Using an in vitro autologous human system, the immunomodulatory function of colonic intraepithelial lymphocytes (IEL) on cytokine production by lamina propria mononuclear cells (LPMNC) has been investigated. In contrast to LPMNC, colonic IEL produced only low amounts of IL-10, interferon-γ and interleukin-2. However, co-culture experiments (IEL + LPMNC) have shown that IEL can enhance the PHA-induced synthesis of IL-2 and interferon-γ, but not IL-10 by LPMNC. Using a transwell filter culture system apparatus, this effect was shown not to require a cell-to-cell interaction. Thus, IEL in vitro may modulate the cytokine synthesis of LPMNC, through the production of soluble factors. This may prove highly relevant in the in vivo immune activation of the gastrointestinal mucosa.

Key words: Interferon-γ, Interleukin-2, Interleukin-10, Intraepithelial lymphocytes

Introduction

Because of their unique location within the epithelium, their peculiar morphological and phenotypic features (different from those of peripheral blood and lamina propria lymphocytes) and their oligoclonal expression of αβ T cell receptors, many studies have focused on IEL function. Although their precise role in vivo remains enigmatic, it has been shown in vitro that, under certain conditions human IEL (hIEL) exert various patterns of cytotoxicity, are capable of proliferation, and perform some regulatory functions. It is believed that they may interact with the adjacent epithelial cells and underlying lamina propria cells. In particular, we have previously shown that colonic hIEL can inhibit the proliferation of autologous IPL. Moreover, hIEL were also able to suppress IgA synthesis. Inhibitory mechanisms were shown to occur through soluble factors, possibly cytokines.

While some cytokines have a critical role in inducing inflammation, others are crucial in reducing inflammation and mediating the healing process. Most studies on murine IEL (mIEL) have used lymphocytes from the small intestine. Freshly isolated mIEL produce interferon-γ, and IL-5. Subsequently, it has been shown that both TCR-αβ and γδ mIEL could produce interferon-γ, IL-2, IL-3 and IL-6 but not IL-4. However, the amounts of cytokines released by γδ mIEL were much lower compared with αβ mIEL. In addition, culture supernatants from anti-TCR antibody-activated γδ or αβ mIEL trigger synthesis of TNF-α, TGF-β and GM-CSF that influence epithelial function. Murine colonic IEL expressed similar levels of IL-1, interferon-γ and TNF-α mRNA, and significantly more IL-2, IL-4, IL-5 and IL-10 mRNA when compared with small intestine mIEL. Recently, human IEL were analysed for cytokine mRNA expression: freshly isolated hIEL expressed high levels of IL-1β, IL-8, IL-2R and IFN-γ mRNA, but significantly less IL-1α, IL-6, IL-7, TNF-α and TNF-β mRNA. After in vitro polyclonal activation with phytohaemagglutinin (PHA) and PMA, IL-2 and IFN-γ mRNA were significantly increased, whereas IL-1 mRNA was significantly decreased. Furthermore, hIEL can produce small amounts of IL-2 and IFN-γ in response to PHA even though there is little proliferation. The production of these cytokines by hIEL is markedly increased by addition of sheep red blood cells to PHA or after stimulation through the CD2 molecule. In hIEL, PHA induced no IL-
production while superantigen stimulation resulted in detectable IL-4.\textsuperscript{9,21}

It is thus reasonable to consider the possibility that IEL may influence LPL function through the secretion of soluble factors. Therefore, colonic hIEL were tested for their ability to produce IL-2, IFN\textgreek{g} and IL-10. Furthermore, the putative immunoregulatory function of IEL on cytokine secretion by autologous LPL was examined.

**Materials and Methods**

**Subjects**

Normal mucosa from colonic specimens was used from 10 control patients who were undergoing resection for a colonic carcinoma. The specimens were taken at least 5 cm away from any macroscopic lesion.

