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4-1BB Enhances Proliferation of Beryllium-Specific T Cells in the Lung of Subjects with Chronic Beryllium Disease

Douglas G. Mack,* Allison K. Lanham,* Brent E. Palmer,* Lisa A. Maier,** Tania H. Watts,† and Andrew P. Fontenot*†‡

In contrast to naive T cells, reactivation of memory cells is less dependent on CD28-mediated costimulation. We have shown that circulating beryllium-specific CD4⁺ T cells from chronic beryllium disease patients remain CD28-dependent, while those present in the lung no longer require CD28 for T cell activation. In the present study, we analyzed whether other costimulatory molecules are essential for beryllium-induced T cell function in the lung. Enhanced proliferation of a beryllium-responsive, HLA-DR-restricted T cell line was seen after the induction of 4-1BB ligand expression on the surface of HLA-DR2-expressing fibroblasts. Following beryllium exposure, CD4⁺ T cells from blood and bronchoalveolar lavage of chronic beryllium disease patients up-regulate 4-1BB expression, and the majority of beryllium-responsive, IFN-γ-producing CD4⁺ T cells in blood coexpress CD28 and 4-1BB. Conversely, a significant fraction of IFN-γ-producing bronchoalveolar lavage (BAL) T cells express 4-1BB in the absence of CD28. In contrast to blood, inhibition of the 4-1BB ligand-4-1BB interaction partially blocked beryllium-induced proliferation of BAL CD4⁺ T cells, and a lack of 4-1BB expression on BAL T cells was associated with increased beryllium-induced cell death. Taken together, these findings suggest an important role of 4-1BB in the costimulation of beryllium-responsive CD4⁺ T cells in the target organ. The Journal of Immunology, 2008, 181: 4381–4388.

Optimal activation of naive T cells requires signals through the TCR following engagement with cognate Ag and MHC on the surface of APCs as well as a second signal, termed costimulation (1, 2). This costimulatory signal is most frequently mediated by engagement of the CD28 molecule on the T cell with its ligands, B7-1 or B7-2, on the APC. CD28 is constitutively expressed on almost all human CD4⁺ T cells. However, in the setting of chronic inflammation or advanced age, a subset of circulating CD4⁺ T cells lose CD28 expression (3–7). Down-regulation of CD28 expression has also been observed on bronchoalveolar lavage (BAL) CD4⁺ T cells from patients with sarcoidosis (8) and chronic beryllium disease (CBD) (9). In fact, blockade of CD28-B7 interaction with CTLA-4Ig had no effect on either the beryllium-induced proliferation or Th1-type cytokine expression of BAL CD4⁺ T cells from CBD patients (9). These findings suggest a functional independence of beryllium-responsive CD4⁺ effector memory T (TEM) cells in lung from CD28 costimulation and raise the possibility that other costimulatory ligand-receptor interactions may assume a critical role in activating and sustaining the T cell response in the lung.

4-1BB is an inducible costimulatory molecule of the TNF receptor family that is expressed on activated CD4⁺ and CD8⁺ T cells (10, 11), while its ligand, 4-1BBL, is expressed on activated macrophages, dendritic cells, and B cells (11–13). Although some reports have suggested that 4-1BB is primarily involved in the activation of CD8⁺ T cells, recent studies have shown that both CD4⁺ and CD8⁺ T cells respond to 4-1BBL with enhanced T cell proliferation, effector function, and survival (11, 14). In humans, the interaction of 4-1BBL-4-1BB enhances Th1-type cytokine expression (15, 16). In addition, 4-1BB has been shown to be an important costimulatory molecule on T cells lacking CD28 expression (17, 18), suggesting its potential importance in the activation of lung CD4⁺ T cells because a large portion of these cells have lost CD28 expression (9). Although the role of 4-1BBL in the activation of human anti-influenza and HIV-specific CD8⁺ T cells has been studied (19–21), the role of 4-1BBL-4-1BB in the costimulation of Ag-specific CD4⁺ TEM cells in a target organ has not been addressed.

