**Immunopathology and Infectious Diseases**

**CC Chemokine Receptor 4 Contributes to Innate NK and Chronic Stage T Helper Cell Recall Responses during *Mycobacterium bovis* Infection**

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Cysteine-cysteinyl chemokine receptor 4 (CCR4) is reportedly expressed by platelets, natural killer (NK) cells, macrophages, basophils, and dendritic cells (DCs).† Regarding T-cell subsets, naïve T cells do not express CCR4, suggesting a role in memory or effector T-cell function. Initially, CCR4 was purported to be a marker of Th2 cells, but mounting evidence suggests that this is not the case. The CCR4 genetic deletion had no effect in a mouse model of Th2-dependent ovalbumin (OVA)–elicited allergy airway response.‡ We demonstrated that CCR4+ Th1 and Th2 cells were detected in mouse models of Th1 and Th2 cell-mediated pulmonary granulomatous inflammation, respectively, elicited by mycobacterial and helminth antigens.§ In a detailed analysis,§ human peripheral blood T cells, CCR4 expression was detected on isolated human CD4+ memory T cells with Th1 and Th2 characteristics. The specific ligands for CCR4, cysteine-cysteiny ligan (CCL)17, and CCL22 induced migration of both Th1 and Th2 cells in vitro.† In addition, CCR4 is reportedly expressed by human CD4+CD25+ T-regulatory cells and Th17 cells.∥ Other studies∥∥ support the role of CCR4 in T cells the specific capacity to migrate to skin. However, CCR4 appears to participate in modulation of lung and liver granulomatous inflammations and the innate response to peritoneal bacterial sepsis.§‡ In view of its expression among varied leukocyte populations, CCR4 is likely involved in multiple aspects of both innate and adaptive immunity.

The requirement for CCR4 in mounting a maximal Th1 response to mycobacterial antigens using experimental models of pulmonary granulomas elicited by challenge with bead-immobilized mycobacterial antigens was recently reported.∥ This finding was unexpected and provided further evidence that CCR4 expression and function were not restricted to Th2 cells. However, the role of CCR4 in the lung response to a live mycobacterial agent was unknown. In the...
present study, we further investigated CCR4 contribution to the antimycobacterial response in a model of live Mycobacterium bovis infection. By using CCR4 knockout mice, our study examined multiple parameters of the immune response after pulmonary exposure to mycobacteria. These included innate resistance, induction of effector cells in draining lymphoid tissues, mobilization of effector cells to infected lungs, and ability to mount a recall inflammatory response to mycobacterial antigens. The results show effects on the innate NK cell response and adaptive Th1 and Th17 memory responses, with complete sparing of γδ T-cell responses. The CCR4 appeared to be required for sustaining the local pulmonary memory effector response in the long-term but not early bacterial elimination stage of infection. Therefore, CCR4 may play a central role in homeostatic and late-stage organ-based effector responses. The findings have important implications for antimycobacterial vaccine design and treatment of chronic inflammation.

Materials and Methods

Animals

Mice lacking the CCR4 gene (CCR4<sup>−/−</sup>) were provided by Tularik Inc., (South San Francisco, CA) were generated as previously described,<sup>2</sup> and were bred onto a C57BL/6 background. Knockout status was confirmed by RT-PCR analysis using gene-specific primers and probes. The C57BL/6 wild-type (WT) mice were obtained from Jackson Laboratory, Bar Harbor, ME. CD90.1 (B6.PL–Thy1<sup>a</sup>) congenic and C57BL/6–Tg(TcrαTcrβ)425Cbn/J T cell receptor (TCR) ovalbumin transgenic (OT-II) mice were purchased from Jackson Laboratory. The CD4<sup>+</sup> T cells from the OT-II mice are specific for OVA peptide of amino acids 323–339 (EKLTEWTSSNVMEER) in the context of major histocompatibility complex antigen.<sup>26</sup> The OT–II TCR transgenic mice expressing CD90.1 on a C57BL/6 background were bred in house using male OT–II and female B6.PL–Thy1a/CyJ mice. All mice were maintained under specific pathogen–free conditions and provided with food and water ad libitum in an Association for Assessment and Accreditation of Laboratory Animal Care (AAALAC) approved facility. All animal studies were approved by an institutional review board.

M. bovis BCG Strains and Colony–Forming Unit Determinations

M. bovis BCG, Pasteur strain (Trudeau Institute, Saranac Lake, NY), was propagated in liquid 7H9 medium supplemented with 0.5% glycerol, 10% oleic acid–albumin–dextrose catalase and 0.05% polysorbate detergent (Tween 80). Organisms were harvested in mid–log growth stage, usually 16 to 20 days of culture at 37<sup>°</sup>C in a humidified 6% carbon dioxide atmosphere. Aliquots of BCG were stored at −80<sup>°</sup>C in PBS–glycerol (1:1). Preparations were washed with PBS before in vivo administration.