**Isolation of mucosal lymphocytes**

IEL were isolated as previously described.\textsuperscript{10} Briefly, to isolate colonic IEL, mucosal strips were immersed in RPMI-supplemented with fetal calf serum (FCS), antibiotics (gentamicin 50 \mu g/ml, penicillin 100 U/ml, streptomycin 100 \mu g/ml). The washed specimens were then stirred in the same medium with the addition of 1 mM dithiothreitol (Sigma) for 10 min. After further washing, they were then incubated with calcium- and magnesium-free HBSS and 0.75 mM ethylenediaminetetraacetate (EDTA) in a shaking water bath at 37°C for three periods of 30 min each at 140 oscillations per minute. This step was repeated twice and the resulting supernatants were stored overnight at 4°C. Further purification was achieved by passing this crude preparation down a glass wool column. The filtrate was then centrifuged at 800 g for 30 min through a discontinuous 44/67.5% Percoll bilayer. The IEL were recovered from the interface. To isolate the lamina propria mononuclear cells (LPMNC), the treated tissue was submitted to three more 45-min incubations with calcium- and magnesium-free HBSS and 5 mM EDTA. The mucosal strips were digested overnight at 37°C with 100 ml RPMI 1640 supplemented as above plus 30 mg \textit{Clostridium histolyticum} collagenase (Boehringer Mannheim, Germany). The digestate was passed through a 100 \mu m nylon mesh filter. After separation on Ficoll-paque gradient, the LPL were washed and resuspended in RPMI-supplemented as above. The viability of the cells was assessed using 0.1% trypan blue.

**Preparation of cytokine-containing cell culture supernatants**

hIEL or LPMNC at a concentration of $1 \times 10^6$ were cultured in 1 ml aliquots in 24-well plates with or without 10 \mu g/ml PHA for 72 h at 37°C in 5% CO\textsubscript{2}. The supernatants were then harvested, centrifuged and passed through a 0.22 \mu m filter and stored at –20°C for subsequent cytokine analysis.

**Influence of IEL on the cytokine production by autologous LPL**

LPMNC ($10^6$) were cultured with IEL ($10^6$) for 72 h with or without PHA (10 \mu g/ml) in 24-well plates in a final volume of 1 ml at 37°C in 5% CO\textsubscript{2}. The control cultures consisted of IEL or LPMNC alone, with or without PHA. At the end of the culture, the supernatants were harvested, centrifuged, passed through a 0.22 \mu m filter and stored at –20°C for subsequent cytokine analysis (IL-2, IFN\textgreek{g} and IL-10). In addition, IEL were co-cultured with LPMNC in a Transwell apparatus (Costar Cambridge, MA) (6.5 mm diameter, 3 \mu m pore size) which physically separated IEL from LPMNC. The purpose of this series of experiments was to investigate whether the immunoregulatory function of IEL is mediated by a soluble factor or requires interaction between IEL and LPMNC.

**Cytokine assays**

Gamma-interferon, interleukin-10 (IL-10) and interleukin-2 (IL-2) were measured by a sensitive enzyme amplified immunoassay (Medgenix Diagnostics, Fleurus, Belgium). Optical densities were measured by an automated dual beam ELISA reader at 450 nm for \textgamma-IFN, IL-10 and IL-2. Concentrations of these cytokines were determined by reference to the serially diluted standards included in each plate. The minimum detectable concentrations of these assays were: \textgamma-IFN 0.03 IU/ml; IL-10 1 pg/ml; IL-2 0.1 IU/ml. In these assays, duplicate values varied by < 5% from their mean value.

**Results**

**Cytokine production by colonic mucosal lymphocytes**

The median concentrations of cytokines secreted by IEL and LPMNC alone were respectively 0.3 IU/ml (range 0–0.6) and 1.5 IU/ml (range 1–8) for IFN\textgreek{g}, 1 IU/ml (range 0–20) and 2.1 IU/ml (range 0.8–2.4) for IL-2, 8 pg/ml
(range 1–18) and 17 pg/ml (range 3.5–39) for L-10. Following stimulation with PHA, the median IL-2 concentrations by control IEL and LPMNC cultured alone were respectively 2.5 IU/ml (range 1–26) and 9 IU/ml (range 2–450), the median IFN-γ levels were 27 IU/ml (range 1–105) and 26.5 IU/ml (range 2–242), the median IL-10 concentrations 3 pg/ml (range 1–10) and 4 pg/ml (range 1–100).

Influence of IEL on the cytokine production by autologous LPMNC

After 3 days of culture, the effect of IEL on cytokine synthesis in co-culture with autologous LPMNC was studied. Addition of 10^6 IEL only minimally affected IL-10 production. The median secretion of IL-10 secreted by PHA-stimulated LPMNC was 4 pg/ml (range 1–100). Following the addition of IEL, the concentrations increased to 7.5 pg/ml (range 2–340). In contrast, they enhanced significantly the IL-2 and IFN-γ production by LPMNC. After 3 days of co-culture, IL-2 and IFN-γ levels were respectively 39.5 IU/ml (range 8–450) and 104 IU/ml (18–500) (Figs 1, 2 and 3). Using the Transwell culture apparatus, similar results were obtained (Table 1).

