Up-Regulation of MUC2 and IL-1β Expression in Human Colonic Epithelial Cells by *Shigella* and Its Interaction with Mucins

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**Abstract**

**Background:** The entire gastrointestinal tract is protected by a mucous layer, which contains complex glycoproteins called mucins. MUC2 is one such mucin that protects the colonic mucosa from invading microbes. The initial interaction between microbes and mucins is an important step for microbial pathogenesis. Hence, it was of interest to investigate the relationship between host (mucin) and pathogen interaction, including *Shigella* induced expression of MUC2 and IL-1β during shigellosis.

**Methods:** The mucin-*Shigella* interaction was revealed by an in vitro mucin-binding assay. Invasion of *Shigella dysenteriae* into HT-29 cells was analyzed by Transmission electron microscopy. *Shigella* induced mucin and IL-1β expression were analyzed by RT-PCR and Immunofluorescence.

**Results:** The clinical isolates of *Shigella* were found to be virulent by a congo-red binding assay. The in vitro mucin-binding assay revealed both *Shigella dysenteriae* and *Shigella flexneri* have binding affinity in the increasing order of; guinea pig small intestinal mucin < guinea pig colonic mucin < Human colonic mucin. Invasion of *Shigella dysenteriae* into HT-29 cells occurs within 2 hours. Interestingly, in *Shigella dysenteriae* infected conditions, significant increases in mRNA expression of MUC2 and IL-1β were observed in a time dependent manner. Further, immunofluorescence analysis of MUC2 shows more positive cells in *Shigella dysenteriae* infected HT-29 cells than untreated cells.

**Conclusions:** Our study concludes that the *Shigella* species specifically binds to guinea pig colonic mucin, but not to guinea pig small intestinal mucin. The guinea pig colonic mucin showed a greater binding parameter (R), and more saturable binding, suggesting the presence of a finite number of receptor binding sites in the colonic mucin of the host. In addition, modification of mucins with TFMS and sodium metaperiodate significantly reduced mucin-bacterial binding; suggesting that the mucin-*Shigella* interaction occurs through carbohydrate epitopes on the mucin backbones. Overproduction of MUC2 may alter adherence and invasion of *Shigella dysenteriae* into human colonic epithelial cells.

**Introduction**

Mucosal surfaces employ a number of protective strategies to defend against noxious substances and pathogens found within the intestinal lumen. The mucosal surface contains mucins, which are complex glycoproteins synthesized and secreted by epithelial cells of various organs to lubricate and protect luminal surfaces of the human body [1]. However, excessive mucin secretion is a hallmark of the pathogenesis of several diseases, including infectious diseases such as inflammatory bowel diseases and Shigellosis [2–4]. The exact role of mucins in gut protection is not completely understood. Although adherence is recognized as an important initial step in other bacterial infections [5–7], relatively little information is available on the mechanisms of attachment of *Shigella* to cells on the mucosal surface. A specific site on the host may be involved in the binding of pathogenic bacteria; for example, *Hemophilus influenzae*, a respiratory pathogen, specifically binds to respiratory mucins but does not bind or cause disease in the intestinal tract. Also, enterotoxigenic *E. coli* infects and causes diseases only in the intestinal tract [8,9].

Infection of epithelial cells with bacterial pathogens can induce the excessive production of intracellular adhesion molecules (ICAM-1) and mucins (MUC2 and MUC5AC) through the
activation of TNFα and Interleukin-1 secretion [10–14]. In vivo inoculation of Shigella species altered the expression of MUC2 and MUC5AC in rabbit intestinal epithelial cells, through production of the inflammatory cytokine TNFα [4]. Proinflammatory cytokines such as TNFα and IL-1β can induce the production of MUC2 and MUC5AC in human intestinal and airway epithelial cells [15]. Shigella flexneri stimulates differential mucin gene expression in human colon cancer cells, thereby reducing the production of inflammatory cytokines [16]. The production of different mucins with different levels of expression might involve protection of host epithelium from Shigella flexneri induced inflammation [16].

Shigellae can induce acute intestinal inflammation through the production of many inflammatory cytokines and chemokines from infected epithelial cells [17]. However, the exact relationships between bacterial infection and types of mucin gene expression under shigellosis have yet to be examined. The present study aimed to examine the interaction between Shigella species and intestinal mucin in an attempt to identify region-specific initial binding sites for Shigella species. Furthermore, to evaluate the involvement of inflammatory cytokines in the hyper production of MUC2 during shigellosis.