Recombinant BCG–OVA was prepared using a construct containing the green fluorescent protein gene driven by a mycobacterial heat shock protein 60 promoter and carrying a kanamycin resistance gene, which was provided by Glenn Fennelly (Albert Einstein Yeshiva University, New York, NY), as previously described.<sup>14</sup> Ovalbumin peptides recognized by the OT-I and OT-II TCR-transgenic T cells were cloned by PCR into the C-terminal end of the GFP molecule using extended primers. The altered construct was checked by sequencing, then BCG-Pasteur was transfected with the OVA peptide construct and selected in kanamycin-containing media. The transfected BCGs were GFP positive, and in <i>in vitro</i> splenic cultures both OT-I and OT-II cells were activated by the recombinant bacteria or bacterial lysates. The production of peptides was also confirmed by Western blotting. The course of BCG infection and the bacterial load determined by colony-forming units (CFUs) were similar in C57BL/6 mice infected with the WT or the recombinant GFP-OVA-BCG.

Colonies were determined by counting numbers of colonies developing from serial dilutions distributed on Middlebrook 7H10 agar (100-mm) plates with oleic acid–albumin–dextrose catalase supplement. Plates were cultured at 37<sup>°</sup>C in a humidified 6% carbon dioxide atmosphere for 2 to 3 weeks. For lung CFUs, whole lungs were homogenized in 20 ml of sterile PBS using a Waring blender. Tissue slurries were pelleted by centrifugation and resuspended in 2 ml of sterile water, and serial dilutions were applied to Middlebrook agar described for colony counting.

M. bovis BCG Infection

Mice were exposed to 1 million <i>M. bovis</i> BCG CFUs i.t. Briefly, mice were subjected to ketamine or xylazine anesthesia and then secured by soft restraint to an operating board to expose the neck. After iodine field disinfection, a 5- to 6-mm midline incision was made to expose the trachea. By using a 26-gauge needle and syringe, 1 × 10<sup>6</sup> organisms in a 0.025-ml volume were injected in the tracheal lumen, followed by closure of the skin wound with a 4-0 suture (Softsilk) on a C-13 cutting needle. For recovery, mice were placed on a water-jacketed heating pad to maintain body temperature.

Induction of Synchronized Anamnestic Granulomas

Secondary type 1 Ag-bead granulomas were generated as previously described.<sup>15</sup> Briefly, BCG-infected mice were challenged i.v. with 6000 beads (Sepharose 4B) covalently coupled to <i>M. bovis</i>–purified protein derivative (PPD) (Department of Agriculture, Veterinary Division, Ames, IA). For histological analysis, individual excised lung lobes were inflated and fixed with 10% buffered formalin for morphometric analysis. The granuloma area was measured in a blinded fashion in hematoxylin-eosin–stained sections of paraffin-embedded lungs using computer-assisted morphometry. A minimum of 20 lesions was measured per lung.
Lymph Node Cell Isolation and Culture

After perfusion with cold RPMI 1640 medium (Sigma Aldrich, St Louis, MO), lungs, excluding trachea and major bronchi, and mediastinal lymph nodes were excised. Lymph nodes were teased into a single-cell suspension and then cultured in RPMI 1640–supplemented medium (RPMI 1640 medium with 20-mmol/L HEPES buffer, 10% fetal bovine serum, 100-U penicillin, and 100-µg/ml streptomycin) at 5 × 10^6 cells/ml in the presence or absence of 10-µg/ml PPD. Cultures were incubated at 37°C with 5% carbon dioxide for 48 hours. Supernatants were collected by centrifugation and measured by enzyme-linked immunosorbent assay (ELISA).

Cytokine and Chemokine Measurement

Murine IL-17A and interferon (IFN) γ were measured in culture supernatants by ELISA using commercially available reagents and standards (Pharmingen, San Diego, CA; R&D Systems, Minneapolis, MN; Peprotecht Inc, Rocky Hill, NJ). Sensitivities were between 15 and 50 pg/ml. Murine CCL22 was measured in aqueous lung extracts by ELISA using commercial reagents (R&D Systems). Aqueous extracts were prepared as previously described.16

Immunohistochemical Staining

Paraffin-embedded lung sections were deparaffinized in 100% xylene for 3 minutes and in a 1:1 solution of xylene and ethanol for 3 minutes. Sections were rehydrated in 100%, 95%, 75%, and 50% ethanol solutions, 3 minutes each, and finally rinsed in PBS. Antigen retrieval used a 1:20 dilution of decloaker (Bullseye; Biocare Medical, Concord, CA) and incubation of sections at 125°C for 30 seconds, followed by a 90°C incubation for 10 seconds in a decloaking chamber (Biocare Medical, Concord, CA). Slides were rinsed in cold tap water after antigen retrieval and placed in PBS wash for 3 minutes. An avidin and biotin blocking kit (Biogenex, San Ramon, CA) and goat serum block (Biogenex) were used according to the manufacturer’s instructions. Rabbit anti–mouse CCL22 antibody (Abcam, Cambridge, MA), diluted 1:50 in PBS, was incubated on sections for 1 hour 15 minutes at room temperature. Nonimmune rabbit IgG served as a parallel control. Secondary goat anti-rabbit alkaline phosphatase–conjugated antibody was diluted to 1:1000 in PBS and incubated for 20 minutes. Fast red substrate kit (Biogenex) was prepared and incubated at 20 minutes. Hematoxylin was used as the counterstain.

