The aim of this study was to determine the role of CC chemokine CCL6/C10 in acute inflammation. Intraperitoneal injection of thioglycollate increased peritoneal CCL6, which peaked at 4 h and remained elevated at 48 h. Neutralization of CCL6 significantly inhibited the macrophage infiltration (34–48% reduction), but not other cell types, without decreasing the other CC chemokines known to attract monocytes/macrophages. CCL6 was expressed in peripheral eosinophils and elicited macrophages, but not in elicited neutrophils. Peritoneal CCL6 level was not decreased in granulocyte-depleted mice where eosinophil influx was significantly impaired. Thus, CCL6 appears to contribute to the macrophage infiltration that is independent of other CC chemokines. Eosinophils pre-store CCL6, but do not release CCL6 in the peritoneum in this model of inflammation.

Key words: Acute inflammation, CC chemokines, Eosinophils, Monocytes/macrophages, Neutrophils

Introduction

The inflammatory response is characterized by leukocyte infiltration into inflammatory foci, which is initiated and orchestrated by a variety of inflammatory mediators. Of these, chemokines are a group of cytokines that are chemotactic for specific types of leukocyte populations.1,2 Chemokines have been divided mainly into two subfamilies, CXC and CC chemokines, based on their sequence homology and the position of cysteine residues.3,4 CXC chemokines are typically chemotactic for neutrophils, whereas CC chemokines attract and activate monocytes and lymphocytes. The role of CXC chemokine such as CXC8/IL-8 in a variety of inflammatory diseases has been extensively investigated.5 CC chemokines that include CCL2/MCP-1, CCL3/MIP-1α and CCL22/MDC are detected in an early phase of acute inflammation, playing an essential role in the recruitment and activation of monocytes/macrophages.6–10 Murine CCL6/C10, a prototype CC chemokine, was initially identified as a transcript induced in bone marrow cells upon stimulation with granulocyte–macrophage colony-stimulating factor.11 It has shown that CCL6 is present in a variety of chronic inflammatory disorders including experimental demyelinating diseases, allergic airway inflammation, bleomycin-induced lung fibrosis and chronic peritonitis.12–15 These findings demonstrate an important role of CCL6 in chronic inflammation associated with macrophage infiltration. However, the precise role of CCL6 in acute inflammation remains to be elucidated.

The aim of the present study was to characterize the role of CCL6 in acute inflammation. For this purpose, we first examined the production kinetics of CCL6 in a murine model of acute peritonitis induced by thioglycollate injection. We then investigated whether CCL6 could contribute to the leukocyte infiltration in this model by neutralizing the endogenous CCL6 with antibodies against CCL6. Finally, we attempted to determine the major cellular sources of CCL6 using reverse transcription-polymerase chain reaction (RT-PCR), enzyme-linked immunosorbent assay (ELISA) and immunohistochemistry. The present data suggest that CCL6 is an important CC chemokine attracting monocytes/macrophages. Although CCL6 is pre-stored in circulating eosinophils, the CCL6 does not appear to be released in this particular model.

Materials and methods

Mice

Female CD-1 mice (6–8 weeks of age) were purchased from Jackson Laboratory (Bar Harbor, ME, USA).
and housed in the animal care facility unit (University Laboratory of Animal Medicine). The animal use committee at the University of Michigan approved all studies.

Peritonitis model

The mice received intraperitoneal (i.p.) injection of 1 ml of a 2% thioglycollate broth (Difco Laboratories, Detroit, MI, USA). To neutralize CCL6, 0.5 ml of anti-murine CCL6 antiserum was i.p. injected 2 h prior to thioglycollate injection. Polyclonal anti-murine CCL6 antiserum was raised by immunizing rabbits with recombinant murine CCL6 (R&D systems Inc., Minneapolis, MN, USA), as previously described.13,16 The antibodies had a neutralizing activity against murine recombinant CCL6 in vitro chemotaxis assay, and did not cross-react with a number of other murine cytokines and chemokines available (data not shown). As a control, pre-immune rabbit serum (0.5 ml) was used. At appropriate time points following thioglycollate injection, mice were euthanized by cervical dislocation. Peritoneal lavages were collected by washing the peritoneum with 2 ml of sterile saline, cervical dislocation. Peritoneal lavages were collected by washing the peritoneum with 2 ml of sterile saline, centrifuged, and cell-free peritoneal fluids were were centrifuged, and cell-free peritoneal fluids were stored at −20°C for measurements of chemokines. Cell pellets were resuspended in saline, and the cell numbers were counted in a hemocytometer. Differential cell analysis was made after Diff-Quik staining of cytospin preparations.