CBD is characterized by the presence of granulomatous inflammation and progressive fibrosis in the lung (22). Historically, one-third of untreated patients progressed to end-stage respiratory insufficiency (23). Considerable evidence indicates that the accumulation and activation of beryllium-specific CD4⁺ T cells in the lung is central to the pathogenesis of disease (24–27). The ability of peripheral blood and/or BAL CD4⁺ T cells to proliferate in the presence of beryllium salts in vitro forms an essential part of the disease definition (28–31). In the lung, large numbers of beryllium-specific CD4⁺ T cells accumulate (25), with the vast majority expressing an effector-memory phenotype and demonstrating immediate release of IFN-γ, TNF-α, and IL-2 when stimulated with beryllium salts in culture (25, 32). With a known Ag stimulus and access to large numbers of pathogenic T cells from the target organ, CBD is an important organ-specific, immune-mediated disease.
Based on the enhanced proliferation of a beryllium-responsive CD4⁺ T cell line after 4-1BBL expression on the APC, we examined the role of the 4-1BBL-4-1BB interaction in the costimulation of beryllium-specific T cells in blood and lung. In the present study, the results demonstrate that the majority of beryllium-responsive CD4⁺ T cells in blood coexpress CD28 and 4-1BB. Inhibition of the 4-1BBL-4-1BB interaction had no effect on beryllium-induced proliferation, suggesting that the dominant costimulatory molecule in blood is CD28. Conversely, a significant fraction of IFN-γ-producing CD4⁺CD28⁺ T cells in the BAL express 4-1BB, and blockade of 4-1BB costimulation inhibited the proliferation of BAL CD4⁺ T cells in response to beryllium. In addition, 4-1BB expression on BAL CD4⁺ T cells prevented activation-induced cell death after stimulation with Ag. These findings confirm the importance of 4-1BB in the beryllium-induced immune response in the lung and further support a transition within the memory CD4⁺ T cell population from CD28 dependence in blood to a requirement for 4-1BB costimulation in lung.

Materials and Methods

Study population

Seventeen patients with a diagnosis of CBD and eight beryllium-sensitized (BeS) patients were enrolled in this study. Ten healthy, non-beryllium-exposed control subjects were also enrolled. The diagnosis of CBD was established using previously defined criteria, including the presence of granulomatous inflammation on lung biopsy and a positive proliferative response of blood and/or BAL T cells to beryllium sulfate (BeSO₄) in vitro (26, 33). The diagnosis of BeS was established based on a positive proliferative response of PBMCs to BeSO₄ in vitro, and the absence of granulomatous inflammation or other abnormalities on lung biopsy (30, 31). Active smokers were excluded from enrollment. Informed consent was obtained from each patient and control subject, and the protocol was approved by the Human Subject Institutional Review Boards at the University of Colorado Denver and National Jewish Medical and Research Center (Denver, CO).

Preparation of cells and generation of beryllium-specific T cell lines

PBMCs were isolated from heparinized blood by Ficoll-Hypaque density gradient separation, and BAL was performed as previously described (34, 35). The beryllium-specific T cell lines were derived after initial stimulation of BAL cells in 12-well plates (Costar) with RPMI 1640 supplemented with 10% heat-inactivated human serum (Gemini Bio-Products), 20 mM HEPES, 1 mM sodium pyruvate, 100 U/ml penicillin, 100 µg/ml streptomycin, and 2 mM t-glutamine (all from Invitrogen), and the addition of 100 µM of BeSO₄, as previously described (36). After 5 days of culture, cells were washed and 105 cells/ml of BeSO₄ and the lymphoblasts were expanded further in culture medium supplemented with 20 U/ml human IL-2 (R&D Systems). The T cell lines were maintained in culture by cycles of restimulation (every 2–3 wk) with 100 µM of BeSO₄ and autologous, mitomycin C-treated, fibroblasts expressing HLA-DP2 and either B7.1, OX-40L, or 4-1BBL. The transfected fibroblast cells were washed three times with HBSS before use as APCs to ensure the absence of free BeSO₄ in culture.

Lymphocyte proliferation assay

PBMCs (2.5 x 10⁵ cells/well) and BAL cells (1 x 10⁵ cells/well) were cultured in 96-well flat-bottom microtiter plates in the presence of medium alone, 10 µM BeSO₄, or 2.5 µg/ml PHA. After 96 h of culture, the wells were pulsed with 1 µCi of [³H]thymidine for an additional 16 h, and incorporation of radioactivity was determined by β-emission spectroscopy. Proliferation assays were performed in triplicate. Inhibition of proliferation was attempted with addition of indicated amounts of mAbs directed 4-1BBL (M730; a gift from Amgen) at the beginning of culture as described above. Proliferation with and without the indicated mAb was determined and the extent of inhibition was calculated. In some experiments, beryllium-specific T cell lines (1 x 10⁵ cells) were cultured in 96-well flat-bottom microtiter plates with 5 x 10⁵ medium or BeSO₄-pulsed, mitomycin C-treated, fibroblasts expressing HLA-DR and either B7.1, OX-40L, or 4-1BBL. The transfected fibroblast cells were washed three times with HBSS before use as APCs to ensure the absence of free BeSO₄ in culture.

