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## *Salmonella* Degrades the Host Glycocalyx Leading to Altered Infection and Glycan Remodeling

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Complex glycans cover the gut epithelial surface to protect the cell from the environment. Invasive pathogens must breach the glycan layer before initiating infection. While glycan degradation is crucial for infection, this process is inadequately understood. *Salmonella* contains 47 glycosyl hydrolases (GHs) that may degrade the glycan. We hypothesized that keystone genes from the entire GH complement of *Salmonella* are required to degrade glycans to change infection. This study determined that GHs recognize the terminal monosaccharides (*N*-acetylneuraminic acid (Neu5Ac), galactose, mannose, and fucose) and significantly ( $p < 0.05$ ) alter infection. During infection, *Salmonella* used its two GHs sialidase *nanH* and amylase *malS* for internalization by targeting different glycan structures. The host glycans were altered during *Salmonella* association via the induction of *N*-glycan biosynthesis pathways leading to modification of host glycans by increasing fucosylation and mannose content, while decreasing sialylation. Gene expression analysis indicated that the host cell responded by regulating more than 50 genes resulting in remodeled glycans in response to *Salmonella* treatment. This study established the glycan structures on colonic epithelial cells, determined that *Salmonella* required two keystone GHs for internalization, and left remodeled host glycans as a result of infection. These data indicate that microbial GHs are undiscovered virulence factors.

Epithelial cells in the human gastrointestinal tract (GIT) are covered with at least two glycan layers, composed of multiple layers of glycoproteins (mucin) and complex oligosaccharides (glycocalyx) that protect cells from the local environment and infection<sup>1</sup>. Mucin is the most distal layer of glycoproteins in the GIT lumen and is directly exposed to the luminal microbiome. The glycocalyx layer is adjacent the epithelial membrane that is composed of trans-membrane glycoproteins and glycolipids and are components of membrane lipid rafts that extend from the membrane, which are specific bacterial and viral receptors used for microbial invasion resulting in transduction of extracellular signals into the cell<sup>2–5</sup>. Glycans represent the first and crucial interface of the cell surface through which microbes interact with the host immune system mediating recognition and communication processes; thus, controlling immunological recognition, cell-cell adhesion, and pathogen binding. Glycans are mixture of structures that include short chains of saccharides, with simple structures to highly branched complex oligosaccharides that are complicated with an array of specific linkages between the monosaccharide residues leading to a large diversity in arrangements during their synthesis to form higher order chemical structures<sup>1</sup>. In the case of bacterial pathogens, this layer provides a barrier to physically exclude microorganisms from gaining access to the epithelial membrane and the associated receptors used for infection and have been suggested to be used for co-evolution with commensal bacteria<sup>2</sup>.

Bacterial interaction with the epithelial surface via the glycan has been recognized for many years with the use of lectins to identify microbial activity<sup>6</sup>. Beyond system level interactions specific interactions with via fucose

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and sialic acid have been implicated to regulate commensal interactions and provide sugar sources in complex communities where some members may cleave the sugar for use by other community members<sup>6–10</sup>. The specific linkages (i.e.  $\alpha$ -1,2/3/4 and  $\alpha$ -2,3/6 linkages), in addition to these specific sugars are also implicated in controlling the microbial interaction<sup>11</sup>. The breath of the microbes that interact with mucin and the underlying glycolyx interactions is large and has been suggested to be a primary underpinning of co-evolution of an individual and their microbiome<sup>11,12</sup>. In fact, recognition of the glycan as the primary interacting surface has also led to detailing the very large and complex carbohydrate digestion enzymes in bacteria that are assembled in CAZymes<sup>13</sup>. Use of the host glycan to provide nutrients that regulate bacterial infection and virulence is increasingly recognized as an important characteristic for individual microbes to penetrate the mucin layers and subsequently gain access to the cell membrane for association and invasion to progress into the disease state<sup>14,15</sup>.

