Dysfunction of Natural Killer T Cells in Patients with Active Mycobacterium tuberculosis Infection

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Natural killer T (NKT) cells are known to play a protective role in the immune responses of mice against a variety of infectious pathogens. However, little is known about the detailed information of NKT cells in patients with Mycobacterium tuberculosis infection. The aim of this study was to examine NKT cell levels and functions in patients with active M. tuberculosis infection, to investigate relationships between NKT cell levels and clinical parameters, and to determine the mechanism responsible for the poor response to α-galactosylceramide (α-GalCer). NKT cell levels were significantly lower in the peripheral blood of pulmonary tuberculosis and extrapulmonary tuberculosis patients, and the proliferative responses of NKT cells to α-GalCer were also lower in patients, whereas NKT cell levels and responses were comparable in latent tuberculosis infection subjects and healthy controls. Furthermore, this NKT cell deficiency was found to be correlated with serum C-reactive protein levels. In addition, the poor response to α-GalCer in M. tuberculosis-infected patients was found to be due to increased NKT cell apoptosis, reduced CD1d expression, and a defect in NKT cells. Notably, M. tuberculosis infection was associated with an elevated expression of the inhibitory programmed death-1 (PD-1) receptor on NKT cells, and blockade of PD-1 signaling enhanced the response to α-GalCer. This study shows that NKT cell levels and functions are reduced in M. tuberculosis-infected patients and these deficiencies were found to reflect the presence of active tuberculosis.

Tuberculosis (TB) remains one of the infectious causes of illness with high morbidity and mortality in humans. In 2009, ~9.4 million new cases of active infection were diagnosed and the disease was responsible for roughly 1.7 million deaths (48). Although 32% of the world’s population has been estimated to be currently infected by Mycobacterium tuberculosis, the majority are latently infected and remain healthy (1, 14). Nevertheless, approximately 5 to 10% of individuals infected with M. tuberculosis will eventually develop active disease. In addition, the risk of developing active tuberculosis is increased by a variety of immunocompromised conditions, such as HIV infection, diabetes, renal failure, immunosuppressant treatment, and malnutrition (29). Accordingly, host immune status might play an important role in the control of progression to active disease. Furthermore, emerging evidence indicates that cellular immunity to M. tuberculosis requires the coordinated responses of the innate and adaptive immune systems (23, 37).

Human natural killer T (NKT) cells are a distinct subset of T lymphocytes and are characterized by the restricted expression of an invariant Vα24-Jα18 T cell receptor (TCR) chain paired with the Vβ11 TCR chain. The pair of TCR chains recognizes glycolipid antigens, such as α-galactosylceramide (α-GalCer), presented by the major histocompatibility complex (MHC) class I-like molecule CD1d (20). In addition, NKT cells play a bridging role between innate and adaptive immunity due to their ability to activate a variety of innate and adaptive immune cells, such as dendritic cells, monocytes, natural killer (NK) cells, T cells, and B cells, by rapidly producing large amounts of Th1 and Th2 cytokines, like gamma interferon (IFN-γ) and interleukin-4 (IL-4) (45). Furthermore, NKT cells are known to serve as regulatory and/or effector cells and are implicated in a broad range of diseases, including autoimmunity, cancer, and infectious diseases (19, 47).

NKT cells are known to play a protective role in the immune responses of mice against a variety of infectious pathogens, including bacteria, viruses, and protozoan parasites, although some of these results are controversial (18, 28). More specifically, several studies have shown that murine or human NKT cells may mediate protection against M. tuberculosis (4, 17, 37, 38, 46). For example, it was demonstrated in a recent study that α-GalCer administration, alone or in combination with classic chemotherapy, can improve the clinical outcomes of M. tuberculosis infection in mice (38). It has also been shown that α-GalCer incorporation into bacillus Calmette-Guérin vaccine enhances the host immune response by modulating T cell priming via murine NKT cell activation (46). However, much less is known about human NKT cell levels and functions in patients with M. tuberculosis infection. A numerical deficiency of NKT cells has been found in pulmonary tuberculosis (PTB) patients (33, 40, 42), but the NKT cell level has not been determined in extrapulmonary tuberculosis (EPTB) patients. Furthermore, the response of NKT cells to α-GalCer has been studied.
TABLE 1 Clinical and laboratory characteristics of the TB patients and HCs

