Expression and Function of Inducible Costimulator on Peripheral Blood CD4⁺ T Cells in Behçet’s Patients with Uveitis: A New Activity Marker?

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PURPOSE. Inducible costimulator (ICOS) is an important co-stimulatory molecule involved in T-cell activation. In this study, the role of ICOS in the pathogenesis of uveitis in Behçet’s disease (BD) was investigated.

METHODS. Peripheral blood mononuclear cells (PBMCs) were obtained from BD patients with uveitis in the active or remission phase and in healthy subjects. Total RNA was isolated from PMBCs, and mRNA expression was analyzed on an oligonucleotide microarray. ICOS expression on CD4⁺ T cells was determined by flow cytometry, and the functional costimulatory effect of ICOS/B7RP-1 interaction was assessed on stimulation with concanavalin A (conA) or IRBP in the presence or absence of anti-ICOS mAb.

RESULTS. As the result of microarray analysis, ICOS in PBMCs showed the greatest difference in expression in BD patients with uveitis compared with healthy control subjects. ICOS expression on CD4⁺ T cells in BD patients with uveitis was significantly higher than that in healthy individuals, both before and after conA stimulation. Among the BD patients, ICOS expression on CD4⁺ T cells was significantly higher in those with active uveitis than in those with remitted uveitis. Blockade of ICOS/B7-related protein-1 (B7RP-1) interaction by anti-ICOS mAb significantly decreased IFN-γ, IL-17, and TNF-α production by PBMCs when stimulated with conA or IRBP in BD with active uveitis.

CONCLUSIONS. High ICOS expression in BD patients with uveitis contributed to the upregulation of IFN-γ, IL-17, and TNF-α production, suggesting that abnormal ICOS costimulation may play an immunopathologic role in the pathogenesis of uveitis in BD. (Invest Ophthalmol Vis Sci. 2010;51:5099–5104) DOI: 10.1167/iovs.10-5286

Endogenous uveitis such as Behçet’s disease (BD), Vogt-Koyanagi-Harada syndrome, sympathetic ophthalmia, birdshot retinochoroidopathy, and sarcoidosis is a potentially blinding disease in humans and is responsible for 10% to 15% of the acquired blindness in Japan. Although endogenous uveitis covers a spectrum of clinical entities, all forms are believed to share immunohistologic similarities characterized by the infiltration of mainly T cells. Behçet’s disease (BD) is a systemic inflammatory disease characterized by oral and genital ulcers as well as ocular, cutaneous, arthritic, vascular, and neurologic lesions. Many factors have been implicated in the development of inflammation and severity of disease. Although genetic factors, infections, and autoimmune responses have been implicated in BD, much about its pathogenesis remains uncertain.

An increasing number of reports indicate that aberrant cellular immunity, such as pathogenic CD4⁺ T-cell-mediated autoimmunity via the Th1/Th17 pathway, plays a key role in the pathophysiological process in BD with uveitis. Although the mechanisms of ocular inflammation in BD remain largely unknown.

During recent years, the understanding of immunologic mechanisms involved in uveitis has advanced greatly through investigations of experimental autoimmune uveoretinitis (EAU), an animal model of human uveitis that can be induced by immunization of susceptible animals with interphotoreceptor retinoid-binding protein (IRBP) or with an eye-specific retinal antigen or by adoptive transfer of CD4⁺ T cells specific for retinal antigens. EAU resembles certain human uveitic conditions in various aspects and the major mediator is the CD4⁺ T cell.

