Interaction between enteric epithelial cells and Peyer’s patch lymphocytes in response to Shigella lipopolysaccharide: Effect on nitric oxide and IL-6 release

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AIM: To investigate the effect of interaction between enteric epithelial cells and lymphocytes of Peyer’s patch on the release of nitric oxide (NO) and IL-6 in response to Shigella lipopolysaccharide (LPS).

METHODS: Human colonic epithelial cells (Caco-2) were mixed cocultured with lymphocytes of Peyer’s patch from wild-type (C57 mice) and inducible NO synthase knockout mice, and challenged with Shigella F2a-12 LPS. Release of NO and mIL-6 was measured by Griess colorimetric assay and enzyme-linked immunosorbent assay (ELISA), respectively.

RESULTS: In the absence of LPS challenge, NO was detected in the culture medium of Caco-2 epithelial cells but not in lymphocytes of Peyer’s patch, and the NO release was further up-regulated in both cocultures with lymphocytes from either the wild-type or iNOS knockout mice, with a significantly higher level observed in the coculture with iNOS knockout lymphocytes. After Shigella F2a-12 LPS challenge for 24-h, NO production was significantly increased in both Caco-2 alone and the coculture with lymphocytes of Peyer’s patch from the wild-type mice but not from iNOS knockout mice. LPS was found to stimulate the release of mIL-6 from lymphocytes, which was suppressed by coculture with Caco-2 epithelial cells. The LPS-induced mIL-6 production in lymphocytes from iNOS knockout mice was significantly greater than that from the wild-type mice.

CONCLUSION: Lymphocytes of Peyer’s patch maintain a constitutive basal level of NO production from the enteric epithelial cell Caco-2. LPS-induced mIL-6 release from lymphocytes of Peyer’s patch is suppressed by the cocultured epithelial cells. While no changes are detectable in NO production in lymphocytes from both wild-type and iNOS knockout mice before and after LPS challenge, NO from lymphocytes appears to play an inhibitory role in epithelial NO release and their own mIL-6 release in response to LPS.

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Key words: Shigella F2a-12 LPS; Colon epithelial cells (Caco-2); Peyer’s patch lymphocyte; Coculture; Nitric oxide; Interleukin-6

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INTRODUCTION

The intestinal mucosa is an important route of entry for microbial pathogens. The single layer of enteric epithelial cells that lines the intestinal mucosa is the initial site of interaction between the host and entero-invasive microbial pathogens. Following bacterial entry, intestinal epithelial cells rapidly initiate the innate and acquired immune response. Within the first few hours after bacterial invasion, the intestinal mucosa produces mediators that orchestrate the onset of an early inflammatory response. Characteristics of this program include the increased production and release of chemokine(s), cytokines and nitric oxide (NO). These molecules can act as early signals to activate an acute mucosal inflammatory response and enhance the ability of epithelial cells to produce cytokines that regulate mucosal immune responses.

NO is generated by the conversion of L-arginine to L-citrulline by NO synthase (NOS), which exists in three
isoforms, encoded by a separate gene. The expression of inducible NO (iNOS, encoded by NOS2, one of the isoforms) is regulated in various cell types and can be upregulated by stimulation with several cytokines or with bacterial lipopolysaccharide (LPS). iNOS can also be expressed constitutively, in some specialized cells, such as human lung epithelial cells. In human colon epithelial cell lines, the iNOS expression and production of NO are upregulated following stimulation with a combination of interferon-γ (IFN-γ) and interleukin-1 (IL-1) or tumor necrosis factor-α (TNF-α). NO can mediate a wide array of physiological effects in the intestine that are relevant to neuronal regulation of vascular function, and epithelial barrier integrity. Moreover, NO produced by intestinal epithelial cells could play a protective role in the host response to enteric microbial pathogens. Enteroinvasive bacteria directly activate expression of iNOS and NO production in intestinal epithelial cells. The NO in the intestine has been implicated in the suppression of microbial entry into epithelial cells by increasing the protective barrier through the release of epithelial mucins and the induction of intestinal fluid secretion. In addition to direct innate immune action, NO has immuno-regulatory effects relevant to the control of infection on modulating the subsequent acquired immune response.