Immunofluorescence staining and analysis for intracellular cytokine expression

For cytokine expression, ex vivo PBMCs (2 x 10⁶ cells/tube) and BAL cells (1 x 10⁶ cells/tube) were placed in polypropylene tubes (12 x 75 mm; Fisher Scientific) containing 1 ml of complete culture medium plus one of the following experimental conditions: medium alone, 10 ng/ml staphylococcal enterotoxin B, or 100 µM of BeSO₄. Cells were incubated for a total of 20 h at 37°C in a humidified 5% CO₂ atmosphere with 10 µg/ml brefeldin A added after six hours of stimulation. This length of stimulation was necessary to allow for the induction of 4-1BB expression on the CD4⁺ T cell surface. In some experiments, inhibition of Th1-type cytokine expression was attempted with the addition of indicated amounts of 4-1BBL mAb at the beginning of culture. After stimulation, cells were washed and stained with mAbs directed against CD3 (PE-Texas Red, Beckman Coulter) and CD4 (APC-Cy7, BD Biosciences) for 30 min at 4°C, as previously described (9, 25). Cells were washed with PBS containing 1% BSA and permeabilized using a permeabilization medium (Caltag Laboratories) for 15 min at room temperature. Following washing with PBS containing 1% BSA, cells were added to permeabilization medium (Caltag Laboratories), and stained with mAbs directed against IFN-γ (FITC, BD Biosciences), IL-2 (APC, Caltag Laboratories), TNF-α (Alexa Fluor 700, Caltag Laboratories), 4-1BB (PE, BD Biosciences), and CD28 (PE-Cy5, BD Biosciences) for 30 min at 4°C. Fluorescence minus one and isotype controls were used in all staining. Cells were analyzed using a FACSria flow cytometer (BD Immunocytometry Systems) (37). The number of events collected ranged between 1 and 3 million. Electronic compensation was performed with Ab capture beads (BD Biosciences) that were stained separately with individual mAbs used in the test samples. The data files were analyzed using Diva software (BD Biosciences). Lymphocytes were gated based on their forward and side scatter profiles.

PBMCs or BAL cells were stained with anti-CD14-FITC (BD Biosciences) and PE-labeled anti-4-1BBL (BD Biosciences). Cells were first incubated with FcR-Blocking reagent (Miltenyi Biotec) to reduce nonspecific staining. Because 4-1BBL expression was not bimodal, expression levels on CD14⁺ T cells were determined by subtracting the mean fluorescence intensity (MFI) of the fluorescence minus one control from the 4-1BBL MFI on the population of interest. Fluorescence intensity was analyzed using a FACSCalibur cytometer (BD Immunocytometry Systems), as previously described (25, 38).

Apoptosis assay

BAL T cell lines from CBD patients were stimulated with medium, 10 µg/ml anti-CD3, or 100 µM of BeSO₄ for 24 h. Rates of apoptosis were determined on CD3⁺ CD4⁺ T cells with and without 4-1BB expression by staining with annexin-V and 7-aminoactinomycin D (7-AAD) (both from Molecular Probes), and the cells were immediately analyzed by flow cytometry as described above.

Statistical analysis

The Mann-Whitney U test and the Kruskal-Wallis ANOVA were used to determine significant differences between subject groups. A p value of <0.05 was considered statistically significant.
Results
Beryllium-responsive CD4+ T cells in blood produce more IFN-γ per cell than CD4+ T cells in BAL

The MFI of each functional parameter within a given sample is related to the quantitative expression of that function on a per cell basis (39). As a result, the relative amount of each cytokine per cell can be measured for each functional CD4+ T cell population. In addition, previous studies have shown that MFI measured by intracellular cytokine staining directly correlates with cytokine secretion as measured by ELISA (40). We compared IFN-γ MFI on beryllium-responsive CD4+ T cells in blood and in the target organ. Despite a 14-fold greater frequency of beryllium-responsive CD4+ T cells in BAL compared with blood (41), circulating IFN-γ-expressing beryllium-responsive CD4+ T cells were more functional than their counterparts in the lung, expressing 2.5 times more IFN-γ per cell (p = 0.0006) (Fig. IA).