Flow Cytometry and Intracellular Cytokine Staining

Single-cell suspensions were prepared from lungs as previously described.17 Briefly, after perfusion with cold RPMI 1640 medium, lungs were excised, placed in cold RPMI 1640 medium, and then homogenized for 15 seconds in a blender (Waring). Homogenates were incubated with digestion medium containing RPMI 1640 medium, 10% fetal bovine serum (Atlanta Biologicals, Norcross, GA), and 250-µM type IV collagenase (Sigma Aldrich) for 10 minutes at 37°C. The digest was sieved through a 40-µm cell strainer (BD Biosciences, San Jose, CA) and washed twice by centrifugation in RPMI 1640 medium. Single-cell suspensions were also prepared from freshly harvested spleens and lymph nodes by mechanical dissociation, as previously described.18 Total cell yields were determined by standard hemocytometric counting.

Monoclonal antibodies used included the following: fluorescein isothiocyanate–, phosphatidylethanolamine–, phosphatidylethanolamine-Cy5–, Pacific blue–, and anti–presenting cell (APC)–conjugated anti-CD4 (GK1.5), anti-CD8a (S3-6.7), anti–TCRαβ (H57-597), anti–TCRγδ (UC7-13D5), anti-NK1.1 (PK136), anti–IFNγ (R46A2 and XMG1.2), anti–IL-4 (11B11 and BVD6–24G2), anti–IL-17 (TC11-18H10), anti–Foxp3 (FJK-16s), anti-CD62L (MEL-14), anti-CD44 (IM7), anti–CD90.1 (HIS51), and anti–CD90.2 (S3-2.1) (BD Biosciences; or eBioscience, San Diego). All isotype controls and anti-CD16/CD32 (2.4G2) were from BD Biosciences. After blocking with anti-CD16/CD32 for 5 minutes, cells were stained with fluorescent-labeled antibodies or isotype control antibodies in 2% fetal bovine serum–PBS staining buffer for 25 minutes at 4°C. A multicolor flow cytometer (FACScan LSRII; BD Biosciences) with software (FlowJo; Tree Star, Ashland, OR) was used for data acquisition and analysis. Invariant NK T cells were identified using either fluorescein isothiocyanate–labeled (NIH Tetramer Core Facility, Atlanta GA) or phosphatidylethanolamine–fluorescent–labeled (Prommune, Bradenton, FL) CD1d tetramer (loaded and unloaded). Briefly, 100 µL of staining buffer and 0.5 µL of either loaded or unloaded (control) tetramer was added to each anti-CD16/CD32 blocked sample and then incubated for 30 minutes at 4°C. After washing, a second staining for additional surface markers was performed. The NKT cells were defined as CD1tetramer↑+TCRαβ↑+ CD8– cells.

Intracellular flow cytometric analysis of lung and mediastinal lymph node cells was performed as previously described.19 Briefly, single-cell suspensions were incubated for 3 hours in complete medium in the presence of brefeldin A, 1 µg/ml (BD Biosciences). In some cases, cells were stimulated in culture before adding brefeldin A. Cells were harvested and stained for surface antigen expression before staining for intracellular cytokines. Intracellular cytokine staining was performed using a kit (Cytofix/Cytoperm Kit; BD Biosciences) according to the manufacturer’s directions.

In vivo antigen presentation was detected using adoptive transfer of carboxyfluorescein succinimidyl ester–labeled CD4+ T cells (Invitrogen, Carlsbad, CA) from transgenic mice with OVA peptide-specific TCR (OT-II), as previously described.20 The number of analyzed events ranged from 500,000 to 1 million cells for all flow studies.

RNA Purification and Real-Time RT-PCR Analysis

RNA was isolated, reverse transcribed, and used for real-time PCR analyses, as previously described.21 Analysis of the transcripts was performed by real-time PCR using a
to elicit a highly polarized IFN multiple innate immune mechanisms activated in the early mice but is not useful for survival studies. In that stages of antimycobacterial immune responses in provides a means to examine both inductive and elicitation was similar in CCR4 knockout mice. Specifically, expression early after infection. The CCR4 agonist inducement was associated with increasing levels of CCR4 transcript expression at 1 week after infection and declined thereafter. The in-clearance of CCR4 in lungs after BCG exposure, we first assessed the temporal transcript expression of the two known agonists for this chemokine receptor, CCL17 (formerly thymus activated-regulated chemokine) and CCL22 (formerly macrophage-derived chemokine). As shown in Figure 1A, transcripts for both CCR4 agonists showed constitutive levels in preexposure lungs, with an increase after infection. There was significant CCL22 transcript induction at 1 week after infection, with levels increasing by nearly threefold and then declining thereafter with a trend to increased levels at 6 weeks; however, this was not statistically different from constitutive levels. CCL17 showed lower constitutive transcript expression than CCL22, but it also increased by nearly threefold by 1 week after infection and declined thereafter. The increase of CCR4 agonists at 1 week was associated with increasing levels of CCR4 transcript expression (Figure 1B), which peaked at 2 weeks and then declined thereafter. Thus, CCR4 agonists appear to be constitutively expressed chemokines, with BCG infection enhancing expression early after infection. The CCR4 agonist induction was similar in CCR4 knockout mice. Specifically, protein levels for CCL22 were measured in aqueous lung extracts by ELISA and revealed constitutive levels in naïve lungs (ie, 394 ± 40 pg/ml); at 1 week of infection, this increased to 748 ± 273 and 766 ± 264 pg/ml for WT and CCR4 knockout mice, means and standard deviations, respectively.