Discussion

The present data confirm our previous work showing that colonic IEL produce very low amounts of interferon-γ. Although there was a significant increase in interferon-γ production following stimulation with PHA, the absolute amounts of interferon-γ were persistently low. Sperber et al. reported similar findings using IEL stimulated with PHA or superantigens. This was not due to the failure of lectin binding. The amount of interferon-γ produced by stimu-
lated hIEL was of sufficient quantity to induce MHC class II expression molecules on HT-29 cells. Class II bearing epithelial cells can in turn cause further activation of hIEL.

IL-2 is a central regulator of immune and inflammatory responses. It affects T cell proliferation as well as B cells and the function of macrophages, natural killer cells and lymphokine activated killer cell. PHA-stimulated colonic hIEL produce similar low amounts of interleukin-2, as do PHA-stimulated jejunal hIEL. Conversely, Sperber et al. showed that IL-2 production was not observed with PHA- or superantigen-stimulated IEL. Interestingly, induction of low levels of IL-2 does not correlate with the induction of in vitro proliferation. However, addition of exogeneous IL-2 does not restore a normal proliferative response.

IL-10 is a powerful suppressor of cytokines produced by macrophages and Th2 subpopulations of CD4+ T cells. In addition, IL-10 inhibits antigen presentation but supports B cell differentiation and IgA secretion. To the best of our knowledge, this is the first observation that human colonic IEL produce small amounts of IL-10 (as do colonic lamina propria cells). However, the precise action of this cytokine in the intestinal mucosa remains to be defined although a colitis occurs in mice in which there has been targeted deletion of the IL-10 gene. Teitelbaum et al. have shown that neutralizing antibody to IL-10 did not block the inhibitory activity of rat intestinal intraepithelial lymphocytes on the proliferative response of allogeneic spleen-thymus cells stimulated with concanavalin A.

Murine helper T cells can be classified in relation to their cytokine production profile. Thus, Th1 cells produce large quantities of IL-2, IFNγ and TNF-β while Th2 cells produce preferentially IL-4, IL-5, IL-6 and IL-10. It has been suggested that a subset of murine CD8 IEL can produce some type 1 and 2 cytokines. Furthermore, it has been shown that some human CD8+ clones may secrete both patterns of cytokines. It thus appears that these two subsets (Th1 and Th2) are mutually antagonistic. Th1 cells may influence Th2 cell function through these cytokines and vice versa. For example, interferon-γ may be able to down-regulate a Th2 response while IL-10 has powerful inhibitory effects of the synthesis of Th1 cytokines.

The crucial role of cytokines in the mucosal response is well illustrated by a number of animal models of inflammatory bowel disease. IL-2, IL-10 and TGF-β1 knockout mice develop chronic intestinal inflammation. The local functions of these cytokines secreted by colonic hIEL have not been fully investigated. The data presented in this paper indicate that colonic hIEL can enhance the PHA-induced synthesis of IL-2 and interferon-γ by human colonic LPMNC. The upregulated LPMNC cytokine synthesis in hIEL+ LPMNC co-cultures revealed that IEL did not contain cells inhibiting synthesis of these cytokines. Furthermore, absolute amounts of IL-10 produced by PHA-stimulated hIEL and LPMNC cultured alone or in co-culture were persistently low. Thus, the increased release of the Th1-cytokines may prove of importance in the modulation of subsequent immunopathological reactions including both activation of inflammatory cells and expression of class II molecules on epithelial cells.

Additional work must be done to understand the hIEL inhibitory activity previously described. As IL-4 is virtually not produced by hIEL, further studies must be done in order to determine other inhibitory cytokines produced by IEL such as TGF-β and other inhibitory factors (e.g. neuropeptides).

It is important to note that cytokines and neuropeptides might in turn influence hIEL function. It has been suggested that interleukin-7 (IL-7) might be produced in the mucosa by intestinal epithelial cells, especially goblet cells. Interleukin-7 is a B- and T-cell growth factor. It affects the proliferation of activated but not unstimulated human T cells. Recent reports have shown that IL-7 stimulates hIEL proliferation. In contrast, IL-9 or IL-12 did not stimulate the proliferation of IEL. IL-4 secreted by activated lamina propria cell suppresses IEL proliferation and LAK activity. Finally two neuropeptides, substance P and vasoactive intestinal peptide induce no proliferation nor spontaneous cytotoxicity by hIEL. This was due to an absence of substance P and vasoactive intestinal peptide receptors.

In conclusion, human colonic IEL may influence the cytokines produced by LPMNC. This in vitro observation may be of importance in the local in vivo immune activation of the gastrointestinal tract.

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