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Identification of human CCR8 as a CCL18 receptor

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The CC chemokine ligand 18 (CCL18) is one of the most highly expressed chemokines in human chronic inflammatory diseases. An appreciation of the role of CCL18 in these diseases has been hampered by the lack of an identified chemokine receptor. We report that the human chemokine receptor CCR8 is a CCL18 receptor. CCL18 induced chemotaxis and calcium flux of human CCR8-transfected cells. CCL18 bound with high affinity to CCR8 and induced its internalization. Human CCL1, the known endogenous CCR8 ligand, and CCL18 competed for binding to CCR8-transfected cells. Further, CCL1 and CCL18 induced heterologous cross-desensitization of CCR8-transfected cells and human Th2 cells. CCL18 induced chemotaxis and calcium flux of human activated highly polarized Th2 cells through CCR8. Wild-type but not Ccr8-deficient activated mouse Th2 cells migrated in response to CCL18. CCL18 and CCR8 were coexpressed in esophageal biopsy tissue from individuals with active eosinophilic esophagitis (EoE) and were present at markedly higher levels compared with esophageal tissue isolated from EoE patients whose disease was in remission or in normal controls. Identifying CCR8 as a chemokine receptor for CCL18 will help clarify the biological role of this highly expressed chemokine in human disease.
inflammatory reactions and to tissue remodeling and fibrosis in chronic inflammatory diseases. Concordant with the spectrum of AAM activity, the abundance of CCL18 has been found to correlate with disease severity in fibrotic diseases, such as pulmonary fibrosis and scleroderma, and diseases of dysregulated macrophage biology (Schutyser et al., 2005; Prasse et al., 2007, 2009). In allergic diseases, increased CCL18 is also frequently associated with eosinophil infiltration in affected tissues (Pivarcsi et al., 2004; Schutyser et al., 2005; de Nadaï et al., 2006; Lucendo et al., 2011).

CCL18 activity has been detected on peripheral blood lymphocytes (Adema et al., 1997; Hieshima et al., 1997). However, studies using cells transfected with CCR1-7, CXCR1-4, and CXCR1-4 did not reveal a specific CCL18 signaling receptor but did find that CCL18 could antagonize CCR3 (Hieshima et al., 1997; Nibbs et al., 2000). Investigation into atopic dermatitis and allergic diseases enabled the first identification of a specific T cell subset that responded to CCL18, and thus provided clues to its possible chemokine receptor (Günther et al., 2005; de Nadaï et al., 2006). In atopic dermatitis, CCL18 is the most abundantly expressed chemokine in lesional skin (Pivarcsi et al., 2004; Günther et al., 2005). CCL18 bound to blood skin-tractable cutaneous leukocyte antigen expressing (CLA+) memory T cells from individuals with atopic dermatitis and induced migration of CD4+ T cell clones derived from atopic dermatitis lesional skin in vitro and into human skin transplanted in SCID mice in vivo (Günther et al., 2005). The signature homing receptors of skin-tractable T cells are CLA in combination with the chemokine receptors CCR4, CCR10, and CCR8 (Schaerli et al., 2004; Islam et al., 2011). More CCR8-expressing cells are found in inflamed skin of individuals with active atopic dermatitis compared with skin of healthy individuals (Gombert et al., 2005). In studies of human asthma, CCL18 was found to induce the migration of TCR-activated in vitro differentiated human Th2 cells (de Nadaï et al., 2006). CCL18 induced migration of human peripheral blood Th2 cells and regulatory T cells in ex vivo (Bellinghausen et al., 2012; Chenivesse et al., 2012). Th2 but not Th1 cells are programmed to selectively express the chemokine receptors CCR4 and CCR8, a receptor profile shared with regulatory T cells (D’Ambrosio et al., 1998; Wei et al., 2010). Skin-homing CLA+ T cells, Th2 cells, and regulatory T cells thus all express CCR4 and CCR8. However, TCR activation is a prerequisite for functional CCR8 but not CCR4 expression on in vitro differentiated Th2 cells (D’Ambrosio et al., 1998). Here, we report that CCL18 is an endogenous agonist of the human CCR8 receptor.

RESULTS AND DISCUSSION

CCL18 induced migration of CCR8-transfected cells

We assayed the ability of 4DE4 mouse pre-B cell lines transfected with human CCR8 to chemotax to CCL18 and observed peak migration at 10 nM (Fig. 1 a). Untransfected 4DE4 cells exhibited migration to CXCL12 (not depicted), but not to CCL18 (Fig. 1 a) or the known human CCR8 chemokine agonist CCL1 (also called 1-309; Fig. 1 b). CCL1 also induced a peak migration of CCR8 transfectants at 10 nM but with a more robust response (Fig. 1 b). CCL18-induced chemotaxis in CCR8-transfected cells was inhibited by pertussis toxin (PTX), indicating that CCL18 induced the specific coupling of CCR8 to Gac (Fig. 1 c). Checkerboard-type chemotaxis analysis revealed that CCL18 stimulated chemotaxis rather than chemokinesis (Fig. 1 d). CCL18 did not antagonize chemotaxis of CCR8-transfected cells to CCL1 when both chemokines were present in the bottom chamber in transwell chemotaxis assays (Fig. 1 e).