Immunohistochemical detection of agonist expression was also performed. Antibodies suitable for immunohistochemical detection are available only for CCL22 (formerly macrophage-derived chemokine). By using this preparation, we examined expression in the lungs of naïve and BCG-infected mice. As shown in Figure 2A, CCL22 was constitutively expressed in the lungs of uninfected mice among scattered alveolar septal mononuclear cells. One week after infection with BCG, CCL22 was detected in mononuclear cells associated with inflammatory nodules, alveolar macrophages, and interstitial mononuclear cells (Figure 2, C, E, and G). Sections stained with control antibody are shown in Figures 2B, 2D, 2F, and 2H. CCL22 was not observed in epithelial cells, smooth muscle, or endothelium at this time.

**CCR4 Knockout Mice Display Impaired Early Clearance of M. bovis BCG and Reduced Late Stage Inflammation**

Because CCR4 agonist expression was induced in lungs during BCG infection, we next examined the effect of CCR4 gene knockout on infection kinetics. Knockout (CCR4−/−) and control WT (CCR4+/+) mice were exposed to organisms i.t. and then lungs were harvested for quantification of CFUs. As shown in Figure 3A, both CCR4−/− and CCR4+/+ mice were able to largely eliminate organisms by 12 weeks of infection; however, knockout mice showed a statistically significant (nearly 1 log) early impairment of clearance at 2 weeks. The adaptive response to mycobacteria does not manifest in lungs until after 2 weeks, this finding suggested that early innate resistance might be impaired in CCR4−/− mice.
Th1 response is required for elimination of mycobacteria, antimycobacterial response. Although an adaptive CD4+ Th1 response participates in the early-stage antimycobacterial response,24 natural killer CD1d–restricted NK T (invariant NK T) and γδ T cells are thought to retard early mycobacterial expansion.25–28 However, although these populations can slow bacterial growth, they are insufficient for complete resistance to infection-level inoculums. Phenotypic CCR4 expression has been described among all of these innate-stage populations.29–31 Based on these reports, we hypothesized that compromise of an innate component might be responsible for the transient impairment of mycobacterial clearance observed in CCR4−/− mice.

Initial flow cytometric analyses revealed no quantitative defect in recruitment of αβ T, γδ T, invariant NK T, or NK1+ cells to the lungs of knockout mice (Table). Of these populations, NK cells dominated numerically and reached a maximum at 1 week after infection (Figure 4A). Because NK functional activity requires induction of cytokine-producing capacity after interacting with APCs,32 we assessed NK1.1+ cells for expression of intracellular IFNγ and IL-17. As shown in Figure 4, B and C, IFNγ was the dominant cytokine expressed by CD4− NK cells. The IL-17+ NK cells represented a minor population, with no differences detected between control and knockout mice. In contrast, CCR4−/− mice displayed a significant reduction in the activated IFNγ+ population, which persisted during the study period. This effect extended to both CD8+ and CD4− NK cells (data not shown). This finding indicated that CCR4 was required for maximal NK activation but not recruitment during the innate-stage response to M. bovis BCG infection and provided a potential explanation for impaired clearance.

When lungs were examined histologically, inflammation was comparable in both strains during the early elimination stage (2–4 weeks); thereafter, CCR4−/− mice showed less persistent inflammation in the late stage when residual lymphoid-rich inflammation predominated in control BCG-challenged lungs (Figure 3C). The histological impression was supported by quantification of cellularity of lung digests (Figure 3B). Therefore, CCR4 appeared to participate in the late (>4 weeks) and innate (<2 weeks) stages, but no clear effects were apparent during the rapid clearance stage (2–4 weeks).

Figure 2. Immunohistochemical detection of CCL22 in naïve and M. bovis BCG-infected lungs. A: Uninfected mouse lung demonstrating constitutive CCL22 expression (arrow) among scattered septal interstitial mononuclear cells. B: Serial section stained with control rabbit IgG. C: At 1 week, BCG-infected lung showing positive interstitial cell (arrow) near a venule. D: Serial section stained with control rabbit IgG. E: At 1 week, BCG-infected lung showing positive mononuclear cell (arrow) in an inflammatory nodule. F: Serial section stained with control rabbit IgG. G: At 1 week, BCG-infected lung showing positive alveolar macrophage (arrow). H: Serial section stained with control rabbit IgG. Original magnification, ×600. Arrows point to positively staining mononuclear cells.