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>HC</th>
<th>LTBI</th>
<th>EPTB</th>
<th>PTB</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total no.</td>
<td>90</td>
<td>13</td>
<td>18</td>
<td>55</td>
</tr>
<tr>
<td>Sex (no. male/no. female)</td>
<td>46/44</td>
<td>6/7</td>
<td>9/9</td>
<td>35/20</td>
</tr>
<tr>
<td>Mean ± SD age (yr)</td>
<td>54.6 ± 13.0</td>
<td>47.9 ± 9.8</td>
<td>50.9 ± 18.1</td>
<td>55.2 ± 20.9</td>
</tr>
<tr>
<td>No. (%) with previous TB medication</td>
<td>ND⁴</td>
<td>ND</td>
<td>17 (94.4)</td>
<td>44 (80)</td>
</tr>
<tr>
<td>No. (%) with positive sputum AFBb smear</td>
<td>ND</td>
<td>ND</td>
<td>0</td>
<td>35 (63.6)</td>
</tr>
<tr>
<td>No. (%) with cavity on chest radiograph</td>
<td>ND</td>
<td>ND</td>
<td>0</td>
<td>19 (34.5)</td>
</tr>
</tbody>
</table>

No. (%) with the following extent of pulmonary disease:
- Stage 1 (segmental disease): ND ND 0 4 (7.3)
- Stage 2 (lobar disease): ND ND 0 13 (23.6)
- Stage 3 (bi/trilobar disease): ND ND 0 10 (18.2)
- Stage 4 (bilateral disease): ND ND 0 28 (50.9)

No. (%) with the following sites of EPTB:
- Lymph node: ND ND 6 (33.3) 0 (0)
- Bone and joint: ND ND 8 (33.3) 4 (7.3)
- Pericardium: ND ND 1 (5.6) 1 (1.8)
- Peritoneum: ND ND 1 (5.6) 2 (3.6)
- Pharynx: ND ND 1 (5.6) 0 (0)
- Pleura: ND ND 1 (5.6) 0 (0)
- Mean ± SD hemoglobin concn (mg/dl) | 14.0 ± 1.2 | 13.8 ± 1.4 | 11.6 ± 2.3 | 12.0 ± 1.7 |
- Mean ± SD leukocyte count (cells/μl) | 5,592 ± 1,457 | 6,885 ± 1,485 | 5,758 ± 2,046 | 7,669 ± 3,014 |
- Mean ± SD lymphocyte count (cells/μl) | 2,041 ± 719 | 2,102 ± 526 | 1,422 ± 590 | 1,389 ± 668 |
- Mean ± SD platelet count (10⁴ cells/μl) | 243 ± 58 | 228 ± 43 | 242 ± 96 | 297 ± 124 |
- Mean ± SD CRPc concn (mg/dl) | ND | ND | 2.05 ± 2.38 | 5.21 ± 5.42 |

⁴ ND, not done.

b AFB, acid-fast bacilli.

C, reactive protein.

The study was approved by the Institutional Review Board of Chonnam National University Hospital, and written informed consent was obtained from all participants.

MAbS and flow cytometry. The following monoclonal antibodies (MAbs) and reagents were used in this study: fluorescein isothiocyanate (FITC)-conjugated MAbs against CD3, CD4, and CD14; FITC-conjugated annexin V; phycoerythrin (PE)-conjugated MAbs against NKT cell TCR (clone 6B11) and CD1d; peridinin chlorophyll-a protein (PerCP)-conjugated anti-CD3 MAbs; allophycocyanin (APC)-conjugated MAbs against CD3 and CD8; and 7-aminoactinomycin D (AAD) (all were from BD Biosciences, San Diego, CA) and PE-conjugated anti-human programmed death-1 (PD-1) MAb (eBiosciences, San Diego, CA). Cells were stained with combinations of appropriate MAbs at 4°C for 20 min and analyzed on a FACSCalibur flow cytometer using Cell Quest software (BD Biosciences, Mountain View, CA).