Interleukin 6 (IL-6) is a pleiotropic cytokine that is produced by many different cell types and mediates several physiological responses, such as humoral immune response, acute-phase reactions, and hemato- poiesis. In the intestine, IL-6 enhances IgA production in murine Peyer’s patch B cells and human appendix B cells that express the IL-6 receptor. Intestinal epithelial cells may also secrete IL-6 and TGF-β to enhance IgA secretion and suppress IgM production by mucosal B cells, whose function is highly dependent upon local production of T cell cytokines. These facts indicate that IL-6 plays an important role in the mucosal immune response.

We have previously established a coculture system of human colonic epithelial cells (Caco-2) with murine lymphocytes of Peyer’s patch and demonstrated that the cell-cell contact between epithelial and immune cells is important in modulating the epithelial barrier and transport function in response to challenge of Shigella LPS. Although accumulating evidence suggests that the interaction or “cross-talk” between epithelial cells and lymphocytes of the intestine is crucial in the immune response to bacterial invasion, no report has clearly demonstrated how the interaction between epithelial cells and lymphocytes, particularly from the Peyer’s patch, affects the release of important immuno-regulatory mediators, such as NO and IL-6, at rest and in response to bacterial infection. The present study was undertaken using the established coculture system of Caco-2 epithelial cells with lymphocytes of Peyer’s patch to investigate NO and IL-6 release in response to Shigella LPS challenge. We also cocultured Caco-2 epithelial cells with lymphocytes from iNOS knockout mice to investigate the involvement of NO in regulation of IL-6 release in response to Shigella LPS.

### MATERIALS AND METHODS

#### Isolation of Peyer’s patch lymphocytes

Wild-type (C57) mice and iNOS knockout mice of C57 background (SPF, 6-8 wk-old) were obtained from the Animal House of Chinese University of Hong Kong. The lymphoid follicles of the mouse Peyer’s patch were excised from the intestinal serosal side and placed in 10 mL PBS, supplemented with 20 mL/L FBS (Invitrogen Co., Grand Island, NY) and 2% penicillin-streptomycin (Invitrogen Co.). The collected patches were triturated by pipetting up and down a few times and smashing through a metallic grid (mesh: 100). Individual lymphocytes were released in the medium below the metallic grid. The lymphocytes were washed with PBS, in which the distribution of Peyer’s patch T and B cell populations was consistent with previous data when they were checked by flow cytometry, and diluted to a concentration of 1 x 10⁶ cells/mL prior to mixed culture with Caco-2.

#### Enteric epithelial cell culture

Human colonic cell line Caco-2 was purchased from American Type Culture Collection (Rockville, MD). The cells were grown in Dulbecco modified Eagle’s minimal essential medium (DMEM; Invitrogen Co.) with 100 mL/L FBS, 2 mmol/L L-glutamine (Invitrogen Co.), 100 µmol/L non-essential amino acid (Invitrogen Co.), 200 units/mL penicillin and 200 µg/ml streptomycin, at 37 °C, 50 mL/L CO₂. To maintain the growth of an uniform polarized epithelial monolayer, Caco-2 cells were seeded at a density of 3 x 10⁵ cells on a floating permeable support, which was made of a membrane filter (Millipore, 0.45 µm pore size) with a silicone rubber ring attached on top of it for confining the cells (0.45 cm² growth area).

#### Coculture configurations

Three types of culture configurations were established in this study: (1) Caco-2 culture alone: Caco-2 cells were cultured as a homogenous polarized monolayer according to the epithelial cell culture method mentioned above and they served as the epithelial cell control; (2) Coculture: Caco-2 cells (3 x 10⁵) completely mixed with Peyer’s patch lymphocytes (1 x 10⁸), and then were seeded on a permeable support described above and maintained up to the 5th d at 37 °C in a 50 mL/L CO₂ atmosphere; (3) Lymphocytes of Peyer’s patch alone: lymphocytes were cultured in an equal volume of the coculture (1 x 10⁶/well in 96-well plate) at 37 °C in a 50 mL/L CO₂ atmosphere as the lymphocyte control.

