Neutrophils Clear Bacteria Associated with Parasitic Nematodes Augmenting the Development of an Effective Th2-Type Response

John T. Pesce, Zhugong Liu, Hossein Hamed, Farhang Alem, Jeanette Whitmire, Hongxia Lin, Qian Liu, Joseph F. Urban, Jr. and William C. Gause

http://www.jimmunol.org/content/180/1/464

References This article cites 48 articles, 23 of which you can access for free at: http://www.jimmunol.org/content/180/1/464.full#ref-list-1

Subscription Information about subscribing to The Journal of Immunology is online at: http://jimmunol.org/subscription

Permissions Submit copyright permission requests at: http://www.aai.org/About/Publications/JI/copyright.html

Email Alerts Receive free email-alerts when new articles cite this article. Sign up at: http://jimmunol.org/alerts
Neutrophils Clear Bacteria Associated with Parasitic Nematodes Augmenting the Development of an Effective Th2-Type Response

John T. Pesce, Zhungong Liu, Hossein Hamed, Farhang Alem, Jeanette Whitmire, Hongxia Lin, Qian Liu, Joseph F. Urban, Jr., and William C. Gause

Infection with the parasitic nematode Nippostrongylus brasiliensis induces a potent Th2 response; however, little is known about early stages of the innate response that may contribute to protective immunity. To examine early events in this response, chemokine expression in the draining lymph node was examined after N. brasiliensis inoculation. Pronounced increases of several chemokines, including CCL2, were observed. Compared with wild-type mice, elevations in a Gr-1 bright population in the draining lymph node was significantly decreased in CCL2−/− mice after N. brasiliensis inoculation. Further flow cytometric and immunofluorescent analysis showed that in wild-type mice, Gr-1 cells transiently entered and exited the draining lymph node shortly after N. brasiliensis inoculation. The Gr-1 bright population was comprised of neutrophils expressing TGF-β and TNF-α. Following Gr-1 cell depletion, N. brasiliensis infection resulted in transient, but significantly increased levels of IFN-γ, increased serum IgG2a, reduced Th2 cytokines and serum IgE, greatly increased mortality, and delayed worm expulsion. Furthermore, bacteria were readily detected in vital organs. Infection of Gr-1+ cell-depleted mice with N. brasiliensis larvae that were pretreated with antibiotics prevented bacterial dissemination, Th1 inflammatory responses, and decreases in host survival. This study indicates that parasitic nematodes can be an important vector of potentially harmful bacteria, which is typically controlled by CCL2-dependent neutrophils that ensure the optimal development of Th2 immune responses and parasite resistance. The Journal of Immunology, 2008, 180: 464–474.

Received for publication June 18, 2007. Accepted for publication October 18, 2007.

The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked advertisement in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

1 This work was supported by National Institutes of Health Grants AI031678 and AR066188.

2 J.T.P. and Z.L. contributed equally to this work.

3 Address correspondence and reprint requests to Dr. William C. Gause, Department of Medicine, New Jersey Medical School, University of Medicine & Dentistry of New Jersey, 185 South Orange Avenue, Newark, NJ 07101. E-mail address: gausewc@umdnj.edu

4 Abbreviations used in this paper: CLN, cervical lymph node; HPRT, hypoxanthine phosphoribosyltransferase; KO, knockout; WT, wild type.
were strongly chemotactic for immature dendritic cells (17). Parasite-triggered neutrophils are also known to produce TNF-α and induce dendritic cell CD40 and CD86 up-regulation (17). In another report, neutrophils were found to produce IL-4 early in BALB/c mice infected with *Leishmania major*, and were responsible for priming counterprotective Th2 cells (18). Innate cell populations activated following invasion of *N. brasiliensis* larvae may help the development of the protective Th2 adaptive response; however, the initial stages of this response remain largely uncharacterized.

In this investigation, we used quantitative fluorogenic real-time RT-PCR to assess elevations in gene expression of the Th2-type immune response in the draining lymph nodes early after *N. brasiliensis* inoculation. Our studies identified increased levels of CCL2 gene expression shortly after inoculation of wild-type (WT) mice, and an absence of Gr-1<sup>bright</sup> populations in the draining lymph node of *N. brasiliensis*-inoculated CCL2<sup>−/−</sup> mice. The Gr-1<sup>bright</sup> cells were identified as neutrophils that transiently entered the lymph node within 18 h after inoculation. Gr-1<sup>+</sup> cell depletion of *N. brasiliensis*-infected mice induced a pronounced early increase in IFN-γ expression, systemic bacterial infection, decreased host survival and Th2 cell differentiation, and delayed worm expulsion. Furthermore, treatment of infective L3 with an antibiotic mixture before inoculation eliminated associated bacteria and restored the protective Th2 response in anti-Gr-1 Ab-treated *N. brasiliensis*-inoculated WT mice. Our current study reveals a role for neutrophils in limiting inflammation and mortality from nematode parasite-associated bacterial infection and in the generation of optimal antimalnath Th2-type immune responses.

**Materials and Methods**

**Mice**

Female 6- to 10-wk-old BALB/c mice, C57BL/6, and CCL2<sup>−/−</sup> C57BL/6 mice were purchased from The Jackson Laboratory. All the mice were maintained in a specific pathogen-free, virus Ab-free facility during the experiments. Five mice were used per treatment group, if not otherwise indicated. The experiments in this study were conducted according to the principles set forth in the “Guide for the Care and Use of Laboratory Animals,” Institute of Animal Resources, National Research Council, Department of Health, Education, and Welfare (National Institutes of Health).