Isolation of PBMCs, monocytes, and T cells and identification of NKT cells. Peripheral venous blood samples were collected in tubes containing heparin. Peripheral blood mononuclear cells (PBMCs) were isolated by density gradient centrifugation using Ficoll-Paque Plus solution (Amersham Bioscience, Uppsala, Sweden), and monocytes and T cells were purified from PBMCs by magnetically activated cell sorting (MACS) as previously described (9, 36). NKT cells were identified phenotypically as CD3⁺ CD1⁻ 6B11⁺ cells by flow cytometry (3, 9, 16, 34, 43).

NKT cell proliferation assays. The proliferative abilities of NKT cells were assayed by flow cytometry as previously described (9). Briefly, freshly isolated PBMCs were suspended in complete medium (RPMI 1640, 2 mmol/liter l-glutamine, 100 U/ml penicillin, and 100 μg/ml streptomycin) supplemented with 10% fetal bovine serum (Gibco BRL, Grand Is-
Reduced circulating NKT cell numbers in the peripheral blood of EPTB and PTB patients. Freshly isolated PBMCs from 90 age- and sex-matched HCs, 13 LTBI subjects, 18 patients with EPTB, and 55 patients with PTB were stained with PerCP-conjugated anti-CD3 and PE-conjugated anti-6B11 MAb and then analyzed by flow cytometry. Symbols represent subjects, and horizontal lines are median values. (A) Representative NKT cell percentages determined by flow cytometry; (B) percentages of NKT cells in peripheral blood lymphocytes; (C) absolute numbers of NKT cells in peripheral blood samples (per ml of blood). P values were determined by Mann-Whitney U test. *, P < 0.05; **, P < 0.005; ***, P < 0.001.

RESULTS
Reduced numbers of circulating NKT cells in EPTB and PTB patients. The percentages and absolute numbers of NKT cells in the peripheral blood samples of 55 PTB patients, 18 EPTB patients, 13 LTBI subjects, and 90 age- and sex-matched HCs were determined by flow cytometry. Percentages of NKT cells were significantly lower in EPTB and PTB patients than in HCs (medians, 0.07% and 0.03%, respectively, versus 0.10% [P < 0.001, respectively]) (Fig. 1A and B). Absolute NKT cell numbers were calculated by multiplying NKT cell percentages by total lymphocyte numbers per milliliter in peripheral blood. EPTB and PTB patients had significantly lower absolute NKT cell numbers than HCs (medians, 871 and 440 cells/ml, respectively, versus 2,151 cells/ml [P < 0.005 and P < 0.001, respectively]) (Fig. 1C).

Correlation between reductions in NKT cells and CRP in TB patients. To evaluate the clinical relevance of NKT cell levels in EPTB and PTB patients, we investigated relationships between NKT cell numbers and clinical parameters. Percentages and absolute numbers of NKT cells, proliferation indices, and expression levels of CD1d and PD-1 were compared among groups using the Mann-Whitney U test. The Wilcoxon signed-rank test was used to compare changes in the proliferative responses of NKT cells after blockade of PD-1 signaling with anti-PD-1 Abs. All statistical analyses were performed using SPSS (version 17.0) software (SPSS, Chicago, IL), and significance was accepted for P values of <0.05.
lute NKT cell numbers in peripheral blood and clinical parameters using Spearman’s correlation analysis (Table 2). The analysis showed that absolute NKT cell numbers were significantly correlated with lymphocyte counts and C-reactive protein (CRP) levels ($P = 0.002$ and $P = 0.001$, respectively). However, no significant correlations between age, body mass index, smoking, leukocyte counts, or platelet counts and absolute NKT cell numbers were found (Table 2).