It is well known that T-cell activation requires the integration of at least two distinct signals. Signal one results from antigen-specific T-cell receptor engagement by MHC class 2-bound peptide presented by antigen-presenting cells (APCs). Signal two arises from engagement of costimulatory molecules expressed by antigen-presenting cells and T cells. Although CD28 regulation has substantial effects on immunity, its function appears to reside predominantly in the control of primary, but not secondary, immune responses in various autoimmune diseases. The CD28 homolog inductive costimulator (ICOS) has recently been identified as a novel member of the CD28 costimulator family and is expressed by activated T cells in both humans and mice. The ICOS ligand, B7-related protein (B7RP-1), also known as B7 homologous protein (B7h), is constitutively expressed by dendritic cells, macrophages, and B cells, and the degree of expression is increased by IFN-γ, TNF-α, and IL-17. Discovery of the ICOS/B7RP-1 pathway has led to considerable interest in the physiologic and pathologic functions of this pathway. A series of reports support the concept that ICOS is critical in the regulation of Th2 responses. However, recent murine studies indicate that ICOS also regulates Th1- and Th17-mediated diseases, such as experimental allergic encephalomyelitis.
To identify new genes that may cause or contribute to the disease process of ocular BD, we used cDNA microarrays that provide the expression profile of more than 54,000 genes, some of which are immune-related, whereas others are involved in the cell cycle, cell growth, intracellular signaling, cellular adhesion, and transport, and we compared the expression profiles of healthy individuals and patients. One of the genes found to be upregulated in patients with ocular BD is ICOS, an activation marker expressed on activated T cells that binds B7RP-1-expressing monocytes. The engagement of ICOS with B7RP-1 along with an appropriate antigen provide a positive signal that promotes T-cell differentiation, cytokine secretion, and effector function in the absence of CD28.35 This process has been demonstrated in vivo in a murine EAU model, and we have shown that ICOS protein expressed on infiltrating CD4+ T cells and B7RP-1 expressed on infiltrating APCs are upregulated directly after disease onset.36 Therefore, we speculated that in active BD, pathogenic CD4+ T cells that infiltrate the eye express ICOS in the inflamed eye and would be an appropriate target for the treatment of human ocular BD. Moreover, we have demonstrated that blockade of the ICOS/B7RP-1 costimulatory pathway inhibits ocular inflammation in the effector phase of murine EAU.36 Therefore, to investigate the role of ICOS in the pathogenesis of ocular BD, we assessed its expression on peripheral blood CD4+ T cells and functional roles in patients with ocular BD. The results suggest that upregulation of ICOS is associated with the pathogenesis of uveitis in BD.

**Materials and Methods**

**PBMC Samples**

Thirty-five patients with BD (27 men and 8 women, mean age: 37.8 ± 11.1 years) were enrolled in the study (Table 1). All patients fulfilled the diagnostic criteria for ocular BD proposed by the International Study Group for BD. Other etiologies were excluded after a complete ocular and systemic work-up. The samples from the patients were classified as active or inactive intraocular inflammation, according to the presence or absence of active ocular inflammation on the day of sample collection, as previously described.36 Briefly, uveitis in the active phase (active intraocular inflammation) was defined by the presence of any of the following ocular inflammation findings observed by ophthalmoscopy: three or more infiltrating cells in the anterior chamber, worsening vitreous opacity, and retinal exudates. Uveitis in the remission phase (inactive ocular inflammation) was defined as the absence of any of the signs of active disease. Seventeen patients had recurrent uveitis, and active intraocular inflammation was observed in blood samples obtained at the Ophthalmic Center of Tokyo Medical University Hospital. Eighteen patients had inactive ocular BD. Twenty-one healthy individuals (16 men and 5 women, mean age: 42.7 ± 16.6 years) served as control subjects. Patients who had used oral or intra-venous corticosteroids or immunosuppressive agents such as cyclosporine were excluded. All subjects gave informed consent before sample acquisition. The study was approved by the ethics committee of the Tokyo Medical University and complied with the guidelines in the Declaration of Helsinki for research involving human subjects.

Heparinized blood samples were obtained from patients and healthy individuals, and PBMCs were isolated within 2 hours by density gradient centrifugation. The PBMCs were washed twice and resuspended at 1 × 10^6 cells/mL in complete medium (RPMI 1640 supplemented with 10% fetal calf serum, 1 mM L-glutamine, 100 U/mL penicillin, and 100 µg/mL streptomycin). Cell suspension was dispensed in 24-well plates (Falcon; BD Biosciences, Mountain View, CA) at 1 mL/well and incubated with or without concanavalin A (ConA; 10 µg/mL) for 12 hours.