**Parasite infection and Gr-1 depletion**

Parasite eggs shed with the feces of *N. brasiliensis*-infected BALB/c mice were cultured in a mixture of charcoal and sphagnum moss stored in plastic petri dishes. The charcoal and moss culture was autoclaved before use to ensure the elimination of nonparasite-associated bacteria. Eggs hatch and develop from L1 through third stage (L3) infective larvae within 1 wk (19). Mice were inoculated intracutaneously in the ear with a 10<sup>5</sup> suspension of 500 L3 collected from cultures using a modified Baermann apparatus (19). The L3 were washed by centrifugation and replacement of sterile PBS three times before injection. Intracutaneous inoculation of L3 has been shown to produce the same level of infectivity as s.c. inoculation (3) (our unpublished observations). Parasite egg and adult worm numbers were evaluated in infected mice, as described previously (3). In several experiments, Gr-1<sup>bright</sup> cells were depleted in vivo by i.p. administration of anti-Gr-1 mAb (clone RB6-8C5) or isotype-matched control mAb (BD Biosciences) 1 day before larvae inoculation.

**Antibiotic treatment of infective *N. brasiliensis* larvae**

*N. brasiliensis* L3 were inoculated with 400 U of penicillin, 400 μg/ml streptomycin plus 400 μg/ml neomycin (Invitrogen Life Technologies) or sterile PBS for 2 h at room temperature, after which the larvae were washed three times with sterile PBS. Antibiotic-treated and untreated *N. brasiliensis* L3 were placed on blood agar plates (Fisher Scientific) and cultured at room temperature for 24 h. In some experiments, the antibiotic-treated larvae were used to inoculate BALB/c mice in the ear.

**Quantitative measurement of serum IgS**

Blood samples were collected 10 days after *N. brasiliensis* inoculation and processed to obtain serum. Total serum IgE, IgG1, and IgG2a levels were detected by ELISA (3).

**Gene expression by real-time PCR**

Total RNA was extracted from draining lymph nodes using RNAzol B (AMS Biotechnology), and total RNA was then reverse transcribed to cDNA using Superscript II (Invitrogen Life Technologies), as previously described (20). Real-time PCR Taqman (Applied Biosystems) kits were used for assessing IL-4, IFN-γ, IL-4R, IL-10, IL-13, TNF-α, and 18S ribosomal levels. Primers were designed for CCL2, TGF-β, and hypoxanthine phosphoribosyltransferase (HPRT), which were used to quantitate differences in gene expression. All data were normalized to 18S ribosomal or HPRT values. Primer sequences for SYBR Green assays were performed using the following primer pairs: CCL2-CTGTCCACCTCAAGTGGT (forward) and ATCTTGCTGGTGAATGAGTACGA (reverse), and HPRT-TCCTTGGAGATGTCTCAGGAGGA (forward) and AGCCAGTCGCAAAGAACTTATAGC (reverse), TGF-β (21). The Applied Biosystems 7700 sequence detector was used for amplification of target mRNA, and quantification of differences between treatment groups was calculated according to the manufacturer’s instructions.

**Cell sorting and cytokine gene expression by RT-PCR**

For sorting of Gr-1<sup>+</sup> cells, CLN cells were labeled with PE-conjugated anti-Gr-1 Ab (BD Pharmingen). After washing, the cells were applied to FACSVantage high-speed sorter, and Gr-1<sup>bright</sup> and Gr-1<sup>−</sup>cells were sorted and collected following standard protocol. For isolation of CD11c<sup>+</sup> cells, CLN cells were labeled with anti-CD11c microbeads and passed through LS<sup>+</sup> columns (Miltenyi Biotec). Purity of sorted cell populations was checked by FACS analysis. For RT-PCR, total RNA was extracted from purified cell populations with the RNA Isolation Kit (Stratagene), specially developed for isolating small RNA quantities. Total RNA was then reverse transcribed, as previously described. Real-time PCR kits (Applied Biosystems), specific for individual cytokines or rRNA, were used to quantitate differences in gene expression, and all data were normalized to constitutive rRNA values. Sorted cells were morphologically characterized on cytospin glass slides (Shandon Cytospin 4; Thermo Electron) after Wright-Giemsa staining (CAMCO).

**Flow cytometry**

Lymph node cells were harvested, and 1 × 10<sup>6</sup> cells were blocked with Fc Block (BD Pharmingen) and then incubated with anti-Gr-1-FITC, anti-CD11c-PE, and CD11b-PerCP-Cy5.5; anti-Gr-1-FITC, anti-MHC II-PE, and anti-B220-FITC; anti-Gr-1-PE, F4/80-FITC, and DX5-allophycocyanin; or anti-Gr-1-FITC and anti-CCR3-Alexa Fluor 647 (all from BD Pharmingen) for 30 min on ice. After washes, cells were fixed with 1% paraformaldehyde (Fisher Scientific) and data were collected by FACSCanilcub (BD Pharmingen). Expression of cell surface markers was analyzed on gated Gr-1<sup>bright</sup> cells or Gr-1<sup>−</sup>cells using WinList 5.0 software (Verity Software House).