CCR4 Knockout Impairs Late Stage CD4+ Effector T Cell But Not γδ T-Cell Mobilization to Lungs

Because CCR4−/− mice also displayed reduced late stage lung inflammation, we explored the contribution of CCR4 to later-stage cell-mediated antimycobacterial effector responses. To this end, CCR4+/+ and CCR4−/− mice were infected with M. bovis BCG and lungs were dispersed enzymatically, cultured with M. bovis PPD, and then analyzed by flow cytometry. Subpopulations of T cells and intracellular expression of IFNγ and IL-17 among these populations were measured during a 12-week study period. As shown in Figure 5, the proportion of CD8+ and γδ+ T cells reached a maximum at 2 weeks of infection and thereafter declined, with no differences observed between CCR4+/+ and CCR4−/− mice (Figure 5, left column). In contrast, CD4+ T cells reached a maximum at 6 weeks, with reductions observed in CCR4−/− mice at this time. Both IFNγ+ and IL-17+ effector T cells were identified in lungs at 2 and 6 weeks of infection, representing early and late bacterial elimination stages (Figure 5, right column). In control and knockout mice, cytokine-positive cells were most frequent among CD4+ and γδ+ T cells. The IFNγ+ cells were more dominant among CD4+ T (Th1), whereas IL-17+ cells were predominantly γδ T cells; however, CD4+ Th17 and IFNγ+...
ported late stage CD4+ (data not shown). These data suggested that CCR4 suppression was also detected in the minor IL-17 were unaffected. Interestingly, at 6 weeks, a significant CCR4 cell response to BCG was highly polarized toward IFN-γ, and IL-17–producing cells, with no IL-4+ T cells represented only 1% to 2% of lung cells, many (>90%) of γδ T cells were IL-17+. In the early stage (2 weeks), cytokine-positive cell populations were comparable among control and knockout mice. In contrast, by 6 weeks, CCR4-/- mice displayed 70% reductions of IFNγ+ CD4+ T cells (1.1% ± 0.3% versus 0.3% ± 0.1% for CCR4+/+ and CCR4+/- mice, respectively), although IL17+ γδ T cells were unaffected. Interestingly, at 6 weeks, a significant reduction was also detected in the minor IL-17+CD4+ population (0.14% ± 0.03% and 0.06% ± 0.02% for CCR4+/+ and CCR4-/- mice, respectively). The effector cell response to BCG was highly polarized toward IFNγ and IL-17–producing cells, with no IL-4+ cells detected (data not shown). These data suggested that CCR4 supported late stage CD4+ Th1 and Th17, but not γδ T-effector cells, in challenged lungs.

**CCR4 Mice Display Normal CD4+ T-Cell Proliferative Responses to Recombinant BCG-OVA and Effector/Memory Cell Generation**

The reduction in late stage Th effectors was potentially due to an impaired capacity to produce them. To test the induction of the adaptive response of CCR4-/- mice, we initially assessed Ag-presenting capacity using transgenic T-cell transfer and infection with recombinant mycobacteria. Control and knockout mice were infected with recombinant BCG-expressing OVA peptides. To identify the optimal period of T-cell proliferation in draining lymph nodes, groups of mice were infected with 1 million CFUs of recombinant BCG-OVA and administered OT-II cells on days 0, 7, and 13; then, lymph nodes were analyzed on days 6, 14, and 21, respectively. As reported for *M. tuberculosis* infection of mice, our studies showed that optimal Ag presentation is delayed in draining lymph nodes until beyond days 7 to 10 of infection (Supplemental Figure S1 at, see http://ajp.amjpathol.org). Thus, WT and knockout mice were administered carboxyfluorescein succinimidyl ester–labeled CD90 congenic D10.11 OVA-specific transgenic CD4+ T cells on day 13 after challenge with recombinant BCG-OVA. Then, on day 21, mice were sacrificed and draining lymph nodes and spleens were subjected to flow cytometric analysis. Figure 6A shows that CCR4+/+ and CCR4-/- mice displayed similar degrees of proliferation, with at least four generations detected. Moreover, proliferation was most apparent in draining lymph nodes than spleens, consistent with the lung being the primary site of Ag insult. No proliferation was

**Table 1.** Lung T- and NK Cell Populations on Day 8 of *Mycobacterium bovis* BCG Challenge

<table>
<thead>
<tr>
<th>Genotype</th>
<th>TCRγδ T cells</th>
<th>TCRγδ T cells</th>
<th>iNK T cells</th>
<th>NK1+ cells</th>
</tr>
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<tbody>
<tr>
<td>CCR4+/+</td>
<td>443 ± 116</td>
<td>17.4 ± 5.3</td>
<td>9.1 ± 2.4</td>
<td>775 ± 337</td>
</tr>
<tr>
<td>CCR4-/-</td>
<td>496 ± 89</td>
<td>20.7 ± 3.3</td>
<td>9.6 ± 2.4</td>
<td>834 ± 196</td>
</tr>
</tbody>
</table>

NK, natural killer; BCG, Bacille-Calmette-Guerin; TCR, T cell receptor; iNK, invariant NK; CCR4, cysteine-cysteiny1 chemokine receptor 4.