Granulocyte depletion

Granulocytes were depleted using RB6-8C5 monoclonal antibody directed against Ly-6G, previously known as Gr-1, an antigen on the surface of murine granulocytes that includes neutrophils and eosinophils.17,18 A total of 100 µg of RB6-8C5 was i.p. administered 1 day prior to thioglycollate injection. This resulted in peripheral blood granulopenia (< 50 cells/µl) by days 1 and 3 after the administration, with a return of peripheral counts to pretreatment levels by day 5.19 Due to the granulopenia, the treatment effectively inhibited neutrophil and eosinophil infiltration in animal models of inflammation.19,20 Iso-type-matched mouse IgG was used as a control.

Isolation of granulocytes and macrophages

Elicited granulocytes and macrophages were isolated from the peritoneal exudates after thioglycollate injection by Percoll gradient centrifugation. For this, exudates from five mice or two mice were combined in each experiment for granulocyte or macrophage isolation, respectively. Six independent experiments were performed. Peripheral granulocytes were isolated from heparinized untreated mice blood (10 mice each, three independent experiments) by dextran sedimentation, followed by Percoll gradient centrifugation and hypotonic lysis of erythrocytes. Diff-Quik staining of cytospin preparations showed that the purity of granulocytes and macrophages in each experiment was > 92% and > 94%, respectively. Resident peritoneal macrophages were harvested from untreated mice, and the adherent cells were used. Isolated peritoneal macrophages were used for RT-PCR and ELISA. For ELISA assay, cells were extracted with phosphate-buffered saline containing 1% Triton X-100 and protease inhibitors (Roche Applied Science, Basel, Switzerland).

RT-PCR

Total RNA was isolated from cells using Trizol reagent (Life Technologies, Grand Island, NY, USA) according to the manufacturer’s instructions. First-strand cDNA was synthesized from 1 µg of total RNA with oligo (dT)12–18 as primers, and the first-strand cDNAs were then amplified by each PCR in the presence of Taq polymerase (Life Technologies) and specific primers. The primers were designed to amplify murine CCL6 referred to the cDNA sequence from the National Center for Biotechnology Information database. The primers were as follows: CCL6 sense, 5'-GACGTCAATGCGGCTGATAC-3' and antisense, 5'-TCGTOGATCTGGCTCG-3'; GADPH sense, 5'-GGTGAAGGTGCTGT-3' and antisense, 5'-GATGGC-CAAGTTGGTCAATGGATCC-3'. The PCR reaction was conducted at 35 cycles of 94°C for 1 min, 55°C for 1 min, and 72°C for 1 min. Ten microlitres of each PCR product was subjected to electrophoresis on a 2% agarose gel in the presence of ethidium bromide.

ELISA

Murine chemokines were quantitated using a standard method of sandwich ELISA as previously described.13,16,21 The captured antibodies, detection antibodies and the recombinant cytokines were purchased from R&D Systems. The ELISAs employed in this study did not cross-react with other murine cytokines, and constantly detected murine cytokines/chemokines above 10 pg/ml.

Immunohistochemistry

Cytospin preparations were fixed in 100% ethanol for 10 min, and the endogenous neutrophil peroxidase was blocked with 0.3% H2O2 in methanol. The slides were rehydrated in Tris-buffered saline (TBS) and blocked with 5% normal goat serum for 20 min at room temperature. Staining for CCL6 was carried out with anti-CCL6 IgG or control rabbit IgG at 5 µg/ml in TBS–1% bovine serum albumin for 30 min at room temperature. After washing with TBS–Tween 20
(0.1%), the slides were incubated with biotinylated goat anti-rabbit IgG antibodies and rinsed. The slides were then incubated for 30 min with Vectastain Elite ABC Reagent (Vector Laboratories, Burlingame, CA, USA). After washing, the reaction was developed with DAB Substrate Kit (Zymed Laboratories, San Francisco, CA, USA). Counter-staining was done with hematoxylin.

Statistics
Statistical significance was evaluated by an unpaired Student’s t-test. $p < 0.05$ was regarded as statistically significant. All results are expressed as mean ± standard error.

Results
Production of CCL6 during thioglycollate-induced peritonitis
To determine the production kinetics of CCL6 in this model of acute inflammation, cell-free peritoneal fluids were collected at appropriate time points after thioglycollate injection and CCL6 level in the fluids were measured by ELISA. As shown in Fig. 1, levels of CCL6 reached their peak at 4 h and the levels were sustained for 48 h after the injection. Thus, CCL6 increased rapidly in the peritoneum and the level remained elevated even after 48 h post-injection.