#### Shigella F2a-12 LPS pretreatment

When the cells cultured on the membrane filter reached confluence on the 4th d, Shigella F2a-12 LPS (5 µg/mL, obtained from the Immunology Department of Institute of Microbiology & Epidemiology, Academy of Military Medical Sciences) was added on the apical side of the epithelium monolayer growing on the membrane filters and treated for 24 h at 37 °C in a 50 mL/L CO₂ atmosphere.
Detection of NO release (Griess colorimetric assay)
After confluent Caco-2 monolayers on the Millipore filter were challenged by *Shigella F2a-12* LPS pretreatment for 24 h, cell culture supernatants were collected and kept at -20°C until evaluation of NO level in the medium by Griess reagent system (Promega, USA). Briefly, nitrite accumulation was determined by mixing equal volumes of cell culture medium and the Griess reagent (10 mL/L sulfanilamide in 50 mL/L phosphoric acid, and 1 mL/L N-1-naphthylenediamine dihydrochloride in water). The absorbance (A550-A540) was read on a 96-well microplate reader. Standard curves were constructed with known concentrations of sodium nitrite. The values for the amount of NO were converted from the absorbance values based on the standard curve constructed for each assay.

Enzyme-linked immunosorbent assay (ELISA)
After confluent Caco-2 monolayers on the Millipore filter were challenged by *Shigella F2a-12* LPS pretreatment for 24 h, the culture supernatants were collected and kept at -20°C until evaluation of mL-6 level in the culture medium. The IL-6 was measured using a mouse IL-6 quantitative colorimetric sandwich ELISA from Biosource (USA) according to the manufacturer’s instructions. The values were calculated on the basis of a standard curve constructed for each assay.

Statistical analysis
All data are expressed as means ± SE. The number of experiments represents independent measurements on separate monolayers. Statistical analysis was performed by repeated measure analysis of variance (ANOVA), and “P” values less than 0.05 were considered statistically significant.

Results

Constitutive release of NO from Caco-2 epithelial cells was potentiated by the cocultured lymphocytes of Peyer’s patch

In the absence of LPS stimulation, measurements of NO from the culture medium showed a moderate level of NO release from Caco-2 cells (67.64 ± 13.79 µmol/L) but almost negligible amount (3.15 ± 0.46 µmol/L) in lymphocytes of Peyer’s patch from wild-type mice; however coculture of both produced over 2-fold (P < 0.001) increase in NO release (Figure 1).

The increase in NO production in the coculture could be due to either potentiation of epithelial release by the cocultured lymphocytes or induction of NO release from lymphocytes by coculturing with epithelial cells. To distinguish this, lymphocytes from iNOS knockout mice were used. If coculture induced NO production in lymphocytes, this would not be observed with iNOS knockout lymphocytes. However, as shown in Figure 2, high level of NO (158.29 ± 2.62 µmol/L) was still observed in the coculture of Caco-2 with iNOS knockout lymphocytes, confirming the epithelial origin of NO. Moreover, coculture with iNOS knockout lymphocytes produced a level of NO significantly greater than that produced by coculture with lymphocytes of intact iNOS (P < 0.001), indicating an inhibitory role of iNOS from lymphocytes in regulating NO release from epithelial cells.

Shigella LPS induced NO release from Caco-2 epithelial cells but with little effect on either lymphocytes of Peyer’s patch or the coculture with lymphocytes of iNOS knockout mice

As shown in Figure 3, the level of NO in the culture me-
Suppression of mIL-6 release from lymphocytes of Peyer’s patch by cocultured epithelial cells

The release of mIL-6 from different cultures was measured using a mouse ELISA kit. The results showed that mIL-6 release was detected from the lymphocytes of Peyer’s patch alone (55.23 ± 2.98 pg/mL) but not from Caco-2 cells, indicating no cross-reactivity of the mouse antibody with human IL-6. When the lymphocytes were cocultured with Caco-2 cells, mIL-6 release was significantly reduced (Figure 4).