**Immunofluorescence and digital microcopy**

Draining CLN were harvested from sacrificed mice and frozen in liquid nitrogen. Cryostat-cut tissue sections (8 μm) were fixed in acetone and stained, as described previously (3, 22), with the following reagents: FITC-conjugated anti-Gr-1 and Alex647-conjugated anti-B220 (BD Pharmingen). Sections were mounted in Fluormount G (Southern Biotechnology Associates) and viewed with a fluorescence microscope (Leica DMi6000B). Images were acquired on a digital camera and were processed with Image-Pro Plus software (Media Cybernetics).

**Results**

*N. brasiliensis* inoculation leads to rapid recruitment of Gr-1<sup>−</sup> cells within the draining lymph node, which is dependent on CCL2 signaling

Previous studies have shown that a highly polarized Th2 response develops in the draining CLN by day 7 after *N. brasiliensis* intracutaneous inoculation in the ear (3), with elevated IL-4 gene expression in the draining lymph node detected as early as day 3 after inoculation (5). Gene expression changes that precede increased IL-4 production in the draining lymph node were detected using quantitative fluorogenic real-time RT-PCR. Lymph nodes from *N. brasiliensis*-infected mice (5/treatment group) were individually collected for RNA analysis at days 1, 2, 3, and 6 after inoculation. As shown in Fig. 1A, elevations in CCL2 (2/5/treatment group) consistently
peaked in the draining CLN at day 1 after *N. brasiliensis* inoculation, and quickly dropped back to baseline. CCL2 is the ligand for CCR2 that is expressed on populations of granulocytes (23), dendritic cells (12), and also Th1 and Th2 cells (24, 25). It has also been shown to be responsible for recruitment of rat neutrophils in the absence of alveolar macrophages after endotoxin-induced injury (12). Recent studies have also suggested that CCL2 may be required for the development of an effective Th2 response in peripheral tissues, although its actual function is unclear (25–28). Further analysis of lymph node samples collected early after parasite inoculation with Affymetrix oligonucleotide microarrays confirmed up-regulation of this chemokine (our unpublished observations).

Recent studies have shown that the host-protective Th2 response to the intestinal nematode parasite, *Trichuris muris*, is substantially inhibited in CCL2 KO mice (25). Our finding that *N. brasiliensis* triggered increased levels of a chemokine in the lymph node known to recruit immune cells suggested that a nonresident immune cell population may migrate to the draining lymph node at the initiation of an antihelminth response. Identification of such cell population(s) may provide useful information concerning the innate response that influences the development of a protective Th2-type immune response induced by *N. brasiliensis* infection. A common characteristic of many of these different cell populations, including granulocytes (10), neutrophils (11), and subsets of dendritic cells (12, 13), is their expression of the cell surface marker Gr-1. We thus examined whether a Gr-1 bright cell population was recruited to the lymph node at the time of elevated CCL2 expression. In these studies, CCL2 knockout (KO) mice or BL/6 WT controls were inoculated with *N. brasiliensis* and, after 18 h when maximal CCL2 expression is observed, draining CLN were removed for analysis. As shown in Fig. 1B, in marked contrast to WT controls, CCL2KO BL/6 mice inoculated with *N. brasiliensis* showed a marked reduction in the recruitment of the Gr-1 bright cell subset to the draining lymph node. In contrast, the frequency of Gr-1 dull cells was not significantly affected. This result suggests that CCR2-directed Gr-1 bright cells infiltrate the draining lymph node after *N. brasiliensis* infection.

We next examined the dynamics of Gr-1 bright cell infiltration into the draining lymph node at early stages of the Th2-type in vivo immune response. To perform these studies, we collected the draining lymph nodes from *N. brasiliensis*-inoculated WT mice or untreated WT controls, and stained lymph node sections with fluorescence-conjugated anti-Gr-1 and anti-B220 Abs. Individual images were taken using a fluorescent microscope and tiled.
that of untreated controls (Fig. 1). Gr-1
reproducible in multiple independent experiments. We also ex-
treated Gr-1 untreated BALB/c mice were used as
the subcapsular space, sinuses, and interfollicular areas (Fig. 1)
lymph node sections from untreated mice. In contrast, by 4 h
after inoculation, Gr-1 cells had accumulated in large numbers in
the subcapsular space, sinuses, and interfollicular areas (Fig. 1).
The number of Gr-1 cells in the draining lymph node at
24 h was much lower than at 18 h after inoculation, and by 48 h
the level of Gr-1 cells in the lymph node was comparable to
that of untreated controls (Fig. 1C). These results were highly
reproducible in multiple independent experiments. We also ex-
amined Gr-1 cell migration into other peripheral lymph nodes
following inoculation with N. brasiliensis. Gr-1 cells migrated
into lymph nodes draining the site of inoculation, but not
peripheral lymph nodes distant from the site of infection (data not
shown).