*Salmonella*, and other invasive enteric pathogens, developed mechanisms to breach the protective glycan layers of host cells, which permits access to the host membrane receptors, that results in invasive intracellular infection. The diversity and complexity of the entire epithelial glycan barrier mandates that a breadth of enzymatic activities is required to degrade these diverse glycans to gain access of the cell membrane. Degradation of these compounds relies in part on glycosyl hydrolases (GHs). GHs are diverse, widely distributed in bacteria, yet poorly characterized enzymes, which hinder determining their specific role during infection dynamics<sup>16,17</sup>. To date there are no studies that define the specific role of the diverse GHs found in *Salmonella* during infection<sup>16,17</sup>. In this study, we hypothesized that only a portion of the 21 different GH families that encompass 49 total carbohydrate-active enzymes in *Salmonella* may degrade glycans to gain access of the host membrane and the microbe receptors used for invasion. Eilam *et al.* demonstrated that microbe's glycan degradation potential is associated with gut pathogens<sup>18</sup>. It was hypothesized in this study that unique and specific GHs in *Salmonella* are required for invasive infection.

Here, we conducted a detailed study of how *Salmonella* degrades the glycolyx layer of human colonic epithelial cells (Caco-2) during *in vitro* infection and led to host glycan remodeling. We established that two GHs,  $\Delta$ *nanH* and  $\Delta$ *malS*, decrease invasion to non-invasive levels comparable to  $\Delta$ *invA*, suggesting that these GHs may be new virulence factors. The *N*-glycome of Caco-2 cells during infection was profiled and showed how *Salmonella* used its glycan-degrading enzymes, *nanH* and *malS*, to degrade the glycolyx layer. More surprisingly, we identified that the host cell responded to microbial glycan degradation by modifying its own glycans leading to decreased sialylation; however, increased fucosylation, higher-mannose, and more hybrid glycans. This alteration in host glycans was due to induction of *N*-glycan biosynthesis pathways during *Salmonella* association. This study demonstrates that a complex molecular interplay between epithelial cells and pathogens result in altered association, but infection proceeds if bacteria have the appropriate GH compliment to overcome the dynamic changes in the glycan.

