IL-10 and IL-13 Production by Peripheral Blood Mononuclear Cells in Patients with Diabetes Mellitus

Aim: It is generally accepted that proinflammatory cytokines secreted by mononuclear cells are responsible for pancreatic beta (B) cell destruction in animal models of autoimmune type-1 diabetes mellitus (DM). Several studies have shown that markers of inflammation and pro-inflammatory cytokines such as interleukin (IL)-10 and IL-13 associate with the metabolic syndrome, dyslipidemia, type-2 DM and type-1 DM. The aim of this study was to examine the in vitro production of IL-10 and IL-13 in cultures of peripheral lymphocytes obtained from patients with type-2 DM, type-1 DM and healthy controls.

Materials and Methods: Twenty patients (10 females, 10 males) with type-2 DM (median disease duration: 10 years), 10 patients (5 females, 5 males) with type-1 DM (median disease duration: 11 years) and the healthy control group were enrolled in the study. IL-10 and IL-13 in culture supernatants were measured using the enzyme immunoassay method. Statistical analyses were performed using Mann-Whitney U test, Wilcoxon signed rank tests, and Spearman’s rank correlation.

Results: IL-10 and IL-13 levels were similar in culture supernatants incubated with 10 µg/ml phytohemagglutinin (PHA) in all patients and healthy controls (P > 0.05). In type-2 DM, IL-10 levels were significantly lower than in the control group in culture supernatants incubated with 20 µg/ml PHA (P = 0.001 for poor metabolic control and P = 0.048 for good metabolic control, respectively).

Conclusions: According to this study, the alterations in IL-13 did not play an important role in the pathogenesis of type-2 DM and type-1 DM. Furthermore, the deficiency in IL-10 production from peripheral blood mononuclear cells observed with high dosages of PHA suggests that IL-10 may play an important role in the pathogenesis of type-2 DM.

Key Words: IL-10, IL-13, diabetes mellitus
Introduction

Diabetes mellitus (DM) is a syndrome characterized by chronic hyperglycemia, which is accompanied by lipid metabolic disorders. There are two types: non-insulin-dependent diabetes mellitus (type-2 DM) and insulin-dependent diabetes mellitus (type-1 DM). Two major pathogenetic mechanisms are operative in type-2 DM. One of them is impaired islet beta (B) cell function and the other is impaired insulin action. Recently it has been reported that impaired cell-mediated immunity was seen in patients with type-2 DM (1). Insulin-dependent DM is an organ-specific chronic autoimmune disease resulting from T-cell-mediated destruction of pancreatic cells. It is generally accepted that proinflammatory cytokines secreted by mononuclear cells are responsible for pancreatic B cell destruction in animal models of autoimmune type-1 DM (2). In type-1 DM, the infiltration of the islets of the pancreas with mononuclear cells leads to the damage of B cells and causes insulin deficiency. Several studies have shown that markers of inflammation and pro-inflammatory cytokines such as interleukin (IL)-10 and IL-13 associate with the metabolic syndrome, dyslipidemia, type-2 DM and type-1 DM (3-5).

Interleukin-10 is produced by activated Th2 lymphocytes, mast cells, monocytes, B lymphocytes, eosinophils and keratinocytes (6). IL-10 is also produced by activated human natural killer cells and has been identified as a vital modulator of abundant production of inflammatory cytokines (7). It acts to inhibit proinflammatory cytokines, growth factors and chemokine production by mononuclear phagocytes (6,8). It down-regulates various inflammatory events. Monocytes pretreated with IL-10 fail to induce antigen-specific T-cell proliferation (6). In an IL-10 transgenic model, IL-10 enhanced the accumulation of CD8 T cells in the pancreas leading to an earlier onset of diabetes in nonobese diabetic mice (9).

Interleukin-13 is a Th2 cytokine with many of the same activities as IL-4 because they share some of the same surface receptors. IL-13 can suppress the cytotoxic functions of monocytes-macrophages and inhibit the production of macrophage-derived proinflammatory cytokines (10). The data on IL-13 are variable. Kretowski et al.’s (2) findings suggest that IL-13 alterations could play an important role both in the pathogenesis and in the prevention of type-1 DM. It was suggested the IL-13 levels correlate with the activity of the autoimmune process (11). Peripheral blood mononuclear cells (PBMCs) from patients with type-2 DM show reduced proliferative response to phytohemagglutinin (PHA) and other mitogens (1).

The aim of this study was to examine the in vitro production of IL-10 and IL-13 in cultures of peripheral lymphocytes obtained from patients with type-2 DM, type-1 DM and healthy controls.