**FIGURE 2.** LN-infiltrating Gr-1bright cells are TNF-α and TGF-β-expressing neutrophils. A, BALB/c mice were
inoculated with N. brasiliensis L3, and 18 h later CLN cells were harvested and stained with anti-Gr-1-PE, F480
FITC, anti-CD11b-PerCP-Cy5.5, anti-CD11c-allophycocyanin, anti-CCR3-Alexa Fluor 647, or anti-DX5
allophycocyanin. B, In separate experiments, BALB/c mice (30/treatment group) were inoculated with N. brasili-
ensis L3, and 18 h later draining CLN were removed. Pooled cell suspensions were stained with anti-Gr-1 Ab. Gr-
1 bright and dull cells were isolated using high speed cell sorting (FACS). Sorted cells were centrifuged on slides using a
Cytospin and then stained with DiffQuick to analyze morphology. C, RNA was purified from sorted cell
populations, and real-time quantitative fluorogenic RT-PCR was used to assess cytokines and IL-4R gene ex-
pression. Sorted CD11c cells from untreated BALB/c mice were used as controls, and data are expressed as
treated Gr-1 cells/untreated CD11c cells. These experiments were repeated twice with similar results.

Lymph node-infiltrating Gr-1bright cells are neutrophils that express TNF-α and TGF-β

We used flow cytometric analysis to determine the cell lineage of the Gr-1 population that infiltrated the draining lymph node after N. brasiliensis infection. At 18 h after N. brasiliensis inoculation, single-cell suspensions were prepared from the draining CLN of WT mice and stained with fluorochrome-conjugated Abs against an array of cell lineage markers. Forward and side scatter analysis of Gr-1bright cells revealed a characteristic granulocyte phenotype (data not shown). As shown in Fig. 2A, the majority of cells staining Gr-1bright also showed bright staining for CD11b, which is a β2 integrin component that is highly expressed by neutrophil and macrophage populations. Further analysis of gated Gr-
1brightCD11bhbright cells revealed a largely homogeneous population that was primarily negative for CCR3 (eosinophils), DX5 (basophils), CD11c (dendritic cells), and F4/80 (macrophages). Gr-1bright cells were electronically sorted by a high-speed FACSVantage cell sorter. Sorted cells were cytopun on glass slides and stained with Wright/Giems. The stained Gr-1bright cells showed a polymorphonuclear phenotype with clear cytoplasm (Fig. 2B). Taken together, the phenotype analysis of Gr-1bright cells indicated a homogenous population of neutrophils (29).

CD11bhGr-1dull cells were moderately increased in the draining lymph node at 18 h after parasite inoculation, and were more heterogeneous with 16.8% CCR3+ cells, 42.5% CD11c+, and 48.1% F4/80+ cells. Consistent with this cell surface phenotype, electronic cell sorting and Wright/Giems staining of Gr-1dull cells revealed two major populations of dendritic cells/macrophages and eosinophils (Fig. 2A). Taken together, these data indicate that the Gr-1dull cells are composed of a heterogeneous population, including eosinophils, dendritic cells, and macrophages. Migration of Gr-1dull cells to the draining lymph node seems independent of CCL2 expression, because a similar number of Gr-1dull was detected in parasite-infected CCL2KO mice (Fig. 1B).

To further characterize the LN-infiltrating Gr-1+ cells, RNA was isolated from the sorted Gr-1bright and Gr-1dull cells at 18 h after N. brasiliensis inoculation and reverse transcribed into cDNA. Cytokine gene expression by these two cell populations was assessed by quantitative fluorogenic real-time RT-PCR. Because the Gr-1+ cell number in untreated LN is too low for cell sorting, CD11c+ dendritic cells were purified from LN of untreated animals. Cytokine gene expression levels by sorted Gr-1+ populations were compared with the levels of naive CD11c dendritic cells. As shown in Fig. 2C, Gr-1bright cells express no or minimal levels of effector cytokines, including IL-4, IFN-γ, IL-12, IL-13, and IL-6. The only two cytokines that were up-regulated in Gr-1bright cells after N. brasiliensis infection were TNF-α and
TGF-β. Expression of TNF-α by lymph node-infiltrating neutrophils was reported previously (29). In direct contrast, the Gr-1<sup>dull</sup> cells revealed no or very low levels of TNF-α and TGF-β expression; high levels of IL-4, IFN-γ, and IL-6; and lesser increases in IL-13 and IL-10 (Fig. 2C). Our findings now show for the first time that neutrophils infiltrate the draining lymph node transiently after helminth parasite inoculation.

**Neutrophils prevent rapid immune cell activation and IFN-γ elevations following N. brasiliensis inoculation**

To further assess the possible function of Gr-1 neutrophils in the Th2-type immune response induced by *N. brasiliensis* infection, we used an anti-Gr-1 Ab to deplete neutrophils before *N. brasiliensis* inoculation. Because neutrophil infiltration of draining lymph nodes peaked at 18 h postinfection, we selected this time point to collect the draining lymph nodes and further assessed cell activation and cytokine expression. Anti-Gr-1 Ab treatment effectively depleted the Gr-1<sup>bright</sup> population, as demonstrated by FACS analysis of lymph node cells, but left the Gr-1<sup>dull</sup> population unaffected (data not shown). To evaluate the cell activation status, lymph node cells were stained with anti-B220, anti-CD11c, and anti-CD4 or anti-CD8 plus anti-CD69, anti-B7.2, anti-MHC II, or anti-CD40 Abs and analyzed by FACS. Expression of cell surface costimulatory molecules was assessed on gated B220<sup>+</sup> B cells and CD11c<sup>+</sup> dendritic cells. Expression of CD69 was examined on gated CD4<sup>+</sup> T cells and CD8<sup>+</sup> T cells. This experiment was repeated twice with similar results.