Previous studies have shown that vaccinia-induced, multifunctional CD8+ T cells (i.e., secrete IFN-γ, IL-2, and TNF-α) produce more IFN-γ on a per cell basis compared with monofunctional cells (39). In response to BeSO4, we had previously identified two predominant populations of beryllium-responsive CD4+ T cells in blood and BAL: cells capable of expressing both IFN-γ and IL-2 and those expressing only IFN-γ (42). Thus, we sought to determine whether beryllium-responsive T cells that retained the ability to secrete IL-2 were more functional than those cells that had lost the ability to express this cytokine. As shown in the representative density plots of BeSO4-induced IFN-γ and IL-2 expression by CD4+ T cells from blood and BAL, beryllium-responsive CD4+ T cells expressing both IFN-γ and IL-2 were more functional than cells expressing only IFN-γ as measured by the MFI of IFN-γ per cell (Fig. 1B).

Overall, CD4+ T cells expressing IFN-γ/IL-2 in response to BeSO4 produced 3.4 and 2.2 times more IFN-γ per cell than IFN-γ-only-secreting CD4+ T cells in blood and BAL, respectively (Fig. 1C).

4-1BBL enhances beryllium-induced CD4+ T cell proliferation

The decreased functional capacity of the beryllium-responsive CD4+ T cell in the lung raised the possibility of different costimulatory requirements existing between the circulating Ag-specific T cells and those present in the target organ. We have previously shown that beryllium-responsive CD4+ T cells in blood remain CD28 costimulation-dependent, while those in the lung no longer require the engagement of CD28 for effector T cell function with a substantial fraction of these cells having lost CD28 expression (9). In the absence of CD28, we sought to determine whether CD4+ T cells in the lung were costimulation-independent or required another costimulatory molecule, such as OX-40 or 4-1BB, for T cell activation. Using replication-defective recombinant adenoviruses expressing full-length 4-1BBL, OX-40L, B7.1, or no transgene (Control-Adv) as previously described (19, 21), expression of 4-1BBL, OX-40L, and B7.1, respectively, on the surface of murine HLA-DP2-expressing DAP3-L cells was determined 24 h after infection (Fig. 2A). The ability of these BeSO4-pulsed fibroblasts expressing either 4-1BBL, OX-40L, or B7.1 to stimulate a beryllium-responsive, HLA-DP2-restricted CD4+ T cell line derived from the lung of a CBD patient was determined. As shown in Fig. 2B, HLA-DP2-expressing fibroblasts were able to initiate...
beryllium-induced T cell proliferation (1512 ± 135 cpm) compared with background proliferation (236 ± 114 cpm; stimulation index = 6.4). No augmentation of the beryllium-induced T cell proliferative response was seen following the expression of either B7.1 or OX-40L on the HLA-DP2-expressing fibroblasts. Conversely, an 8.4-fold increase in BeSO₄-induced T cell proliferation (12,751 ± 803 cpm) was seen after the expression of 4-1BBL on the surface of the HLA-DP2-expressing fibroblasts. No additive or synergistic effects of combining 4-1BBL costimulation with either OX-40L or B7.1 was seen (data not shown). These findings suggest that the binding of 4-1BB on the surface of the beryllium-specific CD₄⁺ TEM cell to 4-1BBL on the APC may represent a critical costimulatory interaction in the lung of CBD patients.