LPS triggered mIL-6 release from both lymphocytes of Peyer’s patch and cocultures, which was further enhanced with iNOS knockout lymphocytes

As shown in Figure 5, mIL-6 levels in either lymphocytes alone or in cocultured with Caco-2 cells were significantly increased 24 h after *Shigella* F2a-12 LPS treatment. Furthermore, the levels of LPS-induced increases in mIL-6 release were significantly higher in groups with lymphocytes from iNOS knockout mice (Figure 5), indicating an inhibitory role of iNOS from lymphocytes in regulating mIL-6 release.

DISCUSSION

Kerneis S and colleagues[19] have demonstrated that murine Peyer’s patch lymphocytes cocultured with human differentiated Caco-2 cell monolayers are able to induce a phenotypic conversion of enterocytes into cells sharing structural and functional properties with M cells. This phenomenon reveals the profound influence of Peyer’s patch lymphocytes on intestinal epithelial phenotypes[20,21]. And our previous study indicated that Peyer’s patch lymphocytes may modulate intestinal epithelial barrier and ion transport.
function in homeostasis and host defense via cell-cell contact and cytokine signaling\[^{16}\]. The present study has further demonstrated the importance of the interaction between intestinal epithelial cells and lymphocytes of Peyer's patch in regulating the release of NO and IL-6 from Caco-2 epithelial cells and lymphocytes of Peyer's patch, respectively, at rest and in response to *Shigella* LPS challenge.

The present study has demonstrated constitutive release of NO from enteric epithelial cells but not from lymphocytes of Peyer's patch. We also firstly found that the cocultured Peyer's patch lymphocytes potentiated NO release from the enteric epithelial cells. It is known that the constitutive and iNOS activity has the highest level in intestinal villus cells, which is resembled by Caco-2 cells, and the lowest is in crypt cells\[^{22}\]. Further studies have also demonstrated that the enterocytes regulate their own ion transport processes, either in basal condition or in the presence of active secretion, through the activation of a constitutive NOS-NO pathway, functioning as a braking force of cAMP-induced ion secretion\[^{23}\]. It has also been reported that iNOS-derived NO is a key mediator of early villous re-epithelialization following acute mucosal injury\[^{24}\]. These facts indicate that constitutive NO production may have a physiological role in intestinal epithelial cells. It has been proposed that NO synthesis in intestinal epithelial cells is up-regulated through some molecules or cytokines including T-lymphocytes derived cytokines\[^{25}\] during cell-cell contact with lymphocytes\[^{26}\]. The observed potentiation of NO release from epithelial cells by Peyer's patch lymphocytes is consistent with the above notion. Our data also suggest that NO from the lymphocytes, although almost negligible, may be involved in negatively regulating NO release from enteric epithelial cells since lymphocytes from iNOS knockout mice further enhance NO release from the epithelial cells. This also rules out the possibility of coculture-induced NO release from the lymphocytes, confirming epithelial release of NO potentiated by the lymphocytes.

Upon *Shigella* F2a-12 LPS challenge, the present results showed that NO from epithelial cell Caco-2 significantly increased but not from lymphocytes of Peyer's patch alone groups. Witthoff\[^{18}\] has indicated that entero-invasive bacteria directly activate expression of iNOS and NO production in human colon epithelial cells. NO is involved in the protective mechanisms in the gastrointestinal tract. Enterocyte-derived NO changes intestinal monolayer permeability and ion channel via iNOS and COX-2\[^{27-29}\]. In the acute phase response, *shigellae* inflammation is characterized by increased cell turnover in the lamina propria (LP) and the epithelium, increased iNOS expression in the surface epithelium, and apoptosis, which seems to be associated with LP macrophages\[^{30}\]. However, NO production in the coculture with C57 wild type lymphocytes further up-regulated by the LPS treatment, yet there was almost no increase in NO production in the coculture with iNOS knockout lymphocytes following LPS challenge. This observation suggests that the coculture of Caco2 and iNOS knockout lymphocytes has become insensitive to the LPS challenge. The iNOS knockout lymphocytes might lack the additional factors that are crucial for LPS-induced NO release. This is a likely scenario since recent studies demonstrated that host defense function of NO in intestinal mucosal immune response is regulated by IL-12 and/or IFN-γ\[^{31}\], which are deficient in the iNOS knockout mice\[^{32}\].