CCL18 induced calcium flux in CCR8-transfected cells

CCR8 transfectants showed a rapid and robust dose-dependent increase in intracellular calcium upon stimulation with CCL18 (Fig. 2 a). In control experiments, CCL18 did not induce a calcium flux in untransfected 4DE4 cells, although they responded to CXCL12 with a robust calcium flux (Fig. 2 b). Heterologous desensitization of ligand-induced calcium flux is routinely used to assess activity at a shared receptor. Stimulation of CCR8 transfectants with CCL18 inhibited, albeit incompletely, subsequent signaling to CCL1, reflecting...
CCL18-induced partial desensitization of the CCR8 receptor (Fig. 2 c). Stimulation of CCR8 transfectants with CCL1 completely inhibited subsequent signaling to CCL18 (Fig. 2 c). We then assayed the ability of CCL18 to desensitize another well-characterized agonist of the human CCR8 receptor, HHV-8 (human herpes virus 8)–encoded chemokine viral MIP (vMIP) 1 (Dairaghi et al., 1999). vMIP–1–induced calcium signaling was partially desensitized by CCL18, and vMIP-I completely desensitized CCL18–induced signaling (Fig. 2 d). We also assayed the L1.2 murine pre–B cell line stably transfected with human CCR8 for chemotaxis and calcium flux to CCL18 but found that they responded weakly to CCL18 despite responding to CCL1. We do not have a clear explanation for the discrepancy in the magnitude of CCL18 responsiveness between L1.2 and 4DE4 CCR8 transfectants but consequently pursued all our functional studies on 4DE4 CCR8 transfectants.

CCL18 promoted internalization of CCR8

Chemokine receptors typically undergo internalization upon chemokine agonist binding. Robust surface expression of CCR8 on CCR8–transfected cells was detected at baseline but not on untransfected 4DE4 cells (Fig. 3 a). Surface CCR8 expression on CCR8–transfected cells was reduced after treatment with increasing concentrations of CCL18 or CCL1 but not CCL17 (a CCR4 but not CCR8 agonist) for 20 min at 37°C (Fig. 3 b). CCL18 and CCL1 binding to CCR8–transfected cells at 4°C did not affect mAb binding to CCR8. To demonstrate that CCL18 induced CCR8 internalization, we followed the kinetics of surface, total, and internalized (total – surface) CCR8 levels in CCR8–transfected cells after treatment with CCL18 or CCL1 (10−8 M). After treatment with CCL18 or CCL1, surface CCR8 levels decreased contemporaneously with increased internalized CCR8 levels, whereas total CCR8 levels were not affected by either treatment (Fig. 3 c).

CCL18 binds to CCR8

Competitive ligand binding experiments with 125I–labeled CCL18 and 125I–labeled hCCL1 enabled us to determine specific
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CCL18 induced migration and calcium flux of highly polarized Th2 cells