**Methods**

**Bacterial strains, media and growth conditions**

Clinical isolates of *Shigella dysenteriae* (*S. dysenteriae*) and *Shigella flexneri* (*S. flexneri*) obtained from the Department of Medical Microbiology, Christian Medical College, Vellore, India, were routinely grown in Luria-Bertani (LB) broth or Tryptic soy broth (Hi-media, Mumbai, India) at 37°C.

**Isolation and purification of Guinea pig intestinal mucin**

The mucin glycoprotein was isolated from Guinea pig as described previously [18–20]. Briefly, Male Duncan Hartley strains of Guinea pig were fasted for 48 h and sacrificed by cervical dislocation. The intestine was removed and suspended in ice-cold sterile PBS containing protease inhibitors [5 mM EDTA, 1 mM PMSF and 10 mM N-ethyl maleimide (NEM)]. The mucosal surface was gently scraped using a rubber spatula, taking care not to damage the underlying epithelium. The mucus scrapings were homogenized and redissolved in 50 mM Tris-HCl (pH 8.0) containing 4 M guanidium hydrochloride, 0.02% sodium azide, 1 mM PMSF, 5 mM EDTA and 2 mM N-ethyl maleimide, by stirring overnight at 4°C. The solution was centrifuged at 10,000 x g for 20 min at 4°C to remove debris and insoluble material. The mucins were partially purified by Sepharose CL-4B column chromatography. The fractions showing positive reaction with Morgan and Elson assay reagent were pooled, dialyzed and lyophilized. The partially purified mucin samples were stored at −20°C.

**Modification of mucins**

Partially purified mucin from Guinea pig small intestine and colon was subjected to various physicochemical treatments and was used for *in vitro* binding studies.

**Tetramethyl urea (TMU) treatment**

Treatment of bacteria with Tetra methyl urea was carried out as described in [21]. Briefly, Bacterial cells were incubated with 0.5 M TMU (final concentration) for 1 h at 37°C. Bacteria was harvested by centrifugation, washed to remove excess TMU, and then added to wells coated with mucin.

**Oxidation**

50 mg of mucin (dry weight), dissolved in 3 ml of PBS (pH 7.4), was treated with sodium metaperiodate (3.1–100 mM) at 4°C for 1 hr in the dark for nonspecific oxidation of sugar moieties. Oxidized mucin was then dialyzed against distilled water for 24 h and used for binding assays.

**Boiling**

Guinea pig colonic mucin (10 mg/ml) was heated in a water bath at 100°C for 10 min and cooled to room temperature. These boiled mucins were used for binding assays.

**Partial deglycosylation**

Partial deglycosylation of mucin was done as described previously [22]. Guinea pig colonic mucin (~25 mg dry weight) was treated with 1 ml of TFMS-Anisole reagent at 0°C, on ice. The reaction was terminated by adding a two-fold excess of ether; precooled to 0°C, drop wise, avoiding bubbling due to exothermic heat. The upper layer of ether was removed by centrifugation at 1500 rpm/10 min. The aqueous phase was dialyzed against distilled water to remove traces of ether and pyridinium salts. The dialysate was lyophilized to be used as partially deglycosylated mucin for binding studies.

**Enzyme-linked immunosorbent assay**

This assay was done as described in [23]. Briefly, microtiter plates (Nunc, immunomodule Maxisorp Denmark) were coated with serially diluted mucin (150 μg mucin protein/well to 1 μg mucin protein/well) in 0.05 M Carbonate buffer (pH 9.6) and incubated overnight at 4°C. The unbound mucin was removed by three washes with PBS-T (0.05%Tween 20). Residual binding sites were blocked with PBS-2%BSA and incubated at 37°C for 1 h. After removing unbound BSA by three washes with PBS-T, 100 μl containing different concentrations of (cfu/ml) bacteria were added to each well and incubated at 37°C for 1 h. The unbound bacteria were removed by washing three times with PBS-T. 100 μl of mouse monoclonal antibody against *Shigella* species (E-Merck, Mumbai, India) (1:1000 in PBS-1%BSA) was added to each well and incubated at 37°C for 1 h. The unbound antibody was removed by washing three times with PBS-T. 100 μl of Rabbit anti mouse HRP conjugated secondary antibody (1:2000 in PBS-1%BSA) was added to each well and incubated at 37°C for 1 h. The unbound antibody was removed by washing three times with PBS-T. Then, the wells were incubated in the dark with 100 μl of substrate (TMB/H2O2) for 15 min. 100 μl of 1 M H2SO4 was added to arrest the reaction. The colour developed was measured at 450 nm in a Perkin-Elmer micro plate reader. All experiments were done in triplicate and repeated thrice. Specific binding to mucin (binding parameter ‘R’) was calculated by subtracting the value of bacteria bound to BSA from the values of bacteria bound to mucin. A value of >2 indicates specific binding. Pig gastric mucin was (Sigma, USA) used as a negative control.