A, Single-cell suspensions were stained with anti-B220, anti-CD11c, and anti-CD4 or anti-CD8 plus anti-CD69, anti-B7.2, anti-MHC II, or anti-CD40 Abs and analyzed by FACS. Expression of cell surface costimulatory molecules was assessed on gated B220<sup>+</sup> B cells and CD11c<sup>+</sup> dendritic cells. Expression of CD69 was examined on gated CD4<sup>+</sup> T cells and CD8<sup>+</sup> T cells. This experiment was repeated twice with similar results. B, In a separate experiment, the collected CLN were further processed for RNA isolation and analyzed for IFN-γ gene expression using real-time quantitative fluorogenic RT-PCR. All data are expressed as relative to that of untreated controls, and the mean and SE are shown for each treatment group (5 mice/group). This experiment was repeated three times with similar results. C, BALB/c mice were administrated either with anti-Gr-1, anti-TGF-β, anti-TNF-α, or anti-TGF-β plus anti-TNF-α Abs, and 1 day later inoculated in the ear with *N. brasiliensis* L3. Gene expression of IFN-γ was analyzed by real-time quantitative fluorogenic RT-PCR. This experiment was repeated twice with similar results.
molecules on accessory cells after Gr-1 cell depletion correlated with increased activation of both CD4+ and CD8+ T lymphocytes. As shown in Fig. 3A, expression of the early activation marker CD69 increased markedly on CD4+ and CD8+ T cells after N. brasiliensis inoculation of anti-Gr-1 Ab-treated mice compared with isotype Ig-treated mice. These results indicate that the presence of Gr-1 cells in the N. brasiliensis-infected hosts is required to prevent a general activation of accessory cells and lymphocytes.

The increased activation of immune cells in the draining lymph nodes after N. brasiliensis infection in Gr-1-depleted mice may be associated with altered effector functions. Based on this, we were interested in examining the levels of effector cytokines at this early time point after N. brasiliensis inoculation. As shown in Fig. 3B, 18 h after N. brasiliensis infection, IFN-γ mRNA levels did not change in isotype-treated mice compared with untreated controls. However, N. brasiliensis infection induced increased IFN-γ expression after neutrophil depletion. Gene expression of IL-4 remained low in all groups at this time point (data not shown).

Neutrophils have been shown to express proinflammatory cytokines, including TNF-α, that can modify the Ag-specific T cell response (29, 30). TGF-β is one of the effector cytokines of T regulatory cells and was shown to suppress both Th1 and Th2 adaptive responses (31). Because the major cytokines produced by Gr-1 bright neutrophils in this system were TGF-β and TNF-α, we next examined whether these cytokines inhibited IFN-γ expression after N. brasiliensis infection. To address this question, neutralizing Abs were used to block TGF-β and TNF-α bioactivity in vivo in mice inoculated with N. brasiliensis larvae, and IFN-γ expression in the draining lymph node was assessed 18 h later (Fig. 3C).

Blocking TGF-β or TNF-α activity alone or together failed to increase IFN-γ expression in the draining lymph node similar to that observed in anti-Gr-1 Ab-treated and N. brasiliensis-infected mice. In addition, the cell surface expression of activation markers was not increased (data not shown). Thus, the increased immune cell activation and IFN-γ expression induced by N. brasiliensis after anti-Gr-1 Ab treatment were not due to the absence of TGF-β or TNF-α produced by neutrophils.

In the absence of neutrophils, N. brasiliensis infection results in significant mortality and impaired antihelminth Th2-type responses

The Th2-type immune response against N. brasiliensis infection is typically protective. Adult worms are expelled from the gut ≤2 wk after inoculation. After depletion of Gr-1 bright neutrophils, N. brasiliensis inoculation resulted in a rapid and dramatic increase in IFN-γ expression. We hypothesized that this elevated IFN-γ expression during the early time point after N. brasiliensis infection
could affect the development of the antihelminth Th2 immune response. Furthermore, because this Th2 response is critical for host protection, the absence of neutrophils might affect worm expulsion as well. To verify these possibilities, we monitored neutrophil-depleted mice after N. brasiliensis infection. By 48 h postinfection, a significant number of neutrophil-depleted mice died, and the mortality rate reached 60–80% by 72 h postinoculation (Fig. 4A). Several other neutrophil depletion experiments were performed in which a lower dose of larvae (300 instead of 500 L3) was administered and similar mortality rates were observed (data not shown). The surviving mice gradually recovered and were healthy at the end of these experiments (day 10 postinfection). In direct contrast, all isotype control Ab-treated N. brasiliensis-inoculated mice were healthy and survived. A slight inflammatory response was observed in the infection site (ear) of the isotype Ab-treated mice that resolved after day 3 postinfection (Fig. 4B); however, the inflammatory response at the infection site was much more severe in anti-Gr-1 Ab-treated mice. Areas of ear necrosis were observed in all of the Gr-1<sup>−/−</sup> cell-depleted mice 3 days post-N. brasiliensis inoculation (Fig. 4B). The mice that were found dead in anti-Gr-1 Ab-treated groups were taken for necropsy (Research Animal Diagnostic Laboratory). A prominent Gram-negative rod bacterium colonized many tissues, including the lungs, liver, spleen, and CLN. Areas of cellular necrosis accompanied bacterial colonization in the spleen, liver, and CLN.