CCR8 induction in murine Th2 cells at levels high enough to detect agonist function occurs transiently after TCR activation of Th2 cells generated by multiple rounds of polarization (D’Ambrosio et al., 1998; Islam et al., 2011). Human Th2 cells also exhibited increased CCR8 expression with successive rounds of Th2 polarization (Fig. 4a) with a concomitant increase in IL-4 and IL-5 expression (Fig. 4b). Highly polarized human Th2 cells generated by three rounds of polarization with subsequent TCR activation (Th2 R3) migrated in response to CCL18 and CCL1, whereas less polarized Th2 cells that expressed lower levels of CCR8 did not (Fig. 4c). Peak migration for both CCL18 and CCL1 was at 10 nM, similar to what was observed on CCR8−transfected cells. CCL18-induced chemotaxis of Th2 R3 cells and untreated controls. (e) Activated human Th2 R3 cells were treated with 1 μg/ml CCR8 mAb and compared with untreated activated Th2 R3 cells in chemotaxis assay to CCL18. a–e are representative of at least three independent experiments (mean ± SEM). (f) Dose−dependent calcium flux response of human Th2 R3 cells to CCL18 stimulation. (g) Heterologous cross-desensitization of the calcium flux response of human Th2 R3 cells to CCL18 and hCCL1 stimulation. f and g are representative of three to five separate experiments. (h) Representative histograms of surface expression of the LFA-1 activation epitope m24 on activated human Th2 R3 cells treated with CCL18, CXCL12, PMA, or not treated (left). Quantitation of m24 expression on activated Th2 R3 cells after CCL18 or CXCL12 stimulation (right). (j) Chemotaxis of activated wild-type and Ccr8−deficient mouse Th2 cells to CCL18. (j) Chemotaxis of mCcr8−transfected and untransfected Ba/f3 cells to CCL18. h–j are representative of at least three independent experiments (mean ± SEM; *, P < 0.05 by unpaired two-tailed Student’s t test for d, e, and h–j).
CCL18 activates mouse CCR8
Activated highly differentiated wild-type mouse Th2 cells migrated to CCL18 in a dose-dependent bell-shaped migration curve with peak migration at 1 nM (Fig. 4 i). Migration was abrogated in Ccr8-deficient Th2 cells in which migration to the positive control CXCL12 remained intact and comparable to wild-type Th2 cells (not depicted). Moreover, mCcr8-transfected but not untransfected Baf/3 murine B cells migrated in response to CCL18 with peak migration at 10 nM (Fig. 4 j). The peak migration induced by CCL18 was comparable to the peak migration induced in these cells by mCCL1 and the recently identified second chemokine ligand for murine CCR8, mouse CCL8 (mCCL8 or mouse monocyte chemotactic protein [MCP] 2; Islam et al., 2011).

CCL18 induces LFA-1 activation
Chemokine signaling can induce “inside-out” activation of integrins. In T cells, chemokine activation of LFA-1 results in a conformational change that can be detected by mAb24, which recognizes the appearance of a new LFA-1 epitope m24 (Schürpf and Springer, 2011). CCL18 induced expression of the m24 epitope, detected by mAb24, which was comparable to that induced by CXCL12 or PMA on activated highly polarized Th2 cells at 37°C. This increase in mAb24 binding was not detected at 4°C or in the absence of stimulation (Fig. 4 h).

Human CCL18 and murine CCL8 are functional analogues
Although mCCL8 was originally given the orthologous designation to human CCL8 (hCCL8), we recently established they are not functional orthologues. hCCL8 is not a CCR8 ligand but is a CCR2 ligand, whereas mCCL8 is a CCR8 ligand but...
is not a CCR2 ligand (Islam et al., 2011). Synten mapping of mCCL8 and hCCL8, which are located in the mouse and human MCP chemokine clusters, respectively, has also revealed they are not orthologues (Zlotnik and Yoshie, 2012). In fact, mCCL8 lacks a human orthologue, and both hCCL8 and CCL18 lack rodent orthologues (Homologene, MGI).

This has led us to hypothesize that mCCL8 and CCL18 may be functional analogues. Consistent with this hypothesis, CCL18 and mCCL8 have been reported to be highly induced in AAMs (Kodelja et al., 1998; Schutyser et al., 2005; Thomas et al., 2012; Egawa et al., 2013). Additionally, mCCL8 was the most abundant chemokine induced in the lung in a bleomycin model of pulmonary fibrosis by microarray analysis (Liu et al., 2011), and CCL18 is the most highly expressed chemokine in idiopathic pulmonary fibrosis (Prasse et al., 2009).

We thus determined if cytokines known to regulate CCL18 induction might also regulate mCCL8 induction in AAMs in an analogous fashion. As expected, human CCL18 and another human Th2-associated chemokine CCL17 were both induced in human AAMs in response to IL-4, but CCL18 was induced to much greater levels (Fig. 5 a). Consistent with published studies (Kodelja et al., 1998), we found that IL-10 weakly induced CCL18 in macrophages. However, as noted by others, a combination of IL-4– and IL-10 synergistically induced CCL18 (Pechkovsky et al., 2010), which is in distinct contrast to IL-4 and TNF (Fig. 5 b). In contrast, IL-10 treatment did not induce CCL17 and instead inhibited IL-4–induced CCL17 expression in AAMs, whereas TNF and IL-4 synergistically induced CCL17 (Fig. 5 b).

In mouse AAMs, we found that Gdβ8 and Cd1l7 were also induced by IL-4. AAM differentiation was confirmed by the specific induction of the mouse AAM markers Arg1, Ym1, and Fizz1 (Fig. 5 c). In a striking parallel to CCL18 induction in human AAMs, IL-4 and IL-10 both independently induced Gdβ8, and importantly, IL-4 and IL-10 in combination, but not IL-4 and TNF, synergistically induced Gdβ8 expression (Fig. 5 d). In contrast, regulation of Cd1l7 induction in mouse AAMs mirrored that of human CCL17 in human AAMs and was distinct from that of CCL18 and Gdβ8. Thus, IL-10, an inhibitory cytokine, paradoxically amplified IL-4 induction of both Gdβ8 and CCL18. These data provide evidence that CCL18 and Gdβ8 are regulated similarly in AAMs, which lends further support to the hypothesis that mCCL8 and CCL18 are functional analogues.