Materials and Methods

All patients with DM were diagnosed in accordance with the Report of the Expert Committee on the Diagnosis and Classification of Diabetes (12). Twenty patients (10 females, 10 males) with type-2 DM (11 patients with poor metabolic control, HbA1C>7.5%, median age: 57 years, and median disease duration:10 years; 9 patients with good metabolic control, HbA1C<7.5%, median age: 63 years, and median disease duration: 11 years), 10 patients (5 females, 5 males) with type-1 DM (median age: 24 years and median disease duration: 11 years) and the control group were enrolled in the study. The control group included 10 healthy subjects (5 females and 5 males, median age: 57 years) matched for age, sex and body mass index (BMI) with type-2 DM patients, but with no clinical or laboratory signs of any disease. The demographic and descriptive details of all patients and controls are shown in Table 1.

<table>
<thead>
<tr>
<th></th>
<th>Patients(^a) (n = 11)</th>
<th>Patients(^b) (n = 9)</th>
<th>Patients(^c) (n = 10)</th>
<th>Controls (n = 10)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (year)</td>
<td>57 (27)</td>
<td>63 (31)</td>
<td>24 (15)</td>
<td>57 (33)</td>
</tr>
<tr>
<td>Sex (M/F)</td>
<td>6/5</td>
<td>4/5</td>
<td>7/4</td>
<td>5/5</td>
</tr>
<tr>
<td>BMI (kg/m(^2))</td>
<td>28 (10)</td>
<td>28 (11)</td>
<td>26 (13)</td>
<td>26 (11)</td>
</tr>
<tr>
<td>Disease duration</td>
<td>10 (12)</td>
<td>11 (12)</td>
<td>11 (14)</td>
<td>-</td>
</tr>
</tbody>
</table>


\(^a\) Patients with NIDDM (poor metabolic control).
\(^b\) Patients with NIDDM (good metabolic control).
\(^c\) Patients with IDDM.
Twelve patients with type-2 DM had hypertension and had taken antihypertensive drugs. Ten patients had hyperlipidemia. All of the type-2 DM patients had taken oral antidiabetic drugs (OAD). All of the type-1 DM patients had taken insulin.

Exclusion criteria included abnormal laboratory test results such as the levels of serum urea, creatinine, liver enzymes, erythrocyte sedimentation rate, C-reactive protein, white and red blood cell counts, and thrombocytes. Cases were also excluded from the trial if they had any autoimmune or infectious diseases. Informed consent was obtained from all patients and healthy controls. The study protocol was approved by the Ethics Review Committee of the Medical School of Gulhane.

**Cell preparations and cultures:** PBMCs were isolated from 20 ml of heparinized venous blood by centrifugation on Ficoll-Hypaque Separating Solution (Seromed®, Biochrom KG Berlin, Germany). Then, PBMCs were washed twice in sterile phosphate buffered saline (PBS) and resuspended in RPMI-1640 medium (Gibco, Invitrogen Corporation, Scotland, UK) supplemented with 10% heat inactivated newborn calf serum (NCS) (Gibco, Invitrogen Corporation, Scotland, UK). These cell suspensions were incubated in culture flasks (Nunc Brand Products, Roskilde, Denmark) at 37 °C in a humidified 5% CO₂ atmosphere for 2 h. Adherent cells were discarded, but nonadherent cells were collected from upper layers of the cell suspensions, washed twice in PBS, and resuspended in RPMI-1640 medium supplemented with 10% NCS, 100 IU/ml penicillin, 10 m/ml streptomycin (Sigma Chemical Co. St Louis, MO, USA), and 2 mM/ml L-glutamine (Gibco, Invitrogen Corporation, Scotland, UK) for cell culture. Viability in the cell preparations was determined by acridine-orange staining method, and found as more than 95%.

Peripheral blood mononuclear cells were cultured at a concentration of 1x10⁶/ml in RPMI-1640 medium supplemented with 10% NCS, 100 IU/ml penicillin, 100 m/ml streptomycin, and 2 mM/ml L-glutamine. Cells were cultured in 24 well plates (Nunc Brand Products, Roskilde, Denmark) in the absence or presence of phytohemagglutinin-M (PHA-M, Gibco, Invitrogen Corporation, Scotland, UK) at 37 °C in a humidified 5% CO₂ atmosphere for 36 h. PHA-M was used at final concentrations of 10 µg/ml and 20 µg/ml in the cultures. Culture supernatants were collected at the end of incubation, then centrifuged at 550g for 5 min and stored at −70 °C until IL-10 and IL-13 analyses.

**Measurement of IL-10 and IL-13:** IL-10 and IL-13 in culture supernatants were measured by the enzyme immunoassay method using commercially available ELISA kits (Cytimmune Sciences Inc, College Park, MD, USA). According to kit prescription, sensitivity was 1.6 pg/ml and 7.6 pg/ml, intraassay variation ±8.4% for both, and interassay variation ± 10.2% and ± 11.9% for IL-10 and IL-13, respectively.