Mesenteric lymph nodes were collected from surviving mice at day 7 after N. brasiliensis inoculation, and processed to assess cytokine expression levels. As shown in Fig. 4C, significantly deceased levels of IL-4 expression were detected in N. brasiliensis-inoculated neutrophil-depleted mice compared with isotype IgG-treated mice; IL-13 expression levels were similarly down-regulated. Although Th2 cytokine expression was decreased after Gr-1 depletion, the immune response as detected on day 7 did not skew in a Th1 direction because IFN-γ expression was low (Fig. 4C, right panel). Similarly, draining CLN IL-4 expression on day 7 after N. brasiliensis infection was significantly decreased in mice administered anti-Gr-1 Ab with no elevation in IFN-γ expression (data not shown).

Consistent with the decreased Th2 response, worm burden and egg counts on day 10 were markedly increased in anti-Gr-1 Ab-treated mice (Fig. 4D). The Th2 response after N. brasiliensis infection is associated with significantly increased serum IgE levels. This IL-4-dependent serum IgE was decreased after anti-Gr-1 Ab treatment (Fig. 4E). Significantly increased levels of IFN-γ-dependent serum IgG2a were detected in N. brasiliensis-inoculated Gr-1-depleted mice compared with controls, providing confirmation of physiologically significant increased levels of IFN-γ. These data indicate that Gr-1<sup>−/−</sup> neutrophils play critical roles in host protection against N. brasiliensis infection.

The selective inhibition of neutrophil recruitment to the draining lymph node of N. brasiliensis-inoculated CCL2KO mice provided the opportunity to confirm findings obtained with anti-Gr-1 Ab treatment. Draining CLN were removed from CCL2KO or control WT mice at 18 h after N. brasiliensis inoculation, and analyzed for Th1/Th2 cytokine mRNA. As shown in Fig. 5A, IFN-γ was markedly up-regulated in N. brasiliensis-inoculated CCL2-deficient BL/6 mice, but not in inoculated WT BL/6 mice by 18 h post-inoculation. In contrast, IL-4 mRNA was not elevated in either CCL2-deficient or WT mice at this early time after N. brasiliensis inoculation. In addition, significantly increased levels of IFN-γ-dependent serum IgG2a were detected in N. brasiliensis-inoculated CCL2-deficient mice compared with inoculated WT controls (Fig. 5B). Furthermore, egg burden markedly increased in CCL2-deficient mice (Fig. 5B). These results are consistent with our previous data, obtained with anti-Gr-1 Ab treatment, suggesting that neutrophils are required for host protection and optimal Th2-type responses after N. brasiliensis infection.

Control of larvae-associated bacteria by neutrophils is critical for host defense and optimal Th2 response development

The findings of necrosis at the infection site and of systemic bacterial spreading in neutrophil-depleted N. brasiliensis-inoculated mice prompted us to investigate the influence of bacterial infection on the host response. It should be noted that we used standard N. brasiliensis propagation techniques, with charcoal and moss autoclaved before use. The larvae-associated bacteria most likely originate from feces in which the eggs developed. Bacteria are most likely ingested by free-living L3 and are also probably associated with their integument. Infective N. brasiliensis L3, along with any associated bacteria, penetrate (or are injected through) the host skin, enter the blood circulation, and 24–48 h later migrate to the lung. After neutrophil depletion, larvae-associated bacteria may spread quickly in the host. To test this hypothesis, we first examined whether bacteria are associated with N. brasiliensis infective larvae. Infective N. brasiliensis L3 were prepared following standard protocols. Five hundred larvae were washed with sterile PBS and then used to inoculate blood agar plates. After overnight culture at 37°C, large numbers of bacterial colonies were recorded (Fig. 6A, left panel). Another 500 N. brasiliensis L3 were treated with a mixture of antibiotics (penicillin plus streptomycin plus neomycin) for 2 h at room temperature. After several washes with sterile PBS, these larvae were inoculated onto another blood agar plate. Few, if any, bacteria colonies were observed on the agar plate after overnight culture in this condition (Fig. 6A, right panel).

Because in vitro treatment with antibiotics could essentially eliminate the larvae-associated bacteria, we next examined the host response to antibiotic-treated N. brasiliensis L3 following administration of anti-Gr-1 Ab. Eighteen hours after ear inoculation, the draining CLN was removed and assessed for cytokine gene expression. Levels of IL-4 mRNA were unchanged in all groups at this early time point (data not shown). We have previously shown the results obtained with anti-Gr-1 Ab treatment, suggesting that neutrophils are required for host protection and optimal Th2-type responses after N. brasiliensis infection.
that inoculation of neutrophil-depleted mice with untreated *N. brasiliensis* L3 significantly increased immune cell activation and IFN-γ expression as detected at 18 h (Fig. 3, A and B). In direct contrast, after antibiotic mixture treatment, *N. brasiliensis* L3 failed to induce expression of immune cell surface activation markers (data not shown) or any elevation of IFN-γ expression in the lymph node (Fig. 6B).