4-1BB expression on BeSO₄-stimulated CD₄⁺ T cells in BAL and blood

Due to the role of the 4-1BBL-4-1BB interaction in enhancing beryllium-induced proliferation of a beryllium-responsive CD₄⁺ T cell line, we analyzed 4-1BB expression on ex vivo BAL CD₄⁺ T cells before and after exposure to 100 µM BeSO₄ in culture. A representative example of the induction of 4-1BB on the surface of BAL CD₄⁺ T cells over a 96-h time period is shown in Fig. 3A. Even though the vast majority of BAL CD₄⁺ T cells express an activated TEM cell phenotype (38), few of these cells express 4-1BB ex vivo (median, 1.0%; range, 0.3–2.9%) (Fig. 3, A and B). After BeSO₄ exposure, 4-1BB was rapidly up-regulated, being expressed by a median of 18% of BAL CD₄⁺ T cells at 24 h (range, 12–30%). In addition, 4-1BB expression peaked at 24 h and remained stable over the 96 h time course. For example, at 96 h, 19% (range, 12–30%) of BAL CD₄⁺ T cells expressed 4-1BB (Fig. 3B). Although the frequency of 4-1BB-expressing T cells in blood was dramatically reduced due to a lower precursor frequency of beryllium-responsive cells, similar findings were seen upon exposure of PBMCs to BeSO₄. For example, a median of 0.5%, 2.3%, 3.0%, 1.5%, and 2.4% of CD₄⁺ T cells expressed 4-1BB ex vivo and after BeSO₄ exposure for 24, 48, 72, and 96 h, respectively (Fig. 3B).

4-1BB expression on beryllium-responsive, IFN-γ-expressing CD₄⁺ T cells on blood and BAL

Next, we analyzed 4-1BB expression on beryllium-responsive, IFN-γ-expressing CD₃⁺CD₄⁺ T cells in the blood and BAL of CBD patients. In the representative histograms shown in Fig. 4A, 84% and 43% of IFN-γ-expressing CD₄⁺ T cells in the blood and BAL, respectively, expressed 4-1BB. Overall, a significantly greater percentage of IFN-γ-producing T cells in blood (median, 80%; range, 48–100%) had up-regulated 4-1BB expression compared with their BAL counterparts (53%; range, 6.5–77%; p = 0.0004) (Fig. 4B). Similar to the decreased expression of IFN-γ per BAL CD₄⁺ T cell compared with blood shown in Fig. 1A, these findings are consistent with a hyporesponsive nature of Ag-specific T cells residing in the target organ.

To determine whether 4-1BB expression enhanced beryllium-induced IFN-γ production, we compared the expression of IFN-γ on a per cell basis in response to 100 µM BeSO₄ on blood and BAL CD₄⁺ T cells with and without surface 4-1BB. In both blood and BAL, a significantly increased IFN-γ MFI was seen on beryllium-specific, CD₄⁺4-1BB⁺ T cells (p < 0.0001 for blood and p = 0.02 for BAL) (Fig. 4C). Interestingly, beryllium-responsive 4-1BB-expressing CD₄⁺ T cells in blood (median, 64819; range, 39,308–152,300) produced significantly more IFN-γ per cell than those cells in the lung (32,745; range, 15,404–78,067; p = 0.01) (Fig. 4C). Based on the dependence of beryllium-specific T cells in

**FIGURE 3.** Induction of 4-1BB expression on the surface of CD₄⁺ T cells from blood and BAL of CBD patients after BeSO₄ exposure. A, A representative example is shown of the time course of 4-1BB induction on the surface of BAL CD₄⁺ T cells ex vivo and after BeSO₄ exposure for 24, 48, 72, and 96 h. B, 4-1BB expression of BAL cells (n = 5) and PBMCs (n = 6) ex vivo and after 100 µM BeSO₄ exposure for up to 96 h is shown.

**FIGURE 4.** Induction of 4-1BB on the surface of IFN-γ-expressing CD₄⁺ T cells from blood and BAL of CBD patients. A, A representative example of 4-1BB expression on IFN-γ CD₄⁺ T cells from blood and BAL is shown, with the percentage of 4-1BB⁺ T cells displayed in the upper right of the histogram. B, Percentage of IFN-γ⁺ CD₄⁺ T cells from blood (n = 17) and BAL (n = 11) expressing 4-1BB is shown. C, The expression of IFN-γ (MFI) on beryllium-responsive CD₄⁺ T cells in blood (left) and BAL (right) with and without the up-regulation of 4-1BB is depicted. Median values for B and C are indicated and shown with solid lines, and statistical comparisons were made using the Mann-Whitney U test. D, The percentage of IFN-γ⁺ CD₄⁺ T cells from blood (left) and BAL (right) in relation to CD28 and 4-1BB expression is shown. Median values are indicated with solid lines, and statistical comparisons were made using the Kruskal-Wallis test.
blood on CD28 costimulation for optimal activation (9), these findings raise the possibility that the continued requirement of blood cells for CD28-mediated costimulation enhanced IFN-γ production compared with lung T cells which no longer require costimulation through the B7-CD28 interaction. When analyzing IFN-γ expression in blood, the majority (median, 75%; range, 57–91%; \( p < 0.0001 \)) of beryllium-responsive CD4+ T cells coexpress CD28 and 4-1BB (Fig. 4D). Conversely, only 39% (range, 34–47%; \( p = 0.02 \)) of beryllium-responsive, IFN-γ-expressing CD4+ T cells in the BAL coexpressed CD28 and 4-1BB, with the majority of the remaining cells being 4-1BB− CD28−. These findings further emphasize the potential importance of 4-1BB in the costimulation of beryllium-specific T cells in the lung.