**Statistical analysis:** Statistical analyses were performed using Mann-Whitney U test, Wilcoxon signed rank tests, and Spearman’s rank correlation in the SPSS for Windows 11.5 statistical software (SPSS Inc. Chicago, IL, USA). A value of p ≤ 0.05 was considered to represent a significant difference between analyzed data.

**Results**

The mean IL-10 and IL-13 levels in culture supernatants incubated with 10 µg/ml PHA were 199 ± 75 pg/ml and 113 ± 31 pg/ml in type-2 DM patients, 209 ± 38 pg/ml and 101 ± 15 pg/ml in type-1 DM patients, and 228 ± 38 pg/ml and 101 ± 15 pg/ml in healthy controls, respectively. The mean IL-10 and IL-13 levels in culture supernatants incubated with 20 µg/ml PHA were 286 ± 38 pg/ml and 185 ± 67 pg/ml in type-2 DM patients, 306 ± 44 pg/ml and 182 ± 24 pg/ml in type-1 DM patients, and 448 ± 166 pg/ml and 225 ± 78 pg/ml in the healthy controls, respectively (Table 2).

In all groups, IL-10 and IL-13 levels were significantly high in culture supernatants incubated with 20 µg/ml PHA compared to levels in culture supernatants incubated with 10 µg/ml PHA (P = 0.002 and P = 0.001 in the patients, P = 0.005 and P = 0.005 in the controls, respectively).

IL-10 and IL-13 levels in all patients and healthy controls were similar in culture supernatants incubated with 10 µg/ml PHA (P < 0.05). IL-13 levels in controls were higher than in the patients in culture supernatants incubated with 20 µg/ml PHA, but without statistical significance. IL-10 levels in type-2 DM patients were
significantly lower than in the control group in culture supernatants incubated with 20 µg/ml PHA (P = 0.001 for patients with poor metabolic control and P = 0.048 for patients with good metabolic control, respectively) (Figure 1.). When we compared IL-10 and IL-13 levels in patients with good and poor metabolic control, there was no statistically significant difference.

**Discussion**

Impaired cell-mediated immunity in diabetics is being reported with increasing frequency. The results of several studies have suggested important roles of cytokines in the pathogenesis of both type-2 DM and type-1 DM (2,5,13,14). It has been reported that long-standing metabolic abnormalities might affect the monocyte function (13). We detected that good and poor metabolic control did not affect IL-10 and IL-13 production by peripheral blood lymphocytes in patients with type-2 DM and type-1 DM. Ragab et al. (15) found no differences between normal individuals and diabetic individuals with good metabolic control when they compared the responses of peripheral blood lymphocytes to either PHA. We also found similar results in our study.

The infiltration of the islets of the pancreas with mononuclear cells leads to the damage of B cells and causes insulin deficiency. In an IL-10 transgenic model, IL-10 enhanced the accumulation of CD8+ T cells in the pancreas leading to an earlier onset of diabetes in nonobese diabetic mice (16). IL-10 production in vitro by lymphocytes from high-risk of type-1 DM patients was spontaneously enhanced both in nonstimulated and insulin-stimulated cultures (14). PHA increased interferon (IFN) gamma, IL-6 and IL-10 production in all groups (14). It is postulated that IL-10 favors the activation of the antigen-presenting cell population and downregulates the autoreactive effector cells at the site of the immune reaction (9). Tchorzewski (14) observed the

---

**Table 2. Comparisons of IL-10 and IL-13 production by PBMCs and some metabolic parameters in patients with NIDDM and IDDM and healthy controls.**