Anti-Gr-1 Ab-treated mice inoculated with untreated *N. brasiliensis* L3 exhibit a markedly increased mortality rate (see Fig. 4A). We next tested whether the larvae-associated bacteria similarly contributed to the death of anti-Gr-1 Ab-treated mice inoculated with *N. brasiliensis*. L3 treated with or without antibiotics were inoculated in the ear of BALB/c WT mice 1 day after anti-Gr-1 Ab administration. Although 60–80% of anti-Gr-1 Ab-treated mice died after infection with untreated L3, none of the anti-Gr-1 Ab-treated mice that were infected with antibiotics-treated L3 showed any sign of illness, and all of them survived (Fig. 6C). In addition, there was no tissue necrosis at the infection site (ear) of the mice inoculated with antibiotics-treated larvae (Fig. 6D). It should be noted that Gr-1+ cells still migrated into the lymph node with a...
similar composition and frequency following injection with antibiotic-treated L3 (data not shown).

Because antibiotic-treated \textit{N. brasiliensis} larvae did not induce a rapid IFN-\gamma response in the anti-Gr-1 Ab-treated host, we next examined whether antibiotic-treated larvae could dampen the protective Th2 response. Inoculation with antibiotic-treated L3 induced a potent IL-4 and IL-13 response in the draining lymph nodes following anti-Gr-1 Ab treatment, similar to that observed in isotype-treated mice inoculated with L3 (mesenteric lymph node, Fig. 6E; CLN, data not shown). In contrast, IFN-\gamma gene expression was not elevated on day 7 after \textit{N. brasiliensis} inoculation. Furthermore, the potent Th2 response that developed in the presence of sterilized larvae was protective with all worms expelled around 2 wk and no eggs present in the feces after inoculation (data not shown). Thus, host protection was sustained following neutrophil depletion in mice infected with antibiotic-treated \textit{N. brasiliensis} L3. These results suggest that the primary function of neutrophils during \textit{N. brasiliensis} infection is to clear larvae-associated bacteria and provide an environment conducive to the development of a Th2-like immune response.

Discussion

\textit{N. brasiliensis} induces one of the most potent and highly polarized Th2-type responses known. However, few studies have examined characteristics of the innate immune response that might influence the adaptive host-protective T cell response. In this study, we found that, shortly after \textit{N. brasiliensis} inoculation, CCL2-dependent neutrophils transiently entered the lymph node, and that depletion of this population led to increased IFN-\gamma expression, mortality, decreased Th2 responses, and delayed worm expulsion. Treatment of infective larvae with antibiotics before inoculation abrogated the requirement of neutrophils, suggesting that neutrophils play a significant role in clearing bacteria associated with parasitic nematode infections, which would otherwise suppress the development of a host-protective Th2-type response.

By examining time points directly after inoculation with \textit{N. brasiliensis}, we were able to identify multiple innate associated immune components, which were up-regulated before the development of IL-4-producing Th2 cells. Of particular interest, we noted the rapid accumulation in the draining lymph node of Gr-1\textsuperscript{bright} neutrophils. Neutrophils comprise about two-thirds of peripheral blood leukocytes and are among the first cells that migrate to sites of inflammation, where they perform effector functions, including phagocytosis and the secretion of cytotoxic compounds. Early studies suggested that neutrophils are transcriptionally inactive due to their terminal differentiation; however, more recent studies have indicated that they are active components of the innate immune response (16). In addition, it has been suggested that neutrophils might carry pathogens/Ags to the lymphoid organs (29, 32, 33). This does not appear to be the case in \textit{N. brasiliensis}-infected animals, because inoculation with larvae plus fluorescence-labeled protein Ag indicated that most of the lymph node-infiltrating neutrophils were not associated with the inoculated Ag (data not shown), although the distribution pattern of neutrophils in lymph nodes after \textit{N. brasiliensis} infection is similar to previously reported data (29, 32, 33).

Neutrophils are also involved in host immunity to parasite infection. They are one of the first cell populations recruited to the site of \textit{T. gondii} infection and are able to release proinflammatory cytokines and chemokines (including TNF-\alpha and CCL2) (34, 35), which in turn contribute to recruitment, maturation, and activation of dendritic cells that can then drive the generation of the Th1-type immune response (17, 36). Although high levels of TGF-\beta and TNF-\alpha were observed in lymph node-infiltrating neutrophils early after \textit{N. brasiliensis} inoculation, our findings suggest that these cytokines did not have a significant impact on the development of the potent Th2-type response induced by \textit{N. brasiliensis} inoculation.

Previous studies using various neutrophil depletion techniques have observed severe immunomodulatory consequences after exposure to fungal, protozoan, or nematode pathogens. Using CXCR2\textsuperscript{−/−} mice, infection with \textit{T. gondii} results in increased parasite burden and diminished Th1-associated immune factors (37), whereas the immune response to infection with the fungal agent \textit{Histoplasma capsulatum} is significantly impaired in the absence of Gr-1\textsuperscript{+} neutrophils (38). Granulocytes directly affect \textit{Strongyloides stercoralis} parasitic L3 housed in diffusion chambers in vivo (39), and neutrophil depletion before infection resulted in increased numbers of migrating larvae and egg production by day 5 after primary inoculation of \textit{Strongyloides ratti} (40). Our findings indicate that during the immune response to \textit{N. brasiliensis}, neutrophils control parasite-associated bacteria, which would otherwise impair the host-protective Th2-type immune response. When interpreting data showing regulatory function(s) for specific Gr-1\textsuperscript{+} cell populations, it is thus important to consider whether bacterial depletion by neutrophils may be an important modulating component of the immune response.