**Increased expression of 4-1BBL in the lung**

4-1BB ligands are expressed on the APC. To examine this, 4-1BB ligand expression was measured by flow cytometry on CD14+ monocyte/macrophages in PBMCs and BAL cells from BeS, CBD, and normal control subjects. There was no difference between groups in 4-1BBL expression on CD14+ cells from blood (Fig. 5). The median 4-1BBL expression (MFI) on CD14+ cells from the peripheral blood of normal control, BeS, and CBD subjects was 25, 7.3, and 11, respectively (Fig. 5). However, 4-1BBL expression on CD14+ cells from the BAL was significantly elevated. Median 4-1BBL expression (MFI) on CD14+ cells in the BAL of BeS (88; range 42–116) and CBD (102; range 55–135) subjects was 12- and 9.2-fold higher than in blood. These data demonstrate that CD14+ APCs in the BAL of BeS and CBD patients have significantly higher levels of 4-1BB ligands than those in blood and suggest that the increased expression of 4-1BBL on CD14+ BAL cells is not specific for CBD patients, but indicative of an activated phenotype of CD14+ cells in the lung.

**Functional dependence of BAL CD4+ T cells on 4-1BB-mediated costimulation**

We used inhibitory concentrations of a 4-1BBL mAb to test whether cells from CBD patients require 4-1BB-mediated costimulation for BeSO4-induced T cell activation in culture. PBMCs and BAL cells from 8 CBD patients demonstrated a positive proliferative response (defined as a stimulation index of \( \geq 2.5 \)) in the presence of 10 \( \mu M \) BeSO4. A representative example of beryllium-induced T cell proliferation of PBMCs from a CBD patient is shown in Fig. 6A, with a peak thymidine incorporation of 33,004 ± 803 cpm compared with background proliferation of 223 ± 138 cpm. Following the addition of 30, 3.0, or 0.3 \( \mu g/ml \) 4-1BBL mAb, no inhibition of beryllium-induced proliferation was seen. In contrast, BAL cells from the same individual showed a 45%, 54%, and 0% inhibition of beryllium-induced T cell proliferation in the presence of 30, 3.0, or 0.3 \( \mu g/ml \) 4-1BBL mAb, respectively (Fig. 6B).

Fig. 6C shows the overall effects of varying concentrations of 4-1BBL mAb on the proliferative responses of PBMCs and BAL cells. These studies again used optimal stimulatory concentrations of BeSO4 (10 \( \mu M \)). Proliferation of BAL cells was partially inhibited with concentrations of 4-1BBL ranging from 0.3 to 30 \( \mu g/ml \) (Fig. 6, B and C). As shown in Fig. 6C, 0.3 \( \mu g/ml \), 3 \( \mu g/ml \), and 30 \( \mu g/ml \) 4-1BBL inhibited the beryllium-induced proliferative response of eight CBD patients by 16 ± 6.4%, 33 ± 5.5%, and 47 ± 7.9%, respectively. In contrast, 4-1BBL mAb had little effect on the proliferation of PBMCs in response to BeSO4, with a 3.8 ± 2.7% inhibition when 30 \( \mu g/ml \) 4-1BBL mAb was added to the culture conditions. Thus, as opposed to PBMCs, these studies indicate that the beryllium-induced proliferative response in the lung requires 4-1BB-mediated costimulation. In separate experiments, the addition of OX-40L mAb had no effect on beryllium-induced proliferation, regardless of the source of the cells (data not shown).

Despite increased IFN-γ expression on beryllium-responsive, CD4+ 4-1BB+ T cells, blockade of the 4-1BBL-4-1BB interaction had no effect on the percentage of IFN-γ+ CD4+ T cells in the BAL after short-term exposure to BeSO4. In the representative example shown in Fig. 7A, ~19% of CD4+ T cells expressed IFN-γ in response to 10 \( \mu M \) BeSO4, irrespective of whether 4-1BBL mAb had been added. Similar findings were seen in two other subjects (Fig. 7B). Thus, despite the link between 4-1BB costimulation and beryllium-induced T cell proliferation, beryllium-stimulated IFN-γ expression in a short-term assay occurs independently of 4-1BB.