Neutralization of endogenous CCL6 reduces macrophage infiltration
Experiments were next conducted to determine the role of CCL6 in this model. To address this, endogenous CCL6 was neutralized by administration of anti-CCL6 antiserum prior to thioglycollate injection. The data in Fig. 2 demonstrate that neutralization of CCL6 resulted in significant decreases in the number of infiltrating macrophages at any time examined, ranging from a 34% reduction to a 48% reduction (43% reduction on average), as compared with the control. The treatment did not affect the numbers of infiltrating neutrophils, eosinophils, and lymphocytes (Fig. 2). These results indicate that endogenous CCL6 is important in the recruitment of macrophages, but not other cell types.

Effects of anti-CCL6 antiserum on the production of other CC chemokines
To determine whether CCL6 could directly or indirectly attract macrophages during inflammation, we examined the peritoneal levels of CCL2, CCL3 and CCL22 known to be involved in the recruitment of macrophages after the treatment with anti-CCL6 antibodies. Consequently, the treatment did not decrease the CCL2 level, but rather augmented CCL2 production, as compared with the control. The CCL22 level was not altered by the treatment (Fig. 3). No appreciable level of CCL3 was detected in the peritoneum (data not shown). Thus, other monocyte chemotactic CC chemokines were not
decreased by neutralization of CCL6, indicating that CCL6 may attract monocytes/macrophages independently of other CC chemokines. Anti-CCL6 treatment failed to alter the peritoneal level of CCL1/TCA3, CCL5/RANTES and CCL11/Eotaxin (data not shown).

Cellular source of CCL6

We next attempted to determine the cellular source of CCL6. First, we isolated peritoneal granulocytes and macrophages following thioglycollate injection, and mRNA expression in the cells was examined by RT-PCR. Non-stimulated granulocytes and macrophages were isolated from peripheral blood and the peritoneum from untreated mice, respectively. As shown in Fig. 4, CCL6 mRNA expression was not detected in elicited granulocytes at 4 h (mean% eosinophils = 4.4%) and 8 h (mean% eosinophils = 13.7%) after the injection. No expression was found at 24 h (mean% eosinophils = 13.7%; data not shown). In contrast, not only CCL6 mRNA (Fig. 4), but also its protein (Fig. 5) were detected in peripheral granulocytes (mean% eosinophils = 24%), and the protein level in the elicited granulocytes was gradually decreased as compared with peripheral granulocytes (Fig. 5). On the other hand, both CCL6 mRNA and protein were detected in resident macrophages, and the levels in the elicited macrophages were increased with time during inflammation (Figs 4 and 5). Immunohistochemically, the cells stained with anti-CCL6 IgG were eosinophils and macrophages. Neutrophils were not stained with the anti-CCL6 IgG (Fig. 6).

The aforementioned findings suggest a possibility that eosinophils may contribute to the early phase of macrophage influx via releasing the pre-stored CCL6 inside the cells. To address this hypothesis, granulocytes were depleted from mice and peritoneal levels...
of CCL6 in the granulocyte-depleted mice were then measured. Depletion of granulocytes resulted in significant decreases in the number of infiltrating neutrophils and eosinophils at 4 h and 8 h after thioglycollate injection (Table 1). Under these conditions, the peritoneal CCL6 levels were measured, which demonstrated that levels of CCL6 were comparable with those in the control (Fig. 7). These findings suggest that CCL6 in eosinophils is not released in the peritoneum in this model.

Discussion

Inflammation is a series of well-coordinated events that depend on sequential arrival of inflammatory leukocytes to the site of inflammation, in which chemokines play a central role. Our present data suggest that CCL6 is unique in its commitment to the development of inflammation. Production of CCL6 peaked at 4 h and the level remained elevated even at 48 h after the injection. Wu et al. demonstrated that high level of CCL6 was detected for up to 10 days in this model. In contrast, CCL2 and CCL22, known to attract monocytes/macrophages, peaked at 4 h after the injection and returned to the basal level by 24 h (own unpublished data). Neutralization of CCL6 resulted in decreases in the number of infiltrating macrophages during the observation periods, but not other cell types, suggesting that CCL6 governs macrophage infiltration from the very beginning to a late period of time, playing a role in the initiation and maintenance of macrophage infiltration.