**Virulence assay**

The virulence nature of *Shigella* species were assessed by Congo-red dye binding assay and the procedure was followed as described previously [24]. Briefly, bacterial strains were grown in Congo red (0.01%) supplemented Tryptic soy broth containing 0.6% yeast extract and 1.5% agar at 37°C for 18 h.

**Cell culture conditions**

HT 29 human colon tumor cell line was obtained from National Centre for Cell Science, Pune, India. HT 29 cells were grown in...
Cells were washed and the non-specific binding sites were blocked permeablized with 0.1% Triton X-100 in PBS for 5 min on ice.

Infection of HT 29 cells with Shigella dysenteriae

HT 29 cells were seeded into 6 well Costar tissue culture plates at a density of 2 x 10^5 cells/ml in volumes of 2 ml per well. At this seeding density, monolayers were sub-confluent (80–90%) at the time of the experiment. Bacteria were grown overnight at 37°C in LB medium and pelleted by centrifugation at 12,000 g for 5 min at 4°C. The cell pellets were washed with PBS (pH 7.4) twice and suspended in antibiotic-free DMEM. Bacteria at 100 cells per epithelial cell (100:1 ratio) were used to infect for 2 h to allow bacterial entry to occur. Monolayers were washed twice to remove extracellular bacteria and the cultures were incubated for different time intervals in the presence of 50 μg/ml of gentamicin to kill the remaining extracellular bacteria.

Investigation of bacterial adhesion and invasion of HT 29 cells by Transmission Electron Microscopy (TEM)

Infection of epithelial cells with Shigella dysenteriae was done as described above. The infection was stopped at 1 h and 2 h time interval for adherence and invasion analysis by TEM as described previously [25]. Infected epithelial monolayers on culture dishes were fixed in 3% (v/v) glutaraldehyde (in 0.1 M sodium cacodylate buffer, pH 7.4) for 2 h. All the samples were washed in PBS and post fixed in 1% osmium tetroxide for 1 h before dehydration in ethanol and embedded in Epon resin, according to standard procedures. Ultrathin sections (60–100 nm) were cut, and uranyl acetate and lead citrate stains were applied prior to examination and photography with a Philips 201C (Netherlands) transmission electron microscope.

Immunofluorescence studies for mucin gene expression

HT 29 cells infected with Shigella dysenteriae were grown for 2 h on cover slips for complete and tight attachment and washed with DMEM in the presence of 50 μg/ml of gentamicin to kill the remaining extracellular bacteria. The infected monolayers were incubated for 0, 1, 3, 6 and 9 h. The infected monolayers were washed with PBS for 5 min and subsequently the cells were permeabilized with 0.1% Triton X-100 in PBS for 5 min on ice. Cells were washed and the non-specific binding sites were blocked with PBS containing 0.1% BSA for 10 min at room temperature before antibody incubation. Cells were washed twice with PBS/BSA and incubated with mouse monoclonal antibody for MUC2 at 1:50 dilution for 1 h at room temperature (E-Merck, Mumbai, India). After the incubation period, monolayers were washed twice with PBS and incubated with secondary antibody (anti-mouse FITC) for 1 h in the dark. After the incubation period, monolayers were washed twice with PBS and the cover slips were mounted with 90% glycerol in PBS. The cells were screened under a fluorescence microscope (Carl Zesis, Germany) with 20X magnifications.