*Data are given as mean ± SE cell number (given in thousands per lung), derived from five to six lungs.
elicited with nonrecombinant BCG-WT organisms. Thus, the CCR4 knockout environment fully supported Ag presentation.

Despite normal Ag presentation, reduced numbers of CD4+ effector cells in BCG-challenged lungs might be secondary to impaired differentiation of effectors in draining lymph nodes. To explore this, we examined the capacity of CCR4−/− mice to generate effector cells in draining mediastinal lymph nodes. Groups of CCR4+/+ and CCR4−/− mice were infected, and their lymph nodes were subjected to intracellular cytokine staining to define the distribution of effector cell populations at 2 weeks. Figure 6B shows that both CCR4+/+ and CCR4−/− mice had similar numbers and types of effectors. The dominant IFN-γ population was CD4+ Th1 cells, with CD8+ and γ/δ T cells comprising the remaining IFNγ producers. The IL-17 producers were largely absent from lymph nodes at this time. Levels of Foxp3+ CD4+ T cells in spleens at 2 weeks (2.9% ± 0.3% and 3.2% ± 0.2% for CCR4+/+ and CCR4−/− mice, respectively; n = 4 per group). Thus, CCR4 knockout did not compromise the induction or spectrum effector cells generated in draining lymph nodes.

We also assessed the circulating Th memory cell compartment by comparing levels of CD4+ CD44hi CD62Llo cell in spleens at 12 weeks after infection, when bacteria are largely eliminated. As shown in Figure 6C, most CD4+ T cells in spleens were CD62Lhi, consistent with naïve T cells, but approximately 10% were CD4+CD62Llo, which were greater than 90% CD44+, consistent with Th memory cells. The CCR4−/− mice showed no gross changes in this compartment. Thus, the late stage defect in knockout mice was likely due to influences on local recruitment, activation, or survival.

Mycobacterium bovis BCG-Infected CCR4 Knockout Mice Display Impaired Late Stage IL-17 Production in Lymph Node Cultures

To assess late stage events in lymphoid tissue, we harvested draining mediastinal lymph nodes over 16 weeks of infection and cultured them with M. bovis PPD to elicit cytokines. At 48 hours, cell-free culture supernates were harvested and cytokine levels were measured by standard enzyme immunoassay as a gross measure of the
Bars are mean PPD-elicited cytokine-producing cells using intracellular cytokine staining. Later, the draining lymph node was harvested, dispersed, and analyzed for staining with isotype control antibody. CD4 generation and emigration of IFN-γ are shown. One million events were analyzed. A: The IFN-γ or IL-17–producing effector populations in draining lymph nodes. The CCR4+/+ and CCR4−/− mice were exposed to 1 million WT BCG; 2 weeks later, the draining lymph node was harvested, dispersed, and analyzed for PPD-elicited cytokine-producing cells using intracellular cytokine staining. Bars are mean ± SD (n = 5–6 mice). The dashed line indicates baseline staining with isotype control antibody. B: Flow cytometric assessment of CD4+ splenic T cells with CD62L−/− and CCR4+/+ and CCR4−/− mice had similar lymph node cell yields during the study period. The pattern of nodal IFN-γ production was similar among the strains, characterized by the highest levels of IFN-γ at 2 weeks when adaptive effectors appear; subsequently, these disappeared from nodes at 4 to 8 weeks, with partial reappearance beyond 8 weeks in the resolution stage (Figure 7B). This pattern is consistent with the generation and emigration of IFN-γ+ effectors, with later appearance of a circulating memory population. Interestingly, IL-17 became more prominent in lymph node cultures after 8 weeks, with CCR4−/− mice showing significantly reduced levels (Figure 7C). The IFN-γ levels also trended lower during this time but did not reach statistical significance. Intracellular cytokine staining of lymph node populations indicated that IL-17 was primarily associated with CD4+ αβ T cells, consistent with Th17 cells (data not shown). These findings suggested that BCG leads to late stage induction of Th17 cells, whose expression was impaired in CCR4−/− mice.

**Figure 6.** The CCR4 knockout does not affect CD4+ T-cell proliferation or effector/memory cell generation after *M. bovis* BCG infection. A: Transgenic T-cell proliferation. The CCR4+/+ and CCR4−/− mice were exposed to 1 million recombinant *M. bovis* BCG-OVA or control BCG i.t.; on day 13, the mice received 5 million carboxyfluorescein succinimidyl ester–labeled CD4+ CD90.1+ OT-II transgenic T cells. On day 21, draining lymph nodes and spleens were subjected to flow cytometric analysis. Representative scattergrams are shown. One million events were analyzed. B: The IFN-γ or IL-17–producing effector populations in draining lymph nodes. The CCR4+/+ and CCR4−/− mice were exposed to 1 million WT BCG; 2 weeks later, the draining lymph node was harvested, dispersed, and analyzed for PPD-elicited cytokine-producing cells using intracellular cytokine staining. Bars are mean ± SD (n = 5–6 mice). The dashed line indicates baseline staining with isotype control antibody. C: Flow cytometric assessment of CD4+ splenic T cells with CD62L−/− and CCR4+/+ and CCR4−/− memory phenotype at 12 weeks of infection. Splenic cell yields were comparable among knockout and controls. Representative scattergrams are shown.