Macrophage infiltration is induced by multiple CC chemokines, leading to a possibility that CCL6 might induce macrophage infiltration indirectly via the production of other CC chemokines known to be involved in the recruitment of macrophages. A recent investigation has demonstrated that CCL6 neutralization decreased IL-13-induced CCL2 and CCL3 production in the lung. Neutralization of CCL6 decreased CCL2 production in a murine model of allergic bronchopulmonary aspergillosis. CCL6 enhanced CCL2 production in a murine model of septic peritonitis. However, the present data demonstrated that neutralization of CCL6 did not decrease CCL2 as well as CCL22 levels, indicating that CCL6 may attract monocytes/macrophages without reference to other CC chemokines in this particular model. Thus, chemokine–chemokine interaction appears to be different among the types of inflammation.

Interestingly, the CCL2 level was rather increased by neutralization of CCL6. The mechanism by which neutralization of CCL6 increased CCL2 in this model remains to be elucidated. CCL6 is induced by anti-inflammatory cytokines interleukin (IL)-4, IL-10 and IL-13, raising a possibility that CCL6 may down-regulate the production of CCL2. However, CCL6 (1–100 ng/ml) failed to decrease the production of CCL2.

Table 1. Thioglycollate-induced leukocyte infiltration in granulocyte-depleted mice

| Time | Treatments   | Neutrophils | Eosinophils | Lymphocytes | Macrophages |
|------|--------------|-------------|-------------|-------------|-------------|
| 4 h  | RB6-8C5 mAb  | 25 ± 0.15   | 0.03 ± 0.01*| 0.72 ± 0.17 | 3.67 ± 0.68 |
|      | Control IgG | 4.57 ± 0.98 | 0.17 ± 0.06 | 0.65 ± 0.12 | 3.68 ± 0.60 |
| 8 h  | RB6-8C5 mAb  | 0.82 ± 0.26*| 0.14 ± 0.08*| 0.71 ± 0.16 | 6.03 ± 0.87*|
|      | Control IgG | 5.02 ± 0.96 | 0.42 ± 0.10 | 0.83 ± 0.14 | 9.03 ± 0.95 |

Mice were treated with RB6-8C5 monoclonal antibody (mAb) or control IgG, 1 day prior to thioglycollate injection. At 4 h and 8 h after thioglycollate injection, mice were euthanized and the infiltrating leukocytes in the peritoneum were counted (10 mice for each time-point).
* p <0.05, ** p <0.001 versus control.
FIG. 7. Peritoneal CCL6 level in granulocyte-depleted mice. Granulocytes were depleted using RB6-8C5 monoclonal antibody. As a control, isotype-matched mouse IgG was used. At indicated intervals after thioglycollate injection, mice were euthanized and the peritoneal fluids were harvested (each group, 10 mice). CCL6 levels in the peritoneal fluids were measured by ELISA.

In summary, we have demonstrated that CCL6 is an important mediator in the recruitment of macrophages in acute inflammation. Thus, the present data together with those in chronic inflammation models suggest that CCL6 may have function in both acute and chronic inflammatory responses. Acute inflammation is tightly linked to T-cell-mediated chronic inflammation, suggesting a broad activity of CCL6 in the development and maintenance of inflammatory disorders.