Impaired proliferative response of NKT cells to $\alpha$-GalCer in EPTB and PTB patients. To examine the proliferative effects of $\alpha$-GalCer on NKT cells, PBMCs from 24 PTB patients, 16 EPTB patients, 9 LTBI subjects, and 30 age- and sex-matched HCs were cocultured with $\alpha$-GalCer for 7 days in the presence of 100 U/ml of IL-2. Percentages of NKT cells and proliferation indices were evaluated by flow cytometry, as described in Materials and Methods. The percentages of NKT cells among PBMCs from an HC subject markedly increased in response to $\alpha$-GalCer, from 0.38% on day 0 to 18.6% on day 7. Similarly, NKT cells from an LTBI subject expanded from 0.38% to 13.9% over the same period. In contrast, NKT cells from a PTB patient and an EPTB patient did not proliferate or proliferated only slightly (Fig. 2A). Although the magnitudes of responses to $\alpha$-GalCer were highly variable among all study subjects, overall proliferation indices were significantly lower in EPTB and in PTB patients than in HCs (medians, 2.9 versus 27.1 [$P < 0.001$] and 0.8 versus 27.1 [$P < 0.001$], respectively) (Fig. 2B). However, no significant differences between HCs and LTBI subjects were observed. In addition, we analyzed the relationship between serum CRP levels and NKT proliferation indices in 24 PTB patients and 16 EPTB patients. It was found that the proliferative response to $\alpha$-GalCer was inversely correlated with CRP level (correlation coefficient $r = -0.559$, $P = 0.001$) (Fig. 2C).

Increased NKT cell death in TB patients. To determine whether the impaired proliferative response of NKT cells to $\alpha$-GalCer was due to a loss of NKT cells in culture, PBMCs from 5 patients, 9 LTBI subjects, and 30 age- and sex-matched HCs were cocultured with $\alpha$-GalCer for 7 days in the presence of 100 U/ml of IL-2. Percentages of NKT cells and proliferation indices were evaluated by flow cytometry, as described in Materials and Methods. The percentages of NKT cells among PBMCs from an HC subject markedly increased in response to $\alpha$-GalCer, from 0.38% on day 0 to 18.6% on day 7. Similarly, NKT cells from an LTBI subject expanded from 0.38% to 13.9% over the same period. In contrast, NKT cells from a PTB patient and an EPTB patient did not proliferate or proliferated only slightly (Fig. 2A). Although the magnitudes of responses to $\alpha$-GalCer were highly variable among all study subjects, overall proliferation indices were significantly lower in EPTB and in PTB patients than in HCs (medians, 2.9 versus 27.1 [$P < 0.001$] and 0.8 versus 27.1 [$P < 0.001$], respectively) (Fig. 2B). However, no significant differences between HCs and LTBI subjects were observed. In addition, we analyzed the relationship between serum CRP levels and NKT proliferation indices in 24 PTB patients and 16 EPTB patients. It was found that the proliferative response to $\alpha$-GalCer was inversely correlated with CRP level (correlation coefficient $r = -0.559$, $P = 0.001$) (Fig. 2C).

Increased NKT cell death in TB patients. To determine whether the impaired proliferative response of NKT cells to $\alpha$-GalCer was due to a loss of NKT cells in culture, PBMCs from 5

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### Table 2: Spearman’s correlation coefficients for absolute NKT cell numbers with respect to clinical and laboratory findings in 73 TB patients