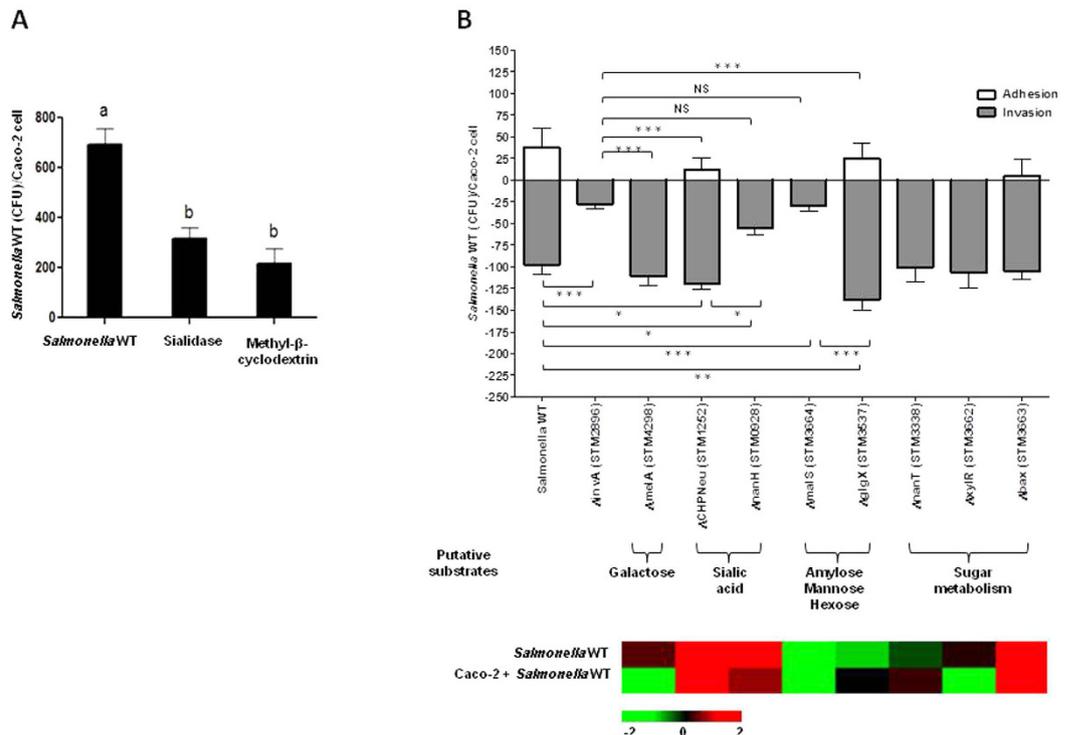
## Results

**Glycan Degrading Enzymes Alter Host Membrane Access during Invasion of *Salmonella*.** To determine if glycans are involved during infection, we first investigated if the removal of Neu5Ac (the outermost monosaccharide on most *N*-glycans of animals) from Caco-2 cells affected *Salmonella* attachment. Caco-2 cells were exposed to sialidase digestion to remove terminal Neu5Ac. As a control, Caco-2 cells were also subjected to methyl- $\beta$ -cyclodextrin (M $\beta$ CD) treatment since M $\beta$ CD inhibits caveolae-dependent endocytosis via disruption of lipid rafts<sup>19</sup>. Depletion of cell surface Neu5Ac led to a significant ( $p < 0.05$ ) reduction in *Salmonella* association (Fig. 1A), further implicating the importance of Neu5Ac in the adherence process. To determine if genes related to metabolism of sugars released from glycan are involved during infection, *nanT* (STM3338; sialic acid transporter), *xylR* (STM3662; xylose operon regulatory protein), and *bax* (STM3663; hypothetical protein similar to ATP-binding protein) were deleted using homologous recombination<sup>20</sup>, characterized in their capacity to infect, and for alteration in invasion *in vitro* with Caco-2 cells (Fig. 1B). Deletion of these genes did not significantly change association compared to *Salmonella enterica* subsp. *enterica* serovar Typhimurium strain LT2, suggesting that genes related to metabolism of sugars are not involved in additional invasion.

Deletion of each gene was verified using whole genome sequencing of the wild type and each deletion mutant (Supplementary Figure S1B and C). Reference based assembly and *de novo* assembly showed that the genomes were isogenic, that only the gene of interest was specifically replaced with the chloramphenicol resistance gene, and that the genome-to-genome distance was not significantly different ( $p > 0.001$ ). Taken together, these data indicate that the gene deletions were adequate to delineate the multi-gene effect demonstrated in this work to digest complex glycans to gain access to the host membrane.

To further test the hypothesis if glycan-degrading GHs alter host membrane access during infection, we analyzed the gene expression of the annotated GHs of *Salmonella* Typhimurium LT2 during *in vitro* infection of Caco-2 cells (Supplementary Table S1). Knowing the most common terminal residues on glycans (i.e. Neu5Ac, galactose, and mannose), GHs that can cleave the most common residues and linkages, along with gene expression analysis, we identified that sialidases (STM0928 and STM1252), galactosidase (STM4298), and amylases (STM3664 and STM3537) likely play an important role during infection. To determine if these GHs are important for infection, these GHs in *Salmonella* Typhimurium LT2 were also deleted (Supplementary Figure S1A) and used to characterize their ability to infect and alter invasion *in vitro* using differentiated (i.e. polarized) colonic epithelial cells (Caco-2 cells) (Fig. 1B). These enzymes targeted different glycans based on their terminal sugar specificity, as seen with the differences in the infectivity of each bacterial mutant (Fig. 1B), further implicating terminal sugar digestion could be important during infection so that the T3SS can gain access to the membrane to initiate injection of effector molecules.  $\alpha$ -Galactosidase (STM4298; *melA*) recognizes terminal  $\alpha$ -galactose molecules of glycans. Deletion of *melA* led to a similar invasion phenotype as the WT.