**Activation-induced apoptosis of BAL CD4+ T cells in relation to 4-1BB expression**

We also investigated whether the expression of 4-1BB on beryllium-specific CD4+ T cells protected these cells from undergoing activation-induced cell death. BAL-derived beryllium-specific T

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**FIGURE 5.** 4-1BB ligand expression is elevated on CD14+ cells in the lung of BeS and CBD patients. 4-1BBL expression was evaluated on CD14+ cells from the peripheral blood of normal \( (n = 11) \), BeS \( (n = 8) \), and CBD \( (n = 16) \) patients and the BAL of BeS \( (n = 5) \) and CBD \( (n = 16) \) subjects. Median values are shown with solid lines. Statistical significance was established using the Kruskal-Wallis test.
we observed a 30-fold increase in the percentage of cells expressing IFN-γ in the presence of varying concentrations of anti-4-1BBL mAb and analysis of BAL CD4+/H9262 T cells from CBD patients with and without the addition of anti-4-1BBL mAb.

Discussion

Our previous work suggests that differences in the functional properties of Ag-specific T cells in blood and BAL of CBD patients: those cells capable of secreting both IFN-γ and IL-2 and those only able to express IFN-γ (25, 42). It is widely accepted that IFN-γ-producing CD4+ T cells are more differentiated than CD4+ T cells that retain the ability to secrete IL-2 (43). However, when we analyzed beryllium-responsive CD4+ T cells from blood and BAL for the expression of markers associated with either T cell senescence (e.g., CD57) or exhaustion (e.g., programmed death-1 (PD-1)), no significant difference in the expression of either PD-1 or CD57 on IFN-γ-only and IFN-γ/IL-2-secreting CD4+ T cells was observed, irrespective of the location of the beryllium-responsive T cell (37, 41). In the present study, we noted that beryllium-specific CD4+ T cells expressing both IFN-γ and IL-2 were more functional (i.e., producing more IFN-γ per cell) than an IFN-γ-only secreting CD4+ T cell in both blood and BAL. In addition, 4-1BB expression in response to beryllium exposure in culture enhanced IFN-γ expression. Thus, polyfunctional (IFN-γ−, IL-2+, and 4-1BB+) Ag-specific CD4+ T cells in blood and BAL express more IFN-γ than their monofunctional counterparts, further supporting the role of 4-1BB costimulation in enhancing the effector function of the beryllium-responsive CD4+ T cell.

In addition to an independence from CD28 costimulation, beryllium-specific CD4+ T cells residing in the target organ are characterized by up-regulation in the expression of negative costimulatory molecules, such as PD-1 (37) and CTLA-4 (9). Furthermore, the percentage of IFN-γ-producing, beryllium-responsive CD4+ T cells expressing 4-1BB in the BAL was significantly diminished compared with IFN-γ-producing cells in blood, and this lung T cell subset was predisposed to undergo activation-induced cell death upon beryllium exposure in culture. Thus, it is likely that the combination of the loss of CD28 expression from the surface of BAL CD4+ T cells, the expression of PD-1 and CTLA-4, and the marginal ability to these cells to up-regulate other costimulatory molecules, such as 4-1BB, contribute to the hyporesponsive state of the beryllium-specific CD4+ T cell in the lung as documented in the current study. Whether the same is true in the target organ of other immune-mediated human diseases is an important unanswered question.

Conversely, there is a continued reliance of memory T cells in blood on CD28 for T cell activation, and the majority of circulating beryllium-specific, 4-1BB-expressing CD4+ T cells coexpress...
CD28. This is in contrast to CD8+ T cells where 4-1BB is predominantly expressed on CD8+ cells (44, 45). In addition, despite the induction of 4-1BB on the surface of beryllium-responsive, IFN-γ-expressing CD4+ T cells in blood, there was no effect of blocking the 4-1BBL-4-1BB interaction on beryllium-induced T cell proliferation. When B7.1 and 4-1BBL were expressed simultaneously on blood monocytes, the kinetics of the anti-influenza CD8+ T cell response paralleled the response with B7.1 alone (19). This is likely due to the constitutive expression of CD28 as opposed to the inducible nature of 4-1BB on the T cell surface. Thus, these findings strongly suggest that despite the induction of 4-1BB on beryllium-responsive CD4+ T cells, CD28 delivers the dominant costimulatory signal in blood.