<table>
<thead>
<tr>
<th>Variable</th>
<th>Correlation coefficient ($r$)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yr)</td>
<td>-0.129</td>
<td>0.279</td>
</tr>
<tr>
<td>Body mass index (kg/m²)</td>
<td>0.142</td>
<td>0.354</td>
</tr>
<tr>
<td>Smoking (no. of pack-years)</td>
<td>0.164</td>
<td>0.175</td>
</tr>
<tr>
<td>Hemoglobin concn (g/dl)</td>
<td>0.208</td>
<td>0.077</td>
</tr>
<tr>
<td>Leukocyte count (cells/μl)</td>
<td>-0.185</td>
<td>0.117</td>
</tr>
<tr>
<td>Lymphocyte count (cells/μl)</td>
<td>0.361</td>
<td>0.002</td>
</tr>
<tr>
<td>Platelet count (10³ cells/μl)</td>
<td>-0.075</td>
<td>0.531</td>
</tr>
<tr>
<td>CRP concn (mg/dl)</td>
<td>-0.428</td>
<td>&lt;0.001</td>
</tr>
</tbody>
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![FIG 2](http://iai.asm.org/) Decreased proliferative response of NKT cells from EPTB and PTB patients to $\alpha$-GalCer. (A and B) Proliferation indices of NKT cells. Freshly isolated PBMCs (1 x 10⁶ cells/well) from 30 age- and sex-matched HCs, 9 LTBI subjects, 16 patients with EPTB, and 24 patients with PTB were cultured for 7 days in the presence of IL-2 (100 U/ml) and $\alpha$-GalCer (100 ng/ml) or 0.1% DMSO as a control. The cells were then stained with PerCP-conjugated anti-CD3 and PE-conjugated anti-6B11 MAb and analyzed by flow cytometry. Proliferation index was defined as the ratio of NKT cell percentage within CD3⁺ T cells on day 7 to NKT cell percentage within CD3⁺ T cells on day 0, and indices are expressed as fold increases. The results shown are representative of the experiments (A). Symbols represent subjects, and horizontal lines are median values (B). (C) The relation between serum CRP level and proliferation index in 24 PTB patients and 16 EPTB patients. (D) Increased apoptosis of NKT cells in TB patients. Freshly isolated PBMCs from 5 age- and sex-matched HCs and 5 TB patients were cultured for 7 days in the presence of IL-2 (100 U/ml) and $\alpha$-GalCer (100 ng/ml). NKT cell apoptosis was analyzed by flow cytometry using FITC-conjugated annexin V, PE-conjugated 6B11 MAb, 7-AAD, and APC-conjugated anti-CD3 MAb. Percentages of apoptotic (annexin V-positive) NKT cells and necrotic (7-AAD-positive) NKT cells on days 0 and 7 are shown. Results are presented as means ± SEMs. P values were determined by Mann-Whitney U test. *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$. 

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TB patients and 5 age- and sex-matched HCs were cultured with \( \alpha \)-GalCer for 7 days in the presence of 100 U/ml of IL-2. NKT cell apoptosis was then evaluated by flow cytometry, as described in Materials and Methods. Apoptotic and necrotic percentages of NKT cells were minimal on day 0, and no significant differences between HCs and TB patients were found. However, after being stimulated with \( \alpha \)-GalCer for 7 days, NKT cell apoptosis and necrosis were greater for TB patients than for HCs (means, 27.4% versus 12.8% \( P < 0.05 \) and 43.7% versus 10.2% \( P < 0.05 \), respectively) (Fig. 2D). These findings suggest that increased NKT cell apoptosis in culture is related to the impaired proliferative response of NKT cells observed in TB patients.

Reduced expression of CD1d in EPTB and PTB patients. CD1d is known to be required for the presentation of glycolipid antigens to NKT cells. Thus, we hypothesized that the poor response to \( \alpha \)-GalCer observed in EPTB and PTB patients might be due to an altered CD1d expression. To determine whether CD1d levels differed in patients and HCs, percentages of CD1d\(^+\) cells and mean fluorescence intensities (MFIs) of CD1d were evaluated by flow cytometry in the peripheral blood samples of 17 PTB patients, 9 EPTB patients, 11 LTBI subjects, and 17 age- and sex-matched HCs. The percentages of CD1d\(^+\) cells among PBMCs were found to be significantly lower in PTB patients than in HCs (medians, 4.5% versus 6.3% \( P < 0.05 \)) (Fig. 3A). Furthermore, the MFIs of CD1d on PBMCs were also significantly lower in EPTB and in PTB patients than in HCs (medians, 4.8 versus 9.4 \( P < 0.001 \) and 5.2 versus 9.4 \( P < 0.001 \), respectively) (Fig. 3C). However, CD1d levels were similar in LTBI subjects and HCs.