**Figure 7.** Effect of CCR4 knockout on draining lymph node cell yields and cytokine production after *M. bovis* BCG infection. After i.t. infection, groups of mice were harvested at designated points and mediastinal draining lymph nodes were collected, counted, and prepared for culture with PPD antigen. After 48 hours, culture supernates were collected by centrifugation and assayed for cytokines by enzyme immunoassay. A: Total lymph node cell yields. The IFN-γ levels. C: The IL-17 levels. Bars are mean ± SD (n = 5–6 mice). *P < 0.05 versus the parallel control (CCR4+/+) group.

**CCR4 Knockout Mice Display an Impaired Adaptive Granuloma Recall Response**

As a live vaccine, BCG should establish long-term adaptive T-effector memory cells with the capacity to elicit a recall granulomatous response. Regarding mycobacterial infection, a hypersensitivity-type granulomatous response in the lungs is generally thought to represent a protective response that contains and destroys organisms. To test the role of CCR4 in the local recall response, we used a well-characterized model in which granulomas are elicited with mycobacterial Ag-coated beads. Unlike rechallenge with bacilli, this approach provides a highly sensitive indicator of inflammation through synchronization and control of lesion numbers. Briefly, mice were exposed i.t. to BCG; then, in the postelimination phase (20 weeks), mice were challenged i.v. with 6000 *M. bovis* PPD-coated beads to elicit synchronized granulomas. We first determined the induction of CCR4 agonist transcripts in lungs of PPD bead-challenged mice. As shown in Figure 8A, during recall granuloma formation,
mice per group). *BCG-sensitized CCR4

tification).

1 day; thereafter, they declined. CCL22 and CCL17 transcripts were induced in lungs by levels in naïve lungs. A representative study is shown.

Figure 8.

In view of rapid agonist induction, BCG-sensitized CD4+ effector/memory Th cells, the findings suggested that CCR4 was required for localization, activation, or survival of CD4+ effector memory T cells within the lung during the organ-based granuloma recall response.

Discussion

Initial optimism about identifying cellular subset-restricted chemokine receptors has been dampened as the complexity and pleomorphic nature of these receptors become more apparent. Yet, several chemokine receptors have clearly important physiological functions. Among these receptors, CCR4 was initially described as a Th2 cell restricted chemokine receptor.33,34 However, accumulating evidence indicates it has wider expression and function. In this vein, a functional role for CCR4 in a murine model of mycobacterial Ag-elicited granuloma formation was previously reported.3 In that study, it was demonstrated that CCR4 was expressed by mycobacterial Ag-elicited Th1 cells. Subsequently, Qiu et al35 demonstrated strong induction of CCR4 in lungs and draining lymph nodes of macaque monkeys infected with M. tuberculosis associated with IFNγ and IL-17 expression but not IL-4 expression. In the present study, we wanted to determine more specifically the functional role of CCR4 in the response to a live M. bovis BCG infection. Our findings are novel and point to a physiological role for CCR4 in supporting organ-based effector cell responses in innate and late stages of the antimycobacterial response.

We first established constitutive and induced CCR4 agonist expression in lungs after M. bovis BCG infection. The source of CCL22 was determined to be mainly large mononuclear cells in interstitium and inflammatory foci, consistent with the description of this chemokine as derived from macrophages and myeloid DCs.36 We then extensively tested the effect of CCR4 gene knockout on various aspects of the host response. Knockout mice displayed transiently impaired innate stage bacterial control, but the early adaptive clearance stage was unaffected. Thereafter, in the late stage when most bacteria were eliminated, CCR4−/− mice displayed reduced smoldering inflammation. These observations were particularly important because they suggested that although CCR4 was not required for effector cell induction, it appeared to support long-term inflammatory and homeostatic effector cell functions in peripheral organs. The presence of both constitutive and induced expression of CCR4 agonists is consistent with such a role.