RT-PCR analysis of Mucin and IL-1β gene expression

Total RNA was extracted from control cells, cells preincubated with actinomycin D (1 μg/ml) for 30 min and Shigella dysenteriae infected HT 29 cells, using Trizol reagent (Sigma–Aldrich, USA). For cDNA synthesis, 2 μg of total RNA were used as the template in a 20 μl RT reaction mixture by using Reverse-iTTM first strand synthesis kit (ABgene, UK). 100 ng of cDNA template in a total volume of 25 μl was used for reverse transcription. PCR analysis of mRNAs of MUC2, IL-1β and β-actin were carried out by using forward and reverse primers listed below; for MUC2 gene, 5′-ACAACTACTGCCTGCTACCC-3′ and 5′-GTTGATCTCGTACTGTTGAGGCA-3′; for IL-1β gene, 5′-AAACAGATGGAAGTGCTCCTTCCAGG-3′ and 5′-TGGAGACACCACCTTGTTGCTCCA-3′; for β-actin gene, 5′-GTTGGGCGCCCCAGGCACCA-3′ and 5′-CTCCTTATGTCCGGACGGATTCC-3′. The PCR reaction conditions for all the genes were as follows: 94°C for 2 min (1 cycle); 94°C for 30 s, 58°C for 30 s, and 72°C for 1 min (30 cycles); and 72°C for 5 min. The PCR products were subjected to electrophoresis (130 V, constant-voltage field) on a 1% agarose gel equilibrated in Tris-borate electrophoresis buffer containing ethidium bromide (1 mg/ml). Gels were photographed under UV light. The band intensity was measured by using Image J program.

Results

Virulent nature of Shigella

The virulent nature of the Shigella species was analyzed by in vitro Congo-red binding assay, appearance of orange/pink color colonies of S. dysenteriae (Fig. 1a) and S. flexneri (Fig. 1b) suggest that these species are highly virulent when compared to pale white colonies of S. flexneri (Fig. 1c) and S. dysenteriae (Fig. 1a) on Congo-red agar plate. Orange-pink colour colonies of Shigella dysenteriae (a) and Shigella flexneri (b) indicate the highly virulent nature and plasmid DNA cured non-virulent Shigella flexneri (c) shows white colour colonies. doi:10.1371/journal.pone.0027046.g001

Figure 1. Congo-red binding assay. Shigella species were grown in Congo-red agar plate. Orange-pink colour colonies of Shigella dysenteriae (a) and Shigella flexneri (b) indicate the highly virulent nature and plasmid DNA cured non-virulent Shigella flexneri (c) shows white colour colonies.
color colonies of plasmid DNA cured non-virulent strain of *Shigella flexneri* (Fig. 1c).

Bacterial binding to native mucin

The binding of *Shigella dysenteriae* and *Shigella flexneri* to human colonic mucin and guinea pig colonic mucin was concentration dependent and saturable. Human colonic mucin protein (12.5 μg per well) and Guinea pig colonic mucin protein (50 μg per well) gave sufficient receptor sites (Fig. 2a and 3a). Bacterial concentrations of 1.6 × 10^7 cfu/well for *S. dysenteriae* and 1.5 × 10^7 cfu/well for *S. flexneri* gave maximal binding to mucins, while bacterial concentrations higher than 1.5 × 10^7 cfu/well, the receptor mucin was saturated (Fig. 2b and 3b). The guinea pig small intestinal mucin and Pig gastric mucin concentration of

**Figure 2. Binding of *Shigella dysenteriae* to mucins.** a. Binding of *Shigella dysenteriae* to varying concentrations of different mucins. b. Binding of different amount of *Shigella dysenteriae* to human and guinea pig mucins. Each value is the mean of triplicate determinations. The assay was carried out as described in materials and methods.
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100 μg/well did not accommodate the same population of bacteria ($1.5 \times 10^7$ cfu/well). The number and affinity of binding sites for *S. dysenteriae* and *S. flexneri* was found to be increasing in the order of guinea pig small intestinal mucin (GPSIM) < guinea pig colonic mucin (GPCM) < human colonic mucin (HCM). The binding results suggest a very high degree of site specificity and

Figure 3. Binding of *Shigella flexneri* to mucins. a. Binding of *Shigella flexneri* to varying concentration of different mucins. b. Binding of different amount of *Shigella flexneri* to human and guinea pig mucins. Each value is the mean of triplicate determinations. doi:10.1371/journal.pone.0027046.g003
species specificity. Table 1 represents the binding parameters of S. dysenteriae and S. flexneri to mucins from different species. None of the clinical isolates bound to unrelated mucin, such as Pig gastric mucin. The human colonic mucin was used as a positive control.