Elevated PD-1 expression in NKT cells in EPTB and PTB patients. PD-1 and its ligands, PD-L1 and PD-L2, deliver inhibitory signals that regulate the activation of conventional T cells (25). Moreover, recent reports have shown that PD-1/PD-L interaction is involved in the induction and maintenance of NKT cell anergy (5, 35). To determine whether the NKT cell dysfunction is related
to NKT cell anergy, we thus examined the expression levels of this receptor in NKT cells in the peripheral blood samples of 6 PTB patients, 6 EPTB patients, 6 LTBI subjects, and 8 age- and sex-matched HCs. (Fig. 5A and B). Interestingly, the percentages of PD-1-expressing cells in NKT cells were found to be significantly higher in EPTB and in PTB patients than in HCs (means, 31.4% versus 9.8% \( P < 0.05 \) and 45.1% versus 9.8% \( P < 0.01 \), respectively) (Fig. 5B). In addition, the MFIs of PD-1 in NKT cells were also significantly higher in EPTB and in PTB patients than in HCs (means, 53.1 versus 27.0 \( P < 0.05 \) and 79.1 versus 27.0 \( P < 0.01 \), respectively) (Fig. 5B). To examine whether PD-1 contributes to the low proliferative response of NKT cells in TB patients, PBMCs from 4 EPTB patients and 3 PTB patients were cultured with \( \alpha \)-GalCer for 14 days in the presence of anti-PD-L1 plus anti-PD-L2 Abs or an isotype control. Although the magnitudes of responses to \( \alpha \)-GalCer were variable among the study subjects, overall proliferation indices were found to be increased after blockade of PD-1 signaling \( (P < 0.05) \) (Fig. 5C).

**DISCUSSION**

It has previously been reported that NKT cell numbers are reduced in PTB patients (33, 40, 42) and that NKT cell levels in LTBI subjects are comparable to those in HCs (40, 42). In the present study, we classified active TB into PTB and EPTB according to infected sites. Our data showed that NKT cell percentages and numbers were reduced in PTB patients and in EPTB patients but not in LTBI subjects. These findings suggest that an NKT cell deficiency reflects the presence of active TB irrespective of tuberculin skin test or IFN-\( \gamma \) release assay reactivity or the site infected.

In the present study, we also found that NKT cell responses to \( \alpha \)-GalCer were markedly suppressed in PTB and EPTB patients. However, NKT cell responses were not found to be diminished in LTBI subjects or HCs. This poor responsiveness of NKT cells has previously been demonstrated in patients with autoimmune, infectious, or malignant diseases, but the precise mechanism involved has not been elucidated (9, 11, 22, 27, 32, 44). Recently, several studies have demonstrated that microorganisms can expand murine NKT cells and subsequently become unresponsive to \( \alpha \)-GalCer-induced activation (7, 10, 26), which suggests that NKT cell function is affected by microbial infection. Our data also demonstrated that NKT cell levels and their proliferative responses to \( \alpha \)-GalCer were inversely correlated with CRP level, which suggests that NKT cell frequency and function are suppressed by chronic microbial infection (e.g., with \( M. \) *tuberculosis*). A negative correlation between NKT cell frequency and CRP level was also observed in patients with chronic autoimmune arthritis (e.g., rheumatoid arthritis) (44).

Although the poor response of NKT cells is not fully understood, it might reflect the lower representation of the most responsive subset in the peripheral blood of TB patients. Human NKT cells can be classified into two or three subsets (e.g., CD4\(^+\), CD8\(^+\), and double-negative NKT cells) according to the expression of CD4 and CD8 (20, 34). NKT cell subset levels in the peripheral blood of HCs and PTB patients were different in our study cohort (see Fig. S2 in the supplemental material). The most responsive subset to \( \alpha \)-GalCer was found to be CD4\(^+\) NKT cells (see Fig. S3 and S4 in the supplemental material). Despite the high frequency of this particular subset (i.e., CD4\(^+\) NKT cells), PTB patients showed low proliferative responses to \( \alpha \)-GalCer. Given the overall lower frequency of NKT cells in PTB patients (>75% absolute decrease compared to HCs), the poor response of NKT