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References

1. Luster AD. Chemokines—chemotactic cytokines that mediate inflammation. N Engl J Med 1998; 338: 436–445.
2. Feng L. Role of chemokines in inflammation and immunoregulation. Inflamm Res 2000; 21: 203–210.
3. Baggioili M. Chemokines and leukocyte traffic. Nature 1998; 392: 565–568.
4. Rot A, Von Andrian UH. Chemokines in innate and adaptive host defense: basic chemokine grammar for immune cells. Annu Rev Immunol 2004; 22: 891–928.
5. Gerard C, Rollins BJ. Chemokines and disease. Nat Immunol 2001; 2: 108–115.
6. Matsukawa A, Miyazaki S, Maeda T, et al. Production and regulation of macrophage chemotactic protein-1 (MCP-1) in lipopolysaccharide- or monosodium urate crystal-induced arthritis in rabbits: roles of tumor necrosis factor alpha, interleukin-1, and interleukin-6. Lab Invest 1998; 78: 973–985.
7. Matsukawa A, Hogaboam CM, Lukacs NW, Lincoln PM, Strieter RM, Kunkel SL. Endogenous monocyte chemotactant protein-1 (MCP-1) protects mice in a model of acute septic peritonitis: cross-talk between MCP-1 and leukotriene B4. J Immunol 1999; 163: 6149–6154.
8. Matsukawa A, Hogaboam CM, Lukacs NW, Lincoln PM, Evanoff HL, Kunkel SL. Proximal role of the CC chemokine, macrophage-derived chemokine, in the innate immune response. J Immunol 2000; 164: 5362–5368.
9. Ajetor MN, Flower RJ, Hannon R, Christie M, Bowers K, Vertay A, Perretti M. Endogenous monocyte chemotactant protein-1 recruits monocytes in the zymosan peritonitis model. J Leukoc Biol 1998; 63: 108–116.
10. Ajetor MN, Das AM, Virag L, Szabo C, Perretti M. Regulation of macrophage inflammatory protein-1 alpha expression and function by endogenous interleukin-10 in a model of acute inflammation. Biochem Biophys Res Commun 1999; 255: 279–282.
11. Orlofsky A, Berger MS, Prystowsky MB. Novel expression pattern of a new member of the MIP-1 family of cytokine-like genes. Cell Regulat 1991; 2: 403–412.
12. Belperio JA, Dy M, Burdick MD, Xue YY, Li K, Elias JA, Keane MP. Interaction of IL-13 and C10 in the pathogenesis of bleomycin-induced pulmonary fibrosis. Am J Respir Cell Mol Biol 2002; 27: 419–427.
13. Hogaboam CM, Gallinat CS, Tash DB, Strieter RM, Kunkel SL, Lukacs NW. Immunomodulatory role of C10 chemokine in a murine model of allergic bronchopulmonary aspergillosis. J Immunol 1999; 162: 6071–6079.
14. Asensio VC, Lassmann S, Pagtenstecher A, Steffensen SC, Henricksen SJ, Campbell IL. C10 is a novel chemokine expressed in experimental inflammatory demyelinating disorders that promotes recruitment of macrophages to the central nervous system. Am J Pathol 1999; 154: 1181–1193.
15. Wu Y, Prystowsky MB, Orlofsky A. Sustained high-level production of murine chemokine C10 during chronic inflammation. Cytokine 1999; 11: 523–530.
16. Steinhauser ML, Hogaboam CM, Matsukawa A, Lukacs NW, Strieter RM, Kunkel SL. Chemokine C10 promotes disease resolution and survival in an experimental model of bacterial sepsis. Infect Immun 2000; 68: 6108–6114.
17. Hedstal K, Russetti FW, Bile JN, et al. Characterization and regulation of BB8-8C5 antigen expression on murine bone marrow cells. J Immunol 1991; 147: 22–28.
18. Hamaguchi-Tsutsu E, Nobumoto A, Hirose N, et al. Development and functional analysis of eosinophils from murine embryonic stem cells. Br J Haematol 2004; 124: 819–827.
19. Tateda K, Moore TA, Deng JC, et al. Early recruitment of neutrophils determines subsequent T1/T2 host responses in a murine model of *Legionella pneumophila* pneumonia. *J Immunol* 2001; 166: 3355–3361.

20. Singer M, Lefort J, Vargaftig BB. Granulocyte depletion and dexamethasone differentially modulate airways hyperreactivity, inflammation, mucus accumulation, and secretion induced by rmIL-13 or antigen. *Am J Respir Cell Mol Biol* 2002; 26: 74–84.

21. Evanoff HL, Burdick MD, Moore NA, Kunkel SL, Strieter RM. A sensitive ELISA for the detection of human monocyte chemoattractant protein-1 (MCP-1). *Immunol Invest* 1992; 21: 39–45.

22. Ma B, Zha Z, Hornet BJ, Gerard C, Strieter R, Elias JA. The C10/CCL6 chemokine and CCR1 play critical roles in the pathogenesis of IL-13-induced inflammation and remodeling. *J Immunol* 2004; 172: 1872–1881.

23. Orlofsky A, Wu Y, Prystowsky MB. Divergent regulation of the murine CC chemokine C10 by Th(1) and Th(2) cytokines. *Cytokine* 2000; 12: 220–228.

24. Lacy P, Moxbel R. Eosinophil cytokines. *Cytokine* 2000; 12: 134–155.

25. Oliveira SH, Tauls DD, Nagel J, et al. Stem cell factor induces eosinophil activation and degranulation: mediator release and gene array analysis. *Blood* 2002; 100: 4291–4297.

26. Melnicoff MJ, Horan PK, Morahan PS. Kinetics of changes in peritoneal cell populations following acute inflammation. *Cell Immunol* 1989; 118: 178–191.

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