Bacterial binding to modified mucin

The possibility of non-specific hydrophobic interactions mediating the binding to BSA and mucin was investigated by using an effective hydrophobic interaction inhibitor, Tetramethyl urea. Bacterial binding to small intestinal mucin of guinea pig and BSA were diminished at 0.5 M concentrations of TMU (data not shown). However, TMU, did not have any significant effect on S. dysenteriae and S. flexneri binding to guinea pig colonic mucin. The adherence of S. dysenteriae and S. flexneri to boiled mucin was significantly higher than the untreated native mucin (Fig. 4a and 4b), whereas, treatments of mucins with periodate and TFMS showed significantly reduced binding to S. dysenteriae and S. flexneri (Fig. 4a and 4b). Guinea pig colonic mucins that were treated with varying concentrations of periodate show markedly decreased binding to S. dysenteriae and S. flexneri, in a concentration dependent manner (Fig. 5a and 5b).

Adhesion and invasion of Shigella dysenteriae to HT29 cells by TEM

The control HT 29 cells showed intact nuclei, cytoplasm and other cell organelles (Fig. 6a). HT 29 cells infected with Shigella dysenteriae (1–2 h) showed the intimate contact between the bacterium and host cell membrane (Fig. 6b). Also, endocytic processes, formation of pseudopod projections and phagocytosed bacteria were located in the vacuoles of Shigella dysenteriae infected HT 29 cells (Fig. 6c).

Expression of MUC2 and IL-1β gene in HT 29 cells by RT-PCR

MUC2 (348 bp) and IL-1β (388 bp) gene expressions were observed at basal levels in control cells, whereas, in the S. dysenteriae infected conditions, higher levels of MUC2 (~4–5 fold for 9 h) and IL-1β (25 fold for 9 h) mRNA were observed in a time dependent manner (Fig. 7). Prior to S. dysenteriae infection, HT 29 cells were pre-incubated with actinomycin D (1 μg/ml) for 30 min, which showed lessIL-1β expression with increasing time of incubation (Fig. 7). Amplification of β-actin (131 bp) was used as an internal control.

Immunofluorescence analysis of MUC 2 expression

HT 29 cells were infected with Shigella dysenteriae for 2 hours and allowed to incubate at different time intervals (0–9 hours) before analysis of MUC2 expression. Approximately 300 cells were positive with green fluorescence of MUC2 expression. Control HT 29 cells showed normal basal level expression of MUC2 (+) up to 9 h of incubation (Fig. 8a), whereas, HT 29 cells infected with Shigella dysenteriae showed increased expression of MUC 2 in a time dependent manner (+, 1 h; ++, 3 h; ++++, for 6 and 9 h) (Fig. 8b–e).

Discussion

The pathogenesis of most of enteropathogens occurs by successful colonization, adherence and invasion on the epithelial cells in the gastrointestinal tract. Previous studies suggest that the invasion of enteropathogens into intestinal epithelial cells is an important step in virulence mechanism of Shigella pathogenesis [26]. During the course of infection, bacterial colonization occurs by the interaction of their adhesin[s] to cognate receptors available on epithelial cells or the extracellular matrix [27]. Reports are available on the mechanism of initial attachment by various pathogens like Vibrio cholerae [28], Salmonella typhimurium [29], and Yersinia enterocolitica [30] to the host mucus layer and intestinal epithelial cells.

Earlier work on the in vitro binding of S. flexneri [31–32] and enteropathogenic E.coli [33] to intestinal epithelial cells showed that the adherence of the bacterium is due to the presence of plasmid DNA (~140 MDa) encoded factors. These are known as Invasive plasmid antigens (Ipa). IpaA, IpaB, IpaC, IpaD, IcsA and IcsB, and are essential virulence factors for attachment, invasion and intracellular spread [34–36]. The virulent nature of S. dysenteriae and S. flexneri were determined by Congo red binding on agar plate, where highly virulent Shigella dysenteriae showed orange/pink color colonies, suggesting that plasmid DNA and its associated virulence factors are intact, whereas, white color colonies indicate non-virulent bacteria [24].