Our findings further showed that bacteria associated with \textit{N. brasiliensis} initiate a potentially lethal bacterial infection following neutrophil depletion, which is most likely not observed in the \textit{Strongyloides} model because parasites in this system are treated with antibiotics (40). Our data showed that antibiotic-treated or untreated larvae similarly induce Th2 immunity in mice with an intact neutrophil population. We interpret this as indicating that neutrophils play an essential role in clearing helminth-associated bacteria, which otherwise would induce a Th1-type response in the absence of neutrophils, thereby impairing the development of an effective Th2-type response. The importance of neutrophils in eliminating \textit{N. brasiliensis}-associated bacteria was demonstrated by significantly increased mortality after anti-Gr-1 Ab treatment. This finding may be important during natural \textit{Strongyloides}, or other similar parasite infections in immunocompromised patients.

As discussed in Results, the \textit{N. brasiliensis} larvae-associated bacteria most likely originate from feces in which the eggs develop. Intestinal nematode larvae naturally develop under similar conditions, and it is thus likely that bacteria are also associated with parasitic nematodes during natural infection. Our experiments with the DO11.10 adoptive transfer system have been established with the intracutaneous ear injection inoculation model, partially because it induces a localized readily detectable response in the draining ear lymph node, while exhibiting a similar infection route and time course to s.c. inoculation. The process of skin penetration during natural infection may result in loss of bacteria to some degree. Further studies using different protocols, including studies in human parasitic nematode infections, are required to verify the significance of our observations.

Recent studies have shown that the host-protective Th2-type response to the intestinal nematode parasite, \textit{Trichuris muris}, is substantially inhibited in CCL2\textsuperscript{−/−} mice (25). We observed that the migration of neutrophils to the draining lymph nodes following inoculation with \textit{N. brasiliensis} is CCL2 dependent. This is consistent with a previous study in which neutralization of CCL2 blocked airway accumulation of neutrophils (41). Although neutrophils failed to migrate to the draining lymph node in CCL2KO mice, they were capable of eradicating the bacteria, because the infected CCL2KO mice did not show increased mortality or ear necrosis (data not shown). It may be that, whereas neutrophils lack the ability to migrate to the lymph node in response to CCR2
signaling, the overall neutrophil population is still intact within the mouse and capable of controlling systemic bacterial infection, as opposed to anti-Gr-1 treatment, which systemically depletes the mice of neutrophils. It is important to note, however, that the inability of neutrophils to enter the lymph node early after infection did lead to significantly increased levels of IFN-γ production. Thus, CCL2-dependent rapid migration of neutrophils in response to infectious agents proves to be an important factor suppressing an alternative Th1-type response.

It should be noted that in the absence of neutrophils, N. brasiliensis inoculation resulted in greatly enhanced mortality despite marked and rapid elevations in IFN-γ production, suggesting that in the absence of neutrophils, the adaptive Th1-type response alone is not sufficient to effectively control bacterial infection. The early increased expression of IFN-γ in neutrophil-depleted mice also affected the N. brasiliensis-induced Th2 immune response. On day 7, IL-4/IL-13-expressing cells developed, but at a lower level, suggesting that early IFN-γ expression suppresses the optimal development of Th2-type immunity, leading to increased egg number and delayed worm expulsion. Our findings are supported by a previous study using exogenously administered IL-12 and anti-IFN-γ Ab, which suggested that Th1 cytokines can inhibit host protection against N. brasiliensis infection (42). The presence of neutrophils ensures an environment for optimal Th2 development by eliminating parasite-associated bacteria that otherwise induce a Th1-type response. Recently, similar studies suggest that Gr-1+CD11b+ cells, which most likely include neutrophils, suppress Th1 responses while inducing Th2 polarization (43).

Our findings also suggest that, even following the early and pronounced Th1-type response resulting from neutrophil depletion, an active N. brasiliensis-dependent process ultimately drives a polarized, albeit attenuated, Th2-type response. This is not consistent with the default hypothesis, which postulates that a Th2-polarized process ultimately drives a Th1-type response, leading to bacterial infection (47, 48). Our results are similar, but also show the importance of innate immune components during N. brasiliensis infection that eliminate pathogenic bacteria exclusive of the pronounced adaptive Th1-type response. Our findings suggest that neutrophil recruitment to the lymph node prevents a bacterial-induced Th1-type response that would interfere with a host-protective Th2-type response to worm infection.

In summary, the current study suggests a previously unknown essential role for neutrophils at early stages of nematode parasite infections. Our findings suggest that clearance of larvae-associated bacteria by neutrophils is critical for wound healing, host protection, and optimal development of the adaptive Th2-type response leading to worm expulsion. In addition to increased mortality, failure to control bacterial spreading significantly affects the developing Th2 immune response and delays worm expulsion. This may be of particular significance in immune compromised individuals infected by helminths or in the development of vaccines that may require stimulation of both Th1-type and Th2-type components of the response to protect against parental helminth infection.

Disclosures

The authors have no financial conflict of interest.

References