<table>
<thead>
<tr>
<th></th>
<th>Patients&lt;sup&gt;‡&lt;/sup&gt; (n = 10)</th>
<th>Patients&lt;sup&gt;†&lt;/sup&gt; (n = 9)</th>
<th>Patients&lt;sup&gt;¶&lt;/sup&gt; (n = 11)</th>
<th>Controls (n = 10)</th>
<th>χ²</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-10* (pg/ml)</td>
<td>203 (317)</td>
<td>185 (196)</td>
<td>235 (149)</td>
<td>221 (120)</td>
<td>2.383</td>
<td>0.497</td>
</tr>
<tr>
<td>IL-10** (pg/ml)</td>
<td>268 (241)</td>
<td>279 (578)</td>
<td>321 (627)</td>
<td>434 (568)</td>
<td>8.720</td>
<td>0.033</td>
</tr>
<tr>
<td>IL-13* (pg/ml)</td>
<td>108 (111)</td>
<td>96 (92)</td>
<td>105 (109)</td>
<td>106 (133)</td>
<td>1.001</td>
<td>0.801</td>
</tr>
<tr>
<td>IL-13** (pg/ml)</td>
<td>200 (173)</td>
<td>187 (238)</td>
<td>208 (221)</td>
<td>219 (215)</td>
<td>2.229</td>
<td>0.526</td>
</tr>
<tr>
<td>HbA1C (%)</td>
<td>8.7 (1.9)&lt;sup&gt;a&lt;/sup&gt;</td>
<td>6.8 (1.4)&lt;sup&gt;b&lt;/sup&gt;</td>
<td>7.0 (3.8)&lt;sup&gt;c&lt;/sup&gt;</td>
<td>6.1 (2.1)&lt;sup&gt;d&lt;/sup&gt;</td>
<td>23.195</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Glucose (mg/dl)</td>
<td>160 (111)&lt;sup&gt;e&lt;/sup&gt;</td>
<td>128 (103)&lt;sup&gt;f&lt;/sup&gt;</td>
<td>121 (120)</td>
<td>101 (44)&lt;sup&gt;g&lt;/sup&gt;</td>
<td>13.669</td>
<td>0.003</td>
</tr>
<tr>
<td>Cholesterol (mg/dl)</td>
<td>208 (109)</td>
<td>186 (72)</td>
<td>199 (75)</td>
<td>192 (93)</td>
<td>1.934</td>
<td>0.586</td>
</tr>
<tr>
<td>Triglyceride (mg/dl)</td>
<td>208 (262)</td>
<td>163 (103)</td>
<td>163 (175)</td>
<td>166 (178)</td>
<td>1.635</td>
<td>0.647</td>
</tr>
<tr>
<td>HDL (mg/dl)</td>
<td>42 (22)</td>
<td>46 (23)</td>
<td>39 (21)</td>
<td>45 (31)</td>
<td>1.564</td>
<td>0.668</td>
</tr>
</tbody>
</table>

Descriptive statistics are presented as median (range). For multiple groups, we used Kruskal-Wallis test. The differences between two groups were evaluated by Wilcoxon signed rank tests or Mann-Whitney U test, where appropriate.


* IL-10 and IL-13 levels in culture incubated with 10 mg/ml phytoagglutinin (PHA).
** IL-10 and IL-13 levels in cultures incubated with 20 mg/ml PHA.
<sup>¶</sup> Patients with NIDDM (poor metabolic control).
<sup>†</sup> Patients with NIDDM (good metabolic control).
<sup>‡</sup> Patients with IDDM.

P < 0.05 for * vs **. in all study groups; P < 0.001 for a vs b, a vs d.
P = 0.004 a vs c, P = 0.001 e vs g and P = 0.021 f vs g.
increase in IL-10 production by lymphocytes of DM patients with high risk, which may implicate the autocrine role of this cytokine in the pathogenesis of islet cells destruction and type-1 diabetes development. In our study, in type-1 DM, type-2 DM and healthy controls, IL-10 levels were similar in culture supernatants incubated with 10 µg/ml PHA, but IL-10 production was significantly higher in culture supernatants incubated with 20 µg/ml in healthy controls compared to levels in the patients with type-2 DM and type-1 DM.

It has previously been shown that IL-13, a recently discovered cytokine, can suppress the cytotoxic function of monocytes/macrophages, inhibit the production of macrophage-derived proinflammatory cytokines, and have a modulatory effect on the B lymphocytes function (17-20). The main anti-inflammatory properties of IL-13 are associated with the downregulation of the secretion of IL-1beta and tumor necrosis factor (TNF) alpha by macrophages/macrophages (2,17). The alteration in IL-13 is also believed to contribute to the development of autoimmune diseases (21-23). In our study, IL-13 production was similar with 10 and 20 µg/ml PHA-stimulated peripheral blood lymphocytes in all groups. We also studied IL-10 and IL-13 production of PBMCs in patients with type-2 DM.

It has been reported that PBMCs from patients with type-2 DM show reduced proliferative response to PHA and other mitogens (1,24). In our study, the proliferative response of IL-10 with 10 µg/ml PHA remained unchanged, but the response was also not enough with 20 µg/ml PHA.

According to our findings, the alterations in IL-13 did not play an important role in pathogenesis of type-2 DM and type-1 DM. Furthermore, the deficiency in IL-10 production from PBMCs with high dosages of PHA suggests that IL-10 may play an important role in the pathogenesis of type-2 DM.

References


Tchorzewski H, Glowacka E, Banasik M, Lewkowicz P, Szalapska-Zawodniak M. Activated T lymphocytes from patients with high risk of type I diabetes mellitus have different ability to produce interferon-gamma, interleukin-6 and interleukin-10 and undergo anti-CD95 induced apoptosis after insulin stimulation. Immunol Lett 2001; 75: 225-34.


Ohno Y, Oaki N, Nishimura A. In vitro production of IL-1, IL-6, and TNF alpha in insulin-dependent diabetes mellitus. J Clin Endocrinol Metab 1993; 77: 1072-7.