*Salmonella* LT2 contains two sialidases in its genome: *nanH* (STM0928) and *CHPNeu* (STM1252; Conserved Hydrolase Putative Neuraminidase), both of which were deleted from the genome in this study. Genetic



**Figure 1. Modulation of carbohydrate-degrading enzymes alter access during invasion of *Salmonella* WT.** (A) Neu5Ac digestion. Methyl- $\beta$ -cyclodextrin (control) disrupts lipid rafts. Sialidase treatment led to reduction in *Salmonella* association. Least Squares Means Differences (LSD) was used for statistical analysis. Levels not connected with the same letter are significantly different,  $p < 0.05$ . (B) *Salmonella* WT knockout strains characterized for the alteration in adhesion and invasion (A/I) using differentiated Caco-2 cells 60 minutes post-infection. White bars represent the CFU of *Salmonella* WT that adhered per Caco-2 cell. The gray bars represent the CFU of *Salmonella* WT that invaded per Caco-2 cell. This was done in combination with transcriptional profiling (bottom panel) of *Salmonella* WT during infection of Caco-2 cells to gain insights about differentially expressed carbohydrate-degrading genes. *Salmonella* WT genes displaying changes in gene expression levels during infection of Caco-2 cells. Colors indicate the expression of each gene induced (red) and repressed (green). LSD was used for statistical analysis. Error bars indicate SEM between 3 biological replications, \* $p < 0.05$ , \*\* $p < 0.001$ , \*\*\* $p < 0.0001$ , Not Significant (NS). The statistics shown at the top indicates the statistical relevance in invasion levels of  $\Delta invA$  compared to the mutant strains.

comparison of these genes found that *nanH* has homology with genes from bacteria and parasites (Supplementary Figure S2A). Gene homology for *CHPNeu* was broadly distributed among bacteria from the gut and environment (Supplementary Figure S2B and S2C). In *Salmonella* there is very little DNA sequence homology between *nanH* and *CHPNeu*; however, the domain structures were conserved, which allowed identification of function between organisms to provide molecular markers to define two different neuraminidases in *Salmonella* that are enzymatically, and likely play different roles different during infection (Supplementary Figure S3). Nitrogenases among microbial genomes also display wide sequence diversity, but retain conservation of enzymatically important domains<sup>21</sup>. Deletion of *nanH* significantly decreased invasion ( $p = 0.0059$ ). The  $\Delta CHPNeu$  mutant significantly increased invasion ( $p = 0.01$ ).

The amylase genes (*mals* and *glgX*) are also divergent in their sequence, but both contain the required domains for amylase activity (Supplementary Figure S4). Deletion of *mals* (STM3664;  $\alpha$ -amylase) significantly decreased *Salmonella* invasion ( $p < 0.0001$ ), resulting in the same level of invasion as  $\Delta invA$  (STM2896; needle complex export protein for T3SS), which is considered non-pathogenic and deficient for adhesion and invasion *in vivo*<sup>22</sup>. Surprisingly, deletion of *glgX* (STM3537; glycogen debranching enzyme), which is an amylase-like enzyme with broader hexose specificity, led to a significant increase in invasion ( $p = 0.0002$ ). Perhaps the deletion of *CHPNeu* and *glgX* affected other genes involved in virulence, such as the *nanH* and *mals*. The invasion levels of *nanH* and *mals* mutants were not significantly different ( $p > 0.05$ ) compared to those found for  $\Delta invA$ ; hence, *nanH* and *mals* may represent new virulence genes in *Salmonella* infections. These observations led to the hypothesis that *Salmonella* targets specific glycan structures during infection that enable the T3SS needle complex access to the membrane since it is not long enough to penetrate the glycan structures on epithelial cells.