Another interesting finding from the present study is that epithelial cells could also suppress mIL-6 secretion by lymphocytes of Peyer’s patch with cell-cell contact. The mIL-6 level in both coculture groups was significantly reduced as compared to that of lymphocytes of Peyer’s patch alone. It is clear that IL-6 is produced by a number of cell types including antigen-presenting cells, lymphocytes and even epithelial cells via stimulating of endotoxin and a number of other stimuli (TGF-β, IL-1β, etc.). However, the mIL-6 in the coculture system is only detectable from mouse lymphocytes of Peyer’s patch in our previous study\[^{33}\]. It should be noted that there is evidence that epithelial cells could inhibit IL-6 secretion from lymphocytes by the interaction between epithelial cells and lymphocytes of Peyer’s patch in homeostasis. After *shigella* LPS treatment for 24 h, mIL-6 level was obviously increased in all groups. More interestingly, the LPS-induced mIL-6 release was significantly greater in groups with lymphocytes from iNOS knockout mice, indicating an inhibitory role of iNOS in regulating mIL-6 release from lymphocytes of their own. Mucosal production of IL-6 is important because this cytokine may regulate a number of local and systemic immune responses, including IgA secretion\[^{34}\], macrophage differentiation, T cell proliferation, and acute phase protein synthesis\[^{35}\] in liver and intestine as well as active transcription of a variety of cytokines and receptors\[^{36}\]. Accumulating evidence\[^{37}\] suggests that NO inhibits the LPS-induced IL-6 production and this effect may be a direct effect or caused by inhibited PGE2 production, or both PGE2 and NO may act in an autocrine or in a paracrine manner. NO from epithelial cells is easy to reconcile NF-κB activity with the needs of host defense through autocrine pathway to produce a rapid response to a pathogenic stimulus that is shut down promptly so as to minimize host damage\[^{38}\]. This study only investigated NO release and mIL-6 secretion from different sources to illustrate the cross-interaction between epithelial cells and lymphocytes in homeostasis and host defense. Further studies are required to identify their regulatory mechanism of NO from intestinal epithelial cells to reduce LPS-induced IL-6 production of Peyer’s patch lymphocytes.

In summary, the cross-interaction between intestinal epithelial cells and lymphocytes of Peyer’s patch regulates host defense mechanism. Lymphocytes of Peyer’s patch can induce a basal constitutive production of NO from epithelial cells. This basal level NO may play an important physiological role in the enteric epithelium in the absence of infection. During infection, NO release from epithelial cells is further upregulated. Conversely, epithelial cells can downregulate IL-6 secretion of Peyer’s patch lymphocytes, while LPS can upregulate IL-6 release. Furthermore, NO from lymphocytes appears to play an inhibitory role in the epithelial NO release and their own IL-6 release in response to LPS. Taken together, NO and IL-6 are important immuno-regulatory mediators in response to bacterial infection through the interaction between epithelial cells and Peyer’s patch lymphocytes.
Conversion by Peyer’s patch lymphocytes of human epithelial barrier and ion transport function by Peyer’s patch and interleukin-2. IgA secretion by intestinal epithelial cell-derived cytokines.

High rate IgA secretion in IgA-committed B cells. Human and murine interleukin 6 induce secretion: studies in rat jejunum in vivo.

Evidence for a cGMP-dependent and a cGMP-independent pathway. Mucin exocytosis from human epithelial cells by nitric oxide: studies in rat jejunum in vivo.

Production in human colon epithelial cells.

Interaction of lipopolysaccharide with human small intestinal lamina propria fibroblasts favors cytokine binding on human intestinal epithelial cells.

Upregulation mediates oxidant-induced disruption of F-actin cytoskeleton.

Regulation of mucosal barrier function and dysfunction.

The two faces of IL-6 on Th1/Th2 differentiation.

The two faces of IL-6 on Th1/Th2 differentiation.

Inhibitory role of nitric oxide on enterocyte-derived cytokines.

Investigation of the intestinal mucosal barrier in human colon epithelial cells: a technical update.