**Analysis of Gene Expression with cDNA Array**

Gene expression profile of human PBMCs was analyzed by microarray, as described previously.77 Briefly, total RNA was isolated from pooled PBMCs obtained from seven patients with BD and active uveitis by an extraction method (Isogen Nippon Gene, Tokyo, Japan), and portions (2 µg) of the preparation were subjected to amplification of mRNA with T7 RNA polymerase. Biotin-labeled cRNA (10 µg) synthesized from the amplified RNA was subjected to hybridization with the Human Whole Genome Bioarray chip (Amersham Biosciences, Piscataway, NJ) that contains oligonucleotides corresponding to a total of approximately 55,585 human genes. Detection and digitization of hybridization signals were performed (G2565; Agilent Technologies, San Diego, CA) and analyzed (CodeLink Expression Analysis software version 4.0; Amersham Biosciences). The microarray data of various costimulatory molecules are summarized in Table 2. Expression levels of each gene were automatically normalized to the median of all spot intensities.

**Reagents**

FITC-conjugated anti-human CD4 (L3T4) was purchased from eBioscience (San Diego, CA), and anti-human ICOS (DX29) from BD Pharmingen (San Diego, CA). ConA was obtained from Vector Laboratories (Burlingame, CA).

**Preparation of IRBP**

Fresh swine IRBP was purified according to the method described by Fukai et al.38 with some modification. Briefly, frozen retinas were thawed in 0.03 M disodium monopotassium phosphate buffer (PB), after which the retinas were removed by centrifugation at 10,000g for 30 minutes. Saturated ammonium sulfate (pH 7.2) was added to the supernatant until 50% saturation was reached, and the mixture was left overnight at 4°C. After centrifugation, the precipitate was dissolved in PB and used as the crude antigen preparation. IRBP was isolated from the preparation according to the method of Redmond et al.39 Briefly, the preparation was purified successively by conA-Sepharose affinity chromatography (HPLC) and ion-exchange HPLC. IRBP in the eluted

**Table 1. Patient Characteristics**

<table>
<thead>
<tr>
<th>Disease Activity</th>
<th>Subjects (n)</th>
<th>Age (y)</th>
<th>Male/Female</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ocular Behçet</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Active</td>
<td>17</td>
<td>36.5 ± 10.1</td>
<td>13/4</td>
</tr>
<tr>
<td>Inactive</td>
<td>18</td>
<td>45.7 ± 15.0</td>
<td>14/4</td>
</tr>
<tr>
<td>Healthy individuals</td>
<td>21</td>
<td>42.7 ± 16.6</td>
<td>16/5</td>
</tr>
</tbody>
</table>

**Table 2. Genes Up- or Downregulated on PBMC with or without ConA in Ocular Behçet Patients**

<table>
<thead>
<tr>
<th>Genes of Costimulatory Molecules</th>
<th>Intensity Difference</th>
<th>GenBank Accession Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>ICOS</td>
<td>1.125</td>
<td>NM_005214</td>
</tr>
<tr>
<td>CD40L</td>
<td>0.274</td>
<td>NM_006139</td>
</tr>
<tr>
<td>4–1BB</td>
<td>0.231</td>
<td>NM_012092</td>
</tr>
<tr>
<td>CTLA4</td>
<td>0.110</td>
<td>NM_00074</td>
</tr>
<tr>
<td>CD28</td>
<td>0.167</td>
<td>NM_001242</td>
</tr>
<tr>
<td>CD27</td>
<td>1.382</td>
<td>NM_019561</td>
</tr>
<tr>
<td>PD-1</td>
<td>3.226</td>
<td>NM_02124</td>
</tr>
</tbody>
</table>

CTLA4, cytotoxic T-lymphocyte-associated antigen 4; PD-1, programmed death-1.
fractions was identified by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and binding of \(^1\)H]retinol. Purification of IRBP to homogeneity, as assessed by SDS-PAGE and silver staining, was accomplished by size-exclusion HPLC in the presence of glycerol.

### Immunofluorescence and Flow Cytometry

The results of ICOS gene expression obtained from microarray were confirmed by flow cytometry. PBMCs were isolated by density gradient centrifugation of heparinized blood samples obtained from patients and healthy individuals. Each PBMC sample was divided into two aliquots: one for direct flow cytometry analysis of ICOS expression and the other for flow cytometry analysis after activation with conA at 10 \(\mu\)g/mL for 12 hours. ICOS analysis was performed by using the following procedures: A cocktail of FITC-conjugated anti-CD4 and PE-conjugated anti-ICOS was added to the PBMCs. After the cells were washed with PBS, the stained cells (live-gated based on forward- and side-scatter profiles and propidium iodide exclusion) were passed through a flow cytometer (FACSCalibur; BD Biosciences, San Jose, CA) and the results analyzed (CellQuest program; BD Biosciences). ICOS expression on human CD4\(^+\) T cells was calculated by gating for the CD4\(^+\) T cell population, and 10,000 cells were analyzed in each experiment.

