), and 13.3 mg/L pepsin (Sigma)<sup>40</sup>. Final pH was adjusted to 2.0 with sterile 6.0 M HCl (Sigma). The recovered bacteria were counted by serial plating on LB agar at each time point with appropriate dilutions in PBS. Because the incubation period was short, experiment was done one group in each independent experiment. The experiment was repeated three times independently. We combined three independent experimental results for further statistical analysis.

**Invasion assay.** MDCK cells were used to generate polarized epithelial cell monolayer.  $1 \times 10^6$  cells were cultured in 6-well plate for 5 days<sup>29</sup>. MDCK and HeLa cells, polar and non-polar, respectively, were infected by *S. Typhimurium* and *S. Choleraesuis* with MOI 100 as previously described<sup>29</sup>. At 1 hour post-infection, cells were washed with PBS and treated with gentamicin (75 µg/ml) for 30 mins. Cells were lysed by lysis buffer (0.5% Triton X-100 in PBS pH 7.4) and plated on LB agar with appropriate dilutions. Invasion experiments were done in triplicate in each independent experiment.

**Penetration assay.** To measure the penetration ability of different strains of *Salmonella*, penetration assays were performed by using MDCK and Caco-2 cell monolayer with MOI 100, as previously described<sup>41</sup>. Penetrated *Salmonella* organisms were retrieved from the basolateral medium and were plated onto the LB agar media at 1, 3 and 6 hours after infection. *E. coli* RDEC-1, a non-invasive strain, was used as a negative control in every experiment to ensure the integrity of the cell monolayers<sup>41</sup>. Experiments were done in triplicate in each independent experiment.

**Intra-macrophage survival assay.** Intra-macrophage survival of *Salmonella* in THP-1 cells was determined by a gentamicin protection assay, using methods described previously<sup>9</sup>. At 4 and 6 hours post-infection, intracellular bacteria were recovered with a lysis buffer (0.5% Triton X-100 in PBS pH 7.4) and were plated onto LB agar with or without appropriate antibiotics. Colonies were counted on the next day to calculate the intracellular survival rate. All the experiments were done in triplicate in each independent experiment.

**Mouse infection model and competitive index assay.** Six- to eight-weeks-old female BALB/c mice were purchased from National Laboratory Animal Center, Taiwan. Mice were monitored daily during the experiments and sacrificed when it showed moribund or pain outcome. All the experiments were approved and followed the national animal care guidelines and the Institutional Animal Care and Use Committee (IACUC) of Chang Gung University (approval No CGU13–112). *Salmonella* were grown to the late-log phase. Bacteria were then washed twice and resuspended in saline. Six to ten mice were grouped and infected orally with *S. Typhimurium* or *S. Choleraesuis* ( $1 \times 10^7$  cfu/mouse). Organs were harvested at 1-, 3-, and 5-day(s) post-infection and homogenized in saline. Bacteria were cultured on LB agar with appropriate dilutions. In competition assays, six- to eight-weeks-old female BALB/c mice were infected by *S. Typhimurium* and *S. Choleraesuis* mixed culture ( $2 \times 10^7$  cfu/mouse). At 5 days post-infection, bacteria were recovered from the organs and plated onto LB agar media with or without chloramphenicol (30 µg/ml) to differentiate between *S. Choleraesuis* SC-B67 (resistant to chloramphenicol), and *S. Typhimurium* SL1344 (susceptible to chloramphenicol). Competitive Index (CI) is defined as the ratio between *S. Choleraesuis* and *S. Typhimurium* within output divided by the ration within the input.

**Statistical analysis.** All the experiments were repeated three times. All the quantitative data in this study was performed mean  $\pm$  standard error. The statistical analysis was calculated by the sigma plot (version 10.0). Unpaired t-test was used to compare the results in the study.

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## Acknowledgements

This study was supported by grants from Chang Gung Memorial Hospital (CMRPG3A1113, CMRPG3D0951, CMRPG3D1721, and CMRPG3E1371). We thank Drs Yi-Ywan M. Chen and Hsin-Chih Lai in Chang Gung University and Chyi-Liang Chen in Chang Gung Memorial Hospital for their advice and suggestions during the study.

## Author Contributions

K.-Y.H. and C.-H.C. designed and performed the study, analyzed the data and drafted the manuscript; Y.-I.W., K.-Y.C. and R.P.J. contributed to collection and analysis of the data; all authors edited the manuscript.

## Additional Information

**Supplementary information** accompanies this paper at <http://www.nature.com/srep>

**Competing financial interests:** The authors declare no competing financial interests.

**How to cite this article:** Huang, K.-Y. *et al.* Hyperinvasiveness of *Salmonella enterica* serovar Choleraesuis linked to hyperexpression of type III secretion systems *in vitro*. *Sci. Rep.* **6**, 37642; doi: 10.1038/srep37642 (2016).

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