***Salmonella* Infection Led to Alteration in Host N-Glycome.** We initiated the glycan definition by determining the glycan structure and composition of uninfected Caco-2 cells to define the N-glycome baseline to allow comparison of the changes made by *Salmonella* during infection. This approach is of particular interest because it extends the specificity beyond binding via lectin-like activities and can be directly linked to enzyme







## Discussion

Use of mucin and the glycocalyx by the microbiome is widely recognized<sup>6</sup>. Degrading the glycocalyx layer disrupts the host's defensive barrier that provides access to the epithelial cell surface and receptors for binding, colonization, and invasion. Because *Salmonella* infections represent a persistent and major health challenge around the world it is important to understand how to control *Salmonella* infections. This study examined the specific enzymes used to degrade the glycan.

The sequence of events leading to *Salmonella* infection in the gut epithelial cells has been elucidated for the T3SS<sup>22</sup>. However, prior to T3SS (*invA*) access to the membrane, *Salmonella* must overcome the protective glycocalyx layer that coats the epithelium. In the gut *Salmonella* must first attach to and secondly degrade the host's protective mucus and glycocalyx layer to gain access to the membrane and subsequently infect the underlying tissue. These results described here first identified that depletion of Neu5Ac from Caco-2 cells led to decreases in *Salmonella* association; therefore, this interaction lends insight into how *Salmonella* reaches the initial goals necessary to establish infection by using Neu5Ac as a potential receptor for adherence process.

In order to breach the glycocalyx layer, pathogens have evolved mechanisms to degrade the glycocalyx, which is common in many organisms<sup>6–10</sup>. *Salmonella* specifically is equipped with 48 enzymes from 21 families of glycosyl hydrolases to overcome this barrier. This study found expression of specific glycan-degrading enzymes in *Salmonella* used to degrade the glycocalyx and alter the host glycans to mediate invasion, which were previously unrecognized genes important to the virulence mechanism.

While the microbe is degrading the host glycan, it is also very important to study the host response during degradation. Efforts were made to improve our understanding of glycan degradation and alteration complexity in its entirety via the use of high resolution mass spectrometry<sup>24</sup> and gene expression profiling. The host genes that hydrolyze mannose, fucose, and Neu5Ac from the glycan; as well as sialyltransferases, were induced during *Salmonella* infection (Fig. 3); supporting the hypothesis that new glycan structures were produced *de novo* (Fig. 2F) during infection. Hooper *et al.* and Bry *et al.* also found *in vivo* that the host produced new glycans containing fucosylation in the ileum that is microbe induced via  $\alpha$ 1,2-fucosyltransferase transcripts<sup>6,7</sup>. Refinement of the glycan by the *Salmonella* during infection was also observed (Fig. 4A). The shift to low Neu5Ac content is supported by these changes in sialidases and sialyltransferases (Fig. 3D,E, Supplementary Table S4). The accumulation of fucosylated glycans is supported by the changes in host fucosidases and fucosyltransferases (Fig. 3C) after infection, as previously observed<sup>6,7</sup>. Pickard *et al.* also observed that rapid fucosylation appears to be a protective mechanism that utilizes the host's resources during host-microbe interactions during pathogen induced stress<sup>25</sup>. These observations confirm the host response to glycan modification during *Salmonella* LT2 association and suggest a dynamic shift in the glycan is caused by *Salmonella* degradation coupled with the host remodeling to produce a glycan of different structures as the infection progresses. This could be a protective mechanism for the host during infection and suggests that the host is actively producing new glycans in response to *Salmonella* infection. The mechanism dictating this phenomenon has yet to be elucidated, which requires additional experimentation to uncover the direct effect of *Salmonella* on host glycan metabolism. Glycan biosynthesis is significantly affected by the disease states and distinct glycan structures could provide information about the specific pathologic states of disease<sup>23</sup>. The host constantly remodels its glycans without altering its intrinsic function, which alters microbiome association<sup>3</sup> and modulates host immune surveillance methods<sup>26</sup>. The host may alter its glycan composition on their cell surfaces to eliminate the expression of a terminal glycan structures in order to limit pathogen binding. The host may discard non-critical glycans to allow its survival<sup>3</sup>. The loss of a particular glycan may involve inactivation of one or more genes involved in glycan biosynthesis<sup>27</sup> and can prevent recognition by pathogens using structure as a receptor. In an effort to alter its glycans to evade pathogens, the host may create new glycan structures either by synthesis or modification.