### Cytokine Assay

Purified PBMCs (5 \(\times\) 10\(^5\)/well) from BD patients with active uveitis were cultured in 96-well microtiter plates, with or without stimulation with conA (3 \(\mu\)g/mL) or IRBP (10 \(\mu\)g/mL) plus anti-ICOS mAb (10 \(\mu\)g/mL) or control IgG at 37°C for 48 hours. The unstimulated PBMCs served as negative control samples. After stimulation, cell-free supernatants were collected at 48 hours and assayed for IFN-\(\gamma\), TNF-\(\alpha\), IL-2, and IL-6 by cytometric bead array (CBA) kits (BD PharMingen) and IL-17 ELISA was 7.8 pg/mL.

### Statistical Analysis

Data were analyzed (JMP 5, SAS Institute, Inc., Cary, NC) and the results expressed as the mean \(\pm\) SD. Statistical analysis was performed by using a paired or unpaired t-test. \(P < 0.05\) was considered significant.

### RESULTS

#### Analysis of Microarray Results in BD Patients with Active Uveitis

To identify the genes involved in the pathogenesis of ocular BD, we compared phosphorescent images of the arrays hybridized with cDNA probes generated from RNA preparations from BD patients with active uveitis and healthy individuals. The array membrane that we used contained 360 positive bacterial super- natants were collected at 48 hours and assayed for IFN-\(\gamma\), TNF-\(\alpha\), IL-2, and IL-6 by cytometric bead array (CBA) kits (BD PharMingen) and IL-17 ELISA (Human IL-17 Quantikine ELISA kit; R&D Systems, Minneapolis, MN), according to the manufacturers' instructions. The minimum levels detected by CBA kits in either the culture supernatant or serum/plasma were 2.6 pg/mL for IL-2, 3.0 pg/mL for IL-6, 7.1 pg/mL for IFN-\(\gamma\), and 2.8 pg/mL for TNF-\(\alpha\); the minimum level in the IL-17 ELISA was 7.8 pg/mL.

In a preliminary study, serial concentrations of conA (1, 3, 5, and 10 \(\mu\)g/mL) or IRBP (1, 5, and 10 \(\mu\)g/mL) were used in the assays. The results showed that optimal concentrations of cytokines were obtained at a concentration of 3 \(\mu\)g/mL for conA and 10 \(\mu\)g/mL for IRBP, as previously described.\(^{39}\) These concentrations were therefore used in subsequent experiments.

#### Immunofluorescence and Flow Cytometry

Next, we compared ICOS expression on CD4\(^+\) T cells in BD patients with active or inactive uveitis. ICOS expression (MFI) on CD4\(^+\) T cells; Fig. 1A, positive rates of ICOS\(^+\)CD4\(^+\) T cells; Fig. 2B) on freshly isolated CD4\(^+\) T cells from BD patients with active uveitis was significantly upregulated compared with that in those with inactive uveitis. After 12 hours of incubation, ICOS expression (MFI and percentage of CD4\(^+\) T cells expressing ICOS) was similar in BD patients with active or inactive uveitis in the presence or absence of conA. These results suggest that higher ICOS expression on CD4\(^+\) T cells in BD patients with uveitis may be involved in the pathogenesis of ocular BD.

### Cytokine Production Stimulated with ConA and Uveitogenic Antigen

To determine whether ICOS has functional costimulatory activity, we first measured the amounts of pathogenic Th1 and Th17 cytokines (IFN-\(\gamma\), IL-17, TNF-\(\alpha\), IL-6, and IL-2) produced by non-specific stimulation (conA) of PBMCs obtained from BD patients with active uveitis. When ICOS stimulation was blocked with anti-ICOS mAb, the amounts of IFN-\(\gamma\), IL-17, and TNF-\(\alpha\) (Figs. 3A–C) produced by PBMCs were both significantly reduced.