The mucosa, containing mucin glycoprotein, is thought to have dual roles, one in protecting the host from pathogens, as well as being able to facilitate the preliminary attachment by the pathogens [37]. On the other hand there are reports emphasizing role of mucin glycoproteins in pathogenesis of Helicobacter pylori induced gastric carcinoma [38] and Pseudomonas aeruginosa infection in cystic fibrosis (CF) patients [39]. To determine whether the mucin layer of the gastrointestinal tract acts as a receptor for Shigella species for firm attachment, and if so, is it the protein core or the oligosaccharide structures, which is involved. The virulent S. dysenteriae and S. flexneri showed preferential binding to human colonic mucin compared with all other mucins. It was clearly evident that the number and affinity of binding sites for Shigella dysenteriae and Shigella flexneri were found to be increasing in order of guinea pig small intestinal mucin < guinea pig colonic mucin <

Table 1. The binding parameters of S. dysenteriae and S. flexneri to mucins from different species.

| Mucin                        | Binding parameter 'R' | S. dysenteriae | S. flexneri |
|------------------------------|-----------------------|----------------|-------------|
| Human colon mucin            | 7.45±0.326–1.055±0.321| 6.769±0.289–0.854±0.19 |
| Guinea pig colon mucin       | 3.357±0.138–0.753±0.228| 2.937±0.129–0.680±0.184 |
| Guinea pig small intestinal mucin | 1.415±0.249–0.693±0.182 | 1.228±0.162–0.746±0.101 |
| PGM porcine type II mucin    | 0.918±0.18–0.662±0.182 | 0.868±0.167–0.332±0.046 |

Microtiter plates were coated with a fixed concentration of mucin (12.5 μg for human; 50 μg for guinea pig). 100 μl of bacteria (~1×10^9 cfu/well) was added. The values are the mean ± S.D triplicate determinations of one representative experiment. (PGM, Pig gastric mucin). doi:10.1371/journal.pone.0027046.t001
human colonic mucin. The binding results suggest a very high degree of site specificity and species specificity of Shigella species. Similar binding patterns were observed in human colonic mucins with Shigella dysenteriae binding [40]. It provides the evidence that the Shigella species is able to bind to human colonic mucin with high affinity as compared with guinea pig colonic mucin.

Various pathogens have their own tissue tropism. Staphylococcus has a binding affinity to ferret nasal and bovine submaxillary mucin [41]. The peptide backbone of the mucin glycoprotein is unique in some aspects, and shows much more uniformity in profile than do oligosaccharide chains [42]. Diversity of mucin-oligosaccharide structure calls for selective affinity of organisms to mucins of different sources. The Guinea pig colonic mucin (GPCM) may harbor more receptor sites than the others, but less than human colonic mucin receptor sites.

Binding of Shigella to guinea pig small intestinal mucin and BSA via the influence of hydrophobic interaction was ruled out, as there was no influence on S. dysenteriae and S. flexneri binding to TMU treated guinea pig colonic mucin. Thus, the receptor-mediated binding of Shigella to GPCM is evident. Further, Mucin pretreated with sodium metaperiodate showed decreased binding, which may be due to oxidation of vicinal hydroxyl groups on carbohydrate moiety [43]. Partial deglycosylation of carbohydrate moieties with TFMS resulted in decreased binding, indicating carbohydrate chains were essential components of the binding sites of GPCM receptor for Shigella species. Similar results were identified in many pathogens [44–46].

Denaturation of mucins by boiling at 100°C resulted in significant increases the pathogen binding, suggesting that dissociation of the polymer into monomeric glycoproteins exposes a number of new receptor sites for the bacteria. Similar observations were reported earlier with Yersinia enterocolitica [47]. It is very clear from this study that S. dysenteriae and S. flexneri bind specifically to the carbohydrate portion of Guinea pig colonic

Figure 5. Adherence of S. dysenteriae and S. flexneri to sodium metaperiodate treated mucins. S. dysenteriae (a) and S. flexneri (b) showed significantly reduced adherence to sodium metaperiodate treated mucins in a concentration dependent manner when compared with untreated mucins. Each value is the mean of triplicate determinations. A p value of less than 0.05 is considered to be significant. Each comparison was made on bacterial adherence to untreated versus treated mucins.

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mucin. Hence, mucin is almost certainly an early site for adherence and establishment of colonization of *Shigella* species.

The pathogenesis of *Shigella* species is mostly contributed by its ability to invade the colonic mucosal epithelium [48]. It still remains unclear whether the initial binding to the colonic mucin would block or facilitate further binding and invasion of *Shigella* to the underlying epithelial cells. Our study shows that most of the bacterial cells entered into the HT 29 cells within 2 h of infection. This invasion was confirmed by transmission electron microscopy and it revealed the occurrence of bacterial adherence, bacteria-induced pseudopod formation, invasion and accumulation in the host cell cytoplasm. This study confirms the previous findings on the invasive nature of *Shigella* species [48].