Our subsequent analyses appeared to support this hypothesis and can be interpreted in the context of an evolving T-cell response. Although many aspects of the
induction of T-cell memory remain unclear, it is known that M. bovis BCG induces organ-based T effector/memory and long-lived lymphoid tissue homing central memory T cells. Our studies demonstrated the induction of effector T cells in draining lymph nodes by 2 weeks after BCG infection, followed by their disappearance. By using OVA-peptide–specific transgenic TCR CD4+ T-cell transfer, we confirmed that the CCR4 knockout environment could support primary adaptive T-cell proliferation. In addition, intracellular cytokine staining demonstrated no defect in the frequency of key effector cell populations within those lymph nodes. During the early adaptive elimination stage (2–4 weeks), effector cells were equivalently delivered to the infected lungs of CCR4+/+ and CCR4−/− mice. The latter was associated with rapid bacterial elimination, which was CCR4 independent. It was only in the late stage (≥6 weeks) that CCR4−/− mice displayed abrogated inflammation associated with reduced IFNγ– and IL-17–producing CD4+ T cells and an impaired recall granulomatous response in lungs. Our observations may relate to CD4+ T-effector/memory cell function at different stages of Ag burden. Specifically, in the early elimination stage, there are abundant Ag and costimulatory cytokines to activate effector cells. Although most effector cells are presumed to die in the target organ after activation, clearly a component persists beyond the early stage when bacteria were largely eliminated. This population was compromised in lungs of CCR4−/− mice. In the skin, CCR4 reportedly promotes development or maintenance of T-effector/memory cells. A similar situation may apply to the lung environment. Maintenance of T effector/memory in mycobacterial infection is purported to depend on low-level Ag persistence involving interactions with organ-based Ag-presenting cells. Moreover, Katou et al reported that CCR4 was required for optimal clustering of DCs and T cells in inflamed skin and lymphoid tissues. These reports are consistent with a recent study indicating that CCR4 expression was required by both APC and CD4+ T cells for optimal T-effector activation. It is reasonable to speculate that in the late elimination stage when Ag burden is low, T-effector/memory function becomes more dependent on constitutive CCR4-mediated interactions with DCs, which are rich sources of CCR4 agonists. Furthermore, the constitutive expression of CCR4 agonists in the lung may promote long-term steady-state T-cell–DC interactions, allowing local activation of tissue homing adaptive or innate effector cells when antigen is present. In addition to conventional TCRαβ+ T cells, M. bovis BCG is known to elicit a potent TCRγδ+ T-cell response in mice. Our finding that TCRγδ+ cytokine-producing T cells were unaffected in CCR4−/− mice is consistent with this population having different migratory and activation requirements than conventional CD4+ T-effector cells. Indeed, TCRγδ+ T cells express predominantly CCR5 and CCR9, respond directly to microbial products, and promote maturation of immature DCs. The TCRγδ+ T cells reached a maximum frequency in lungs at 2 weeks, were broadly activated, and were subsequently supplanted by conventional T cells, consistent with a role in a transition stage between innate and adaptive immunity.

Our finding of late stage appearance of PPD-responsive IL-17–producing cells in lymph nodes of M. bovis–infected mice was novel, and the abrogation of this population in CCR4−/− mice was unexpected. This finding suggests that CCR4 supports sustenance or activation of a lymph node homing memory population. Although preliminary studies indicate that this is a Th17 population, precise characterization regarding central versus effector memory phenotype has yet to be performed. Although we detected few IL-17+ effectors on day 2 of PPD-bead recall granuloma formation, compromise of a Th17 memory response might have relevance to the recall response. The Th17 cells are insufficient for mycobacterial resistance but have been reported to augment the Th1 antimycobacterial recall response. Detailed kinetic studies will be needed to determine whether compromise of Th17 memory cells in CCR4−/− mice contributed to the impaired Th1 mobilization in the late-stage PPD-bead granuloma recall response.

Finally, CCR4−/− mice displayed impaired mycobacterial control in the preadaptive stage of the response (0–2 weeks), which we hypothesized was related to abrogation of innate resistance. It is well established that several innate response mechanisms can control the early expansion of mycobacteria; among these mechanisms, NK cells are thought to be most influential. Indeed, we observed an early mobilization of NK cells to BCG-challenged lungs, reaching a maximum at 1 week and preceding the maximum appearance of TCRγδ+ CD8+, and CD4+ T cells. Although CCR4−/− mice showed no defect in absolute numbers of NK cells, they did display impaired NK activation, manifested by reduced IFNγ expression, providing an explanation for the observed defective innate resistance. Because CCR4 is expressed by NK cells and NK activation, like T-cell effector activation, is thought to involve local interactions with APCs, compromised NK function might be related to impaired interactions with organ-based DCs. Infection of mice with virulent M. tuberculosis overwhelms NK cell– and adaptive T-cell–mediated resistance. However, in humans, innate and adaptive resistance is more effective because only 20% of exposed individuals develop disease. It is conceivable that activated NK cells serve as an early source of IFNγ to promote bacterial killing within infected phagocytes. Based on our findings and published reports, we propose a unified hypothetical model in which CCR4 is not required for organ homing but rather mediates contacts of innate and memory effector cells with DCs in the lung interstitium. Such a scenario would provide a steady-state organ-based sentinel function in which innate-stage NK and late stage CD4+ T-memory/effector cells interact with DCs. The activation of available effectors would occur when DCs are triggered by pathogen-associated stimuli. In the setting of low levels of persisting Ag, such a model could provide a means to sustain chronic inflammation and support organ-based memory populations, also known as “effector lymphoid tissue.” If this model is substantiated, CCR4 would be a potential target for the treatment of chronic smoldering
inflammatory conditions and manipulation of antipathogen effector populations.

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