Interestingly, 4-1BBL blockade had no effect on the percentage of BAL CD4+ T cells expressing IFN-γ after short-term culture in the presence of beryllium salts. This was not entirely surprising given the inducible nature of 4-1BB on activated CD4+ T cells. Despite an increased expression of 4-1BBL on BAL CD14+ cells compared with blood, it is also possible that a greater expression may be required to prime the T cell response. Similar to our findings, 4-1BB has been associated with sustaining as opposed to initiating the immune response, primarily through the induction of Bcl-2, and prevention of apoptosis (11, 14, 46, 47). In the opposite manner, our findings with 4-1BBL blockade are reminiscent of studies in HIV where PDL1-PD-1 blockade had no effect on intracellular Th1-type cytokine expression while enhancing proliferation of HIV-specific CD8+ T cells in blood (48). However, we have previously noted a discordance between proliferation and cytokine-secretion with the maturation state of the beryllium-responsive T cell dictating functional capacity (38). Thus, it remains possible that the beryllium-induced proliferative response of BAL cells is 4-1BB dependent while the Th1-type cytokine response is independent of costimulation.

The partial inhibition of proliferation seen with the 4-1BBL mAb raises the possibility that other inducible costimulatory molecules may be involved in the beryllium-induced immune response. In this regard, the induction of OX-40L expression on APCs had no effect on T cell proliferation, and no synergistic or additive effects were seen when OX-40L was combined with 4-1BBL. In addition, blockade of the OX-40L-OX-40 interaction with an inhibitory OX-40L mAb had no effect on beryllium-induced T cell proliferation, suggesting that OX-40 plays no role in this disease. In preliminary experiments, we have noted that CD40 ligand (CD40L) is up-regulated on BAL CD4+ T cells after BeSO4 exposure in culture, with 32% of BAL CD4+ T cells expressing CD40L at 96 h compared with 0.2% of ex vivo BAL T cells (data not shown). Whether inhibition of the CD40-CD40L interaction also blocks beryllium-induced proliferation of BAL T cells will require additional study.

Due to the presence of beryllium in the lung years after exposure cessation (49), the natural history of CBD is characterized by a gradual decline in lung function, with one-third of untreated patients histologically progressing to end-stage respiratory insufficiency (23). Combined with our previous work (9), our results clearly show that beryllium-induced activation of CD4+ T_{EM} cells in the BAL is independent of CD28 costimulation while requiring the interaction of 4-1BBL with 4-1BB for the proliferative response. In addition, the enhanced proliferation and prevention of activation-induced cell death of pathogenic CD4+ T cells upon 4-1BB costimulation suggest that the 4-1BBL-4-1BB interaction contributes to disease severity. Current therapeutic options for CBD patients are limited to corticosteroids which nonspecifically suppress T cell function. Blockade of the 4-1BBL-4-1BB interaction may down-regulate the function of beryllium-specific CD4+ T_{EM} cells, and as a result, diminish the severity of alveolitis. Thus, the development of targeted immunotherapy directed toward inhibiting 4-1BB costimulation in pathogenic T_{EM} cells in the lung has therapeutic potential, especially in diseases characterized by persistent Ag exposure.

In summary, despite the low frequency of circulating beryllium-specific CD4+ T cells, these cells are more functional than their counterparts in the lung. This enhanced functional capacity is more than likely due to the continued reliance of these cells on CD28 costimulation as opposed to beryllium-responsive T cells in the lung which are independent of CD28 and require 4-1BB engagement for an optimal beryllium-induced proliferative response. These findings further the developing paradigm in CBD and other immune-mediated disorders of a costimulatory transition within the memory CD4+ T cell population from CD28 dominance in blood to a requirement for other inducible costimulatory molecules, such as 4-1BB, on T_{EM} cells in a target organ. The induction of 4-1BB on Ag-specific memory T cells allows for enhanced effector T cell function and survival. In the case of CBD, this translates into continued lung inflammation and disease progression, suggesting the potential utility of inhibition of 4-1BB costimulation as a therapeutic target.

Disclosures
The authors have no financial conflict of interest.

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