The relationship between bacterial infection and over expression of MUC2 and MUC5AC were emphasized especially in the respiratory tract and middle ear epithelial cells [11,49–52]. *Pseudomonas aeruginosa*, a common Cystic fibrosis pathogen, activates MUC2 mucin gene transcription by activation of a Src-dependent Ras-MEK1/2-ERK1/2-pp90rsk-NF-κB pathway [53]. Our previous findings on rabbit ileal loop infection assay, in vivo, with *Shigella* species showed up-regulation of MUC2, MUC5AC and proinflammatory cytokine expressions [4]. The present study shows that the expression of MUC2 transcript was increased in a time-dependent manner in HT 29 cells infected with *Shigella dysenteriae*. Further, its expression at protein level was confirmed by immunostaining infected cells with MUC2 antibody. The results revealed a time-dependent increase of MUC2 expression. This study suggests that the expression of MUC2 was up-regulated at both transcriptional and translational levels. These observations suggest that mucins are normally expressed in human colonic epithelial cells, but under infectious conditions, are upregulated, and may be involved in the mucociliary clearance of enteric pathogens.

The expression of the proinflammatory cytokine IL-1β was observed in *Shigella* infected macrophages [54]. The major function of IL-1β is to recruit the inflammatory polymorphonuclear cells that infiltrate to the infected site and destabilize the epithelium [55]. In addition to inflammation, it can also induce the expression of MUC2 and MUC5AC mRNA in intestinal and airway epithelial cancer cell lines [56–58]. Similar findings were also observed with IL-4, IL-13 and IL-19 on induced mucin gene expression in various epithelial cells [55]. The present study also confirms the previous reports by showing a time-dependent increase in IL-1β and MUC2 expression during *S. dysenteriae* infection of human epithelial cells. Further, IL-1β expression was depleted upon treatment with actinomycin D, which provided additional evidence that *S. dysenteriae* could induce the expression of IL-1β in infected host cells. The over expression of MUC2 may be due to increased expression of IL-1β, consistent with the examples stated above. Further studies are required to identify the mechanism of mucin gene expression and its role in protection against Shigellosis.

**Figure 6.** *Shigella dysenteriae* adherence and invasion of human epithelial cells by Transmission Electron Microscopy. After 1 h and 2 h of infection, cells were fixed and processed for Transmission Electron Microscopy analysis as described in materials and methods. Control cells showed intact nuclei and other cell organelles (a). 1 h of infection of HT 29 cells with *Shigellae* showed the adherence and pseudopod projection formation (b). Endocytic processes and phagocytosis of bacteria was seen in the vacuoles of HT 29 cells at the end of 2 h infection (c). The scale bars indicate 1 μm. [M – Mucin; A – Adherence; P – Pseudopod formation; I - Invasion].

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Figure 7. RT-PCR analysis of MUC2 and IL-1β expression in *Shigella dysenteriae* infected HT 29 cells. HT 29 cells were infected with *Shigella dysenteriae* for different time intervals. After infection, mRNA was isolated for MUC2 and IL-1β gene expression analysis as described in methods. HT-29 control cells showed basal level expression of both MUC2 and IL-1β. Whereas, higher level (intensity) expression of both MUC2 (~4–5 fold higher for 9 h) and IL-1β (~25 fold higher for 9 h) were seen in a time dependent manner upon HT-29 cells infected with *S. dysenteriae*. HT 29 cells pre-incubated with Actinomycin D (1 μg/ml) for 30 min, prior to *S. dysenteriae* infection, showed lower level of expression of IL-1β. The band intensity was measured by using Image J program. Amplification of β-actin was used as an internal control.

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Figure 8. Immunofluorescence analysis of MUC2 expression in *Shigella dysenteriae* infected HT 29 cells. HT 29 cells were infected with *Shigella dysenteriae* in the ratio of 100 bacteria per epithelial cell. Cell monolayers were washed twice to remove extracellular bacteria and the cultures were incubated for different time intervals (0–9 h) in the presence of 50 μg/ml of gentamicin for mucin gene expression analysis. The percentage (%) of cells which expressed MUC2 was classified as follows: –, none; +, light or weak; ++, moderate; ++++, intense. In each experiment 300 cells were counted. Control cells, up to 9 h showed a weak signal (+) and for 1 h incubation with *S. dysenteriae* (+); for 3 h, (++; for 6 h, (+++); and for 9 h incubation (+++). (20× magnification).

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