By combining microbial genomics, glycan profiling, and infection analyses, this study provides unprecedented molecular details for the role of *Salmonella* GHs' during infection of Caco-2 cells. Alterations in the degree of branching, changes in the amount, linkage, and degrees of sialylation and fucosylation in *N*-glycans have been reported as a consequence of diseases<sup>28,29</sup>, but this is the first report to describe how *Salmonella* Typhimurium degraded the glycans that resulted in an infection. Deleting GHs used for glycan degradation resulted in invasion magnitude equal to those associated with T3SS in *Salmonella*, suggesting that these enzymes may be as important as secretion systems. Defining the virulence characteristics and diversity of GHs in *Salmonella* can provide new insights into the importance of glycan structures during host-pathogen interactions. This study further expands our understanding of the infection characteristics and may lead to the host response in glycan modification as mutualistic events between the host production and pathogen degradation. Some of the more encouraging prospects of this research are the potential for new treatments for gastroenteritis caused by *Salmonella*. This work may provide the basis of novel strategies to control enteric infections by targeting glycocalyx-degrading enzymes. The complexity of the pan-genome of *Salmonella*, as well as the multiple methods of infection used by *Salmonella*<sup>30,31</sup> demands the use of multi-omics to define innovative targets to control infection. Use of antibiotics alone is not meeting the needs to control this organism and often leads to increased susceptibility to other pathogens<sup>32</sup>. Use of glycan degradation will preempt *Salmonella* from gaining access of the membrane – a novel method to control infection that may be of use in multiple pathogens in the gut and other epithelial surfaces.

## Methods

**Cell Culture.** All cell *in vitro* experiments were done using colonic epithelial cells (Caco-2; ATCC HTB-37) as described previous<sup>33–37</sup>. Detailed description of the cell growth conditions is available in Supplementary Information.

**Bacterial Strains and Growth Conditions.** *Salmonella enterica* subsp. *enterica* serovar Typhimurium strain LT2 (ATCC 700720; *Salmonella* WT) and the deletion mutants made in this study were used. All isolates

were grown in LB (Difco, BD) at 37 °C with shaking at 220 rpm for 14–16 hours before use. Each biological replicate was done with a new vial of frozen stock after thawing and growth twice as described above.

**Gene Deletion.** Bacterial gene deletions were done as described by Datsenko and Wanner<sup>20</sup>. Detailed description of gene deletions is available in Supplementary Information.

**Genomic sequencing and comparison.** Each isolate was sequenced as described by Ludeke *et al.*<sup>38</sup> as defined by the methods of the 100 K Pathogen Genome Sequencing Project<sup>39–47</sup>. Abyss 1.5.2 was used to assemble the paired end reads using  $k = 64$ <sup>48</sup>. Prokka was used for annotation<sup>49</sup>. Each genome was compared to the wild type by determining the genome distance using Genome-to-Genome Distance Calculator (GGDC) (<http://ggdc.dsmz.de/distcalc2.php>)<sup>50,51</sup>. Whole genome comparisons were done using Mauve under Progressive Mauve as described by Darling *et al.*<sup>52,53</sup>. Contigs were reordered using the *reorder contigs* option in Mauve with default parameters using *Salmonella* Typhimurium LT2 ATCC 700720 (accession number AE006468) as the reference genome. All raw genome sequences generated in this study are available in the NCBI SRA as part of the 100 K Pathogen Genome Project Bioproject Accession PRJNA186441. Accession numbers are listed in Table S5. Single gene analyses were done by extracting sequences from each genome and using MUSCLE through Geneious version 6.1.8 to align sequences<sup>54,55</sup>.