Since PBMCs from BD patients are known to be sensitized to the two most uveitogenic retina-specific antigens, S-antigen and IRBP,\(^1\) we then examined the amounts of pathogenic Th1 and Th17 cytokines produced by stimulation with swine IRBP protein as the specific antigen. Figures 3F–I show the results obtained from five or six representative BD patients with active uveitis. When ICOS stimulation was blocked with anti-ICOS mAb, the amounts of IFN-\(\gamma\) and IL-17 production by IRBP stimulation were reduced in a manner similar to that caused by nonspecific stimulation, except TNF-\(\alpha\). Although no significant production of IL-6 and -2 was observed, both conA- and IRBP-induced productions of IL-6 were apparently reduced, at least in part, by blockade of ICOS costimulation. Thus, these results suggest that signaling through ICOS costimulation is involved in the production of pathogenic Th1 and Th17 cytokines by PBMCs when stimulated with nonspecific or retina-specific antigens.

### DISCUSSION

The precise pathogenesis of uveitis associated with BD is unknown. Accumulated data obtained from animal models such as EAU suggest that Th1 and Th17 cells have major roles...
Studies on autoantigen-induced cellular immune responses in BD patients with uveitis mainly used assays that measure the lymphoproliferative response and cytokine productions. Previous studies showed that Th1 and Th17 cytokine production by polyclonally stimulated PBMCs and activated T cells was significantly elevated in BD patients with active uveitis compared with patients without active uveitis or controls. We therefore further examined the influence on IFN-γ, IL-17, TNF-α, IL-6, and IL-2 production by PBMCs after nonspecific or antigen-specific stimulation when we blocked ICOS costimulation with anti-ICOS mAb. All these findings suggest that the ICOS costimulatory pathway is associated with active uveitis in BD patients and that this pathway has a positive costimulatory role in enhancing the production of Th1 and Th17 cytokines, which are important mediators in the pathogenesis of active uveitis in BD patients. Studies on the manipulation of this pathway may validate its role and also provide a new strategy for the treatment of this disease.

We have previously shown that blockade of the ICOS costimulatory pathway has an ameliorating effect during the effector phase of EAU, by suppressing the expansion and effector function of pathogenic Th1 cells. Considering that blocking Abs to IL-17 ameliorates murine EAU, our data suggest that anti-ICOS mAb may mitigate ocular inflammation.
and systemic autoimmune responses in BD patients with active uveitis by inhibiting Th1 and Th17 cytokine production.

In this study, ICOS was markedly upregulated in CD4+ T cells from BD patients with active uveitis. Both the proportion of ICOS-expressing cells and percentage of ICOS expression in circulating peripheral CD4+ T cells from BD patients with active uveitis were significantly higher in BD patients with active uveitis than in those with inactive uveitis. ICOS expression on CD4+ T cells from BD patients with uveitis may play an immunopathologic role in ocular inflammation. It is important to evaluate the activity of ocular BD. However, currently there is no reliable laboratory marker. C-reactive protein and erythrocyte sedimentation rate, which indicate inflammatory processes nonspecifically, probably cannot reflect the continuously changing magnitude of inflammatory activity in BD.48,49 On the other hand, the level of ICOS expression on CD4+ T cells from BD patients with active uveitis may be a sensitive index for evaluating the disease activity of ocular BD. Our results also suggest that ICOS expression on circulating peripheral CD4+ T cells would be a useful marker for monitoring the therapeutic effects of conventional treatment in patients with BD with active uveitis. In the case of systemic lupus erythematosus (SLE), expression of ICOS on peripheral T cells of patients with active disease was significantly reduced after treatment with high-dose corticosteroid, a strong inhibitor of lymphocyte activation, and the treatment almost completely abolishes the induction of ICOS with ICOS costimulation.50 Results in a recent study of T-cell-mediated autoimmune disease suggested that ICOS expression reflects the aggressiveness of autoreactive effector T cells.51 The higher ICOS expression in BD patients with active uveitis may be associated with a unique immunopathogenesis and the more recurrent or chronic nature of BD. Further studies with prospective follow-up are needed to address this possibility.

In conclusion, our data provide additional evidence of the potential utility of ICOS expression on CD4+ T cells as a marker of disease activity and as a promising therapeutic target for ocular BD via its inhibition of Th1 and Th17 cytokines. As the population studied was small and heterogeneous, further studies are needed to confirm the findings.

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References