**FIG 4** NKT cell abnormalities in EPTB and PTB patients. (A and C) Cocultures of APCs and T cells with \( \alpha \)-GalCer. CD14\(^+\) monocytes and CD3\(^-\) T cells were purified from PBMCs by MACS. Monocytes were used as a source of APCs. Freshly isolated monocytes (1 \( \times \) 10\(^5\) cells/well) were cocultured with T cells (9 \( \times \) 10\(^5\) cells/well) for 7 days as described in Materials and Methods. Cells were then stained with PerCP-conjugated anti-CD3 and PE-conjugated anti-6B11 MAb s and analyzed by flow cytometry. The results shown are representative of the experiments. (B and D) Results were obtained from three independent donors for each group and are presented as means \( \pm \) SDs.

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cells could be due to the absence of the responding cells from the peripheral blood following recruitment at the site of infection.

The loss of NKT cells in culture due to enhanced apoptosis provides another possible explanation for the poor response of NKT cells to \( \alpha \)-GalCer in TB patients. To examine this possibility, we measured apoptotic NKT cell numbers among PBMCs cultured with \( \alpha \)-GalCer by flow cytometry. The data obtained clearly showed that NKT cells from TB patients were more susceptible to apoptosis during \( \alpha \)-GalCer stimulation than NKTs from HCs. Previous studies have demonstrated that mycobacterial infections in mice and humans promote the apoptosis of activated immune cells (e.g., T cells or NK cells) (12, 30, 39). However, in our culture system, the apoptotic proportions of T cells and of NK cells were comparable for TB patients and HCs (data not shown). These findings suggest that the increased apoptosis of NKT cells in culture may reflect a poor response to \( \alpha \)-GalCer in TB patients.

CD1d presents glycolipid antigens, such as \( \alpha \)-GalCer to NKT cells (24). Furthermore, previous studies have shown that blocking CD1d, using MAb or a germ line deletion, inhibits the NKT cell response to \( \alpha \)-GalCer (24, 41, 49). Thus, we considered the possibility that numerical and functional deficiencies of NKT cells in patients might be due to alterations in CD1d expression. As we expected, CD1d expressions on PBMCs and CD14\(^+\) monocytes were found to be downregulated in TB patients. Furthermore, the downregulation of CD1d expression has also been observed in HIV (6, 8) and malignant lymphoma (22). Given that CD1d-restricted NKT cells have unique effector and regulatory functions in innate and adaptive immune responses, these findings suggest that \( M. \) tuberculosis reduces the presentation of CD1d to NKT cells and subverts host immune responses to infected cells.

The poor response to \( \alpha \)-GalCer observed in TB patients could be caused by NKT cell dysfunction as well as by altered CD1d expression. To investigate this possibility, we established co-culture systems using NKT cells and monocytes from HCs and TB patients. Monocytes from patients were found to expand NKT cells from HCs as efficiently as monocytes from HCs, indicating that the monocytes of TB patients are functional. In contrast, NKT cells from patients did not proliferate even when cocultured with monocytes from HCs. Interestingly, similar findings have also been reported for autoimmune and malignant diseases (9, 22, 27). These results provide evidence that the poor response to \( \alpha \)-GalCer is probably due to a defect in the NKT cells of TB patients.

PD-1 is well-known as a coinhibitory molecule on T cells. In conventional T cells, it is not expressed on naïve T cells but is inducibly expressed after T cell activation (25). In recent reports, PD-1 and its ligands have been implicated in the induction and maintenance of NKT cell anergy (5, 35). Our data showed an elevated PD-1 expression on NKT cells in TB patients, suggesting activation of these cells in vivo, findings that have also been observed in HIV-1 infection (32). Interestingly, blockade of the PD-1/PD-L pathway enhanced the response of NKT cells to \( \alpha \)-GalCer.
These findings suggest that PD-1 signaling at least in part contributes to the poor response of NKT cells in TB patients.

Summarizing, our findings show that NKT cells are numerically and functionally deficient in TB patients. In addition, NKT cell deficiencies were found to reflect the presence of active TB and to be correlated with CRP levels. These findings have important implications for the development of α-GalCer-based immunotherapy in M. tuberculosis patients.

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