**Bacterial Association Measurements.** Bacterial association was determined using a modified gentamicin protection assay<sup>56,57</sup> after adding each *Salmonella* treatment in a 96-well plate containing an MOI of 1:1000. The assay was done after incubation for 60 min at 37 °C with 5% CO<sub>2</sub>. Adhered bacteria were measured after the cell culture medium was aspirated and the host/microbe cell mixture was washed once with 200 µl of 1X PBS buffer (pH 7). The host and associated microbes were lysed using 50 µl of commercial Warnex lysis buffer (AES Chemunex) for use in qPCR assays to determine the absolute amount of host and bacteria<sup>57</sup>. All assays were done in three biological replicates.

To calculate number of adhered bacteria, the mean of the number of invaded bacteria was subtracted from the mean of the total number of host associated bacteria. The error for adhered bacteria was propagated using equation  $(\Delta Z)^2 = (\Delta A)^2 + (\Delta B)^2$  where  $\Delta Z$  is the standard error of mean (SEM) for adhered bacteria,  $\Delta A$  is SEM for total host associated bacteria and  $\Delta B$  is SEM for invaded bacteria. One-way ANOVA with Tukey test was done to find significant differences across treatment's and control's group means.

**Gene Expression during Infection.** Samples were collected as previously described<sup>32,34,58,59</sup>. Briefly, Caco-2 cells were cultured in T-75 flasks (BD) and were serum starved 24 h before infection. Respective bacterial treatments with *Salmonella* WT and the deletion mutants, at MOI of 1000, were used to infect epithelial cells as described previously. All treatments were incubated at 37 °C with 5% CO<sub>2</sub> for 60 min. 10 ml of TRIzol LS reagent (Invitrogen, Carlsbad, CA) was added to the cells and mixed with pipette followed by centrifugation at  $7,200 \times g$  for 5 min to pellet the host associated bacteria. TRIzol LS supernatant was stored in a clean tube and further processed for RNA extraction from infected Caco-2 cells. The bacterial pellet was suspended in 2 ml of fresh TRIzol LS, gently mixed and further processed for RNA extraction from host associated bacteria. The experiment was done in two biological replicates.

**Bacterial RNA extraction and gene expression.** Sample preparation for gene expression profiling was performed with RNA isolation, which was done using TRIzol LS reagent (Invitrogen) as described previously<sup>32,34</sup>. Total RNA (10 µg in 20 µl) was reverse transcribed into cDNA with 6 µg of random hexamers and 400 U of Superscriptase II (Invitrogen) according to the manufacturer's protocol. The reaction mixture was cleaned by using the Qiaquick-PCR purification kit (Qiagen, Valencia, CA) according to the manufacturer's instructions and as described previously<sup>34</sup>. The purified single-strand cDNA was eluted from the columns twice with a total of 100 µl of nuclease free water (Ambion, Austin, TX). cDNA fragmentation was done using 0.6 U of DNaseI (Promega, Madison, WI) per µg of cDNA, according to the instructions. The fragmented 1 µg of cDNA was labeled using 2 µl of GeneChip DNA Labeling reagent (Affymetrix, Santa Clara, CA) and 60 U of Terminal Transferase enzyme (New England Biolabs, Ipswich, MA). The samples were denatured prior to hybridization, at 98 °C for 10 min followed by snap cooling at 4 °C for 5 min.

**Hybridization and normalization.** Labeled cDNA was hybridized onto a custom made Affymetrix GeneChip designed against all the annotated coding sequences of *Salmonella* LT2 ATCC 700720 (*Salmonella* WT)<sup>32,34,60</sup>. The chips were hybridized and scanned at the Center for Integrated BioSystems (Utah State University, Logan, UT) as per manufacturer's protocols. Raw data (.cel files) was background corrected, quantile normalized and summarized using MS-RMA<sup>32,61</sup>. The resultant normalized Log<sub>2</sub> transformed intensity matrix was used for further statistical analysis.

**Caco-2 RNA extraction and gene expression.** The TRIzol LS samples containing infected or non infected Caco-2 cells were frozen (Liquid N<sub>2</sub>) and thawed (70 °C) twice. To 750 µl of TRIzol LS sample, 250 µl of water was added. This was further processed for RNA extraction using manufacturer's (TRIzol LS, Invitrogen) instructions. Synthesis of cDNA, biotin labeled cRNA, fragmentation and purification of cRNA were carried out using one-cycle cDNA synthesis kit (Affymetrix, Santa Clara, CA).

**Host hybridization and normalization.** Labeled and fragmented cRNA (10 µg) was hybridized onto the Affymetrix HGU133Plus2 GeneChips as per manufacturer's recommendations at the Center for Integrated BioSystems (Utah State University, Logan, UT). Raw data (.cel files) was background corrected; quantile normalized and summarized using RMA. RMA normalized data was then filtered through the PANP algorithm to make

presence-absence calls for each probe set. Probe sets that were called present in at least one of the samples were included in further statistical analysis.

**Statistical analysis for gene expression.** Gene expression profiles for *Salmonella* WT alone and in the presence of the epithelial cells were obtained 60 min post infection. The data was analyzed as two class unpaired with T statistic, using Significance Analysis of Microarrays (SAM)<sup>62</sup>. All the genes were ranked based on the score from SAM output. This pre-ordered ranked gene list was then used in Gene Set Enrichment Analysis software (GSEA) to detect the coordinate changes in the expression of groups of functionally related genes, upon respective treatments. The gene sets were defined based on the annotations from Comprehensive Microbial Resource (CMR), Cluster of Orthologous Groups of proteins (COGs), and Virulence Factors of pathogenic bacteria Data Base (VFDB).

**Glycan Degradation.** Differentiated Caco-2 cells in T-75 flasks were infected with *Salmonella* WT and its knockouts at MOI 1:1000 and incubated for 60 min at 37°C, 5% CO<sub>2</sub>. Infection samples were washed three times with ice cold 1X PBS to remove non-adherent bacteria and cellular debris. Cells were scraped from the flask with cell scraper and were kept on ice until cell membrane extraction and N-glycan release as detailed in Supplemental Information online. N-glycans were enriched and analyzed using Agilent HPLC-Chip-QTOF MS (Agilent, CA) as detailed in the Supplementary Information. N-Glycans were identified by composition with a retrosynthetic library using accurate mass according to mass tolerance, retention times, and abundance information<sup>24</sup> and further verified by tandem MS (Supplementary Figure S6).

**Ingenuity Pathway Analysis.** QIAGEN'S Ingenuity Pathway Analysis (IPA, QIAGEN Redwood City, www.qiagen.com/ingenuity) software was used to determine biological pathways associated with our gene expression data (Ingenuity Systems, <http://www.ingenuity.com>, IPA release version summer 2014) similar to He *et al.*<sup>33</sup>. Networks representing molecular interaction were constructed based on the IPA database. Determination of pathway associations was determined through IPA (Fisher's exact test).

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## Author Contributions

N.A. designed experiments, constructed deletion mutants, conducted bacterial infection and cell culture experiments, analyzed MS data, glycan, and gene expression data, extracted DNA, analyzed genome sequences, and wrote the manuscript; D.P., C.C.W. performed glycan assays and determined glycan structures; S.F. analyzed the host gene expression; A.M.W. analyzed genome sequences; B.C.H. constructed DNA libraries for sequencing; P.D., J.S. performed enzymatic glycan degradation, conducted gene expression experiments, and constructed deletion mutants; R.J. helped with the analysis of MS data; N.K. constructed deletion mutants; C.B.L. reviewed manuscript; B.C.W. conceived of the hypothesis, planned experiments, critically accessed the data, and wrote the manuscript.

## Additional Information

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