CCL18 and CCR8 are induced in human eosinophilic inflammation

Previously, we found that CCR8 and mCCL8 were essential for the induction of chronic eosinophilic skin inflammation in vivo (Islam et al., 2011). To investigate whether CCL18 and its receptor CCR8 are involved in chronic human eosinophilic inflammation, we examined esophageal biopsy tissue of patients with eosinophilic esophagitis (EoE; Table S1) for expression of Th2-associated chemokines (CCL18, CCL1, and CCL26) and receptors (CCR8, CCR3, and CRTH2). CCL18 and CCR8 mRNA were up-regulated in subjects with active EoE compared with subjects whose disease was in remission and to normal controls (Fig. 5 e). CCL1, CCL26, CCR3, and GRTH2 mRNA were also up-regulated in active EoE. Moreover, CCL18 and CCL1 mRNA levels in esophageal tissue correlated strongly with CCR8 mRNA levels (Fig. 5 f). In contrast, the eosinophil chemoattractant CCL26, which is known to be abundantly expressed in EoE (Blanchard et al., 2007) and is a CCR3 ligand, was not as highly correlated with CCR8 (Fig. 5 f). Thus, CCL18 and its receptor CCR8 are coexpressed in human diseased tissue during active eosinophilic inflammation, consistent with our findings in the mouse.

Phylogeny of CCL18

CCL18 has orthologues only in primates (Homologene). Sequence analysis revealed that CCL18 shares the most amino acid identity (65%) with hCCL3 (MIP-1α), which lacks activity on CCR8. It is hypothesized that the CCL18 gene, which is located within the MIP chemokine gene cluster, arose through duplication and fusion of two CCL3-like genes with deletion and selective use of exons after the speciation of primates and rodents (Schutyser et al., 2005). Consistent with this hypothesis, vMIP-1, the potent virally encoded CCR8 agonist also believed to have arisen from the MIP gene cluster, is more homologous to CCL18 (46% identity) than it is to the other CCR8 agonist hCCL1 (25% identity). Alignment of the mature amino acid sequences of the CCR8 agonists CCL18, CCL1, and vMIP-1 revealed that only CCL18 and vMIP-I share a continuous stretch of 20-aa identity. The human and mouse MIP and MCP chemokines are closely linked gene clusters on human chromosome 17 and mouse chromosome 11. Inflammatory cluster chemokines often tend to evolve rapidly, and tandem gene duplication and diversification are believed to have occurred within the MCP and MIP clusters independently after the speciation of rodents and primates (Zlotnik and Yoshie, 2012). This may explain why these two chemokine clusters are not orthologous between humans and mice and why mouse CCL8 and human CCL18 lack identified orthologues by phylogenetic analysis and synteny mapping. Instead, functional analogues need to be identified based on similar biological function, such as similar regulation and receptor usage, as we have demonstrated for human CCL18 and mouse CCL8.

Our study shows that CCR8 has two endogenous chemokine ligands. Ligands for the same chemokine receptor often have distinct yet cooperative and complementary functions in vivo (Zlotnik and Yoshie, 2012). In the CCR8 system, CCL1 appears to be the more potent agonist. However, CCL18 appears to be more highly expressed in inflammation. CCL18 is detected at two log higher levels than CCL1 in lesional atopic skin, which is analogous to the two log higher expression of mCCL8 compared with mCCL1 in inflamed atopic mouse skin (Pivarcsi et al., 2004; Gombert et al., 2005; Islam et al., 2011). While CCL18 is expressed in dermal dendritic cells and keratinocytes, CCL1 is expressed by epidermal Langerhans cells and skin endothelial cells (Gombert et al., 2005).
CCL1 may therefore promote entry through the cutaneous vasculature while CCL18 may promote retention of CCR8-expressing cells in inflamed tissue. Thus, we predict that CCL1 and CCL18 likely have nonredundant roles in vivo in part dictated by their pattern of expression and their activity on CCR8-expressing cells.

In certain human cancers (e.g., breast and ovarian), tumor-associated macrophages, an AAM-like population which undergoes additional differentiation by T reg cell–derived IL-10, abundantly produce CCL18 (Bonecchi et al., 2011). It is notable that the induction of both CCL18 and mCCL8 is markedly augmented by IL-10. In human breast cancer, CCL18 expression in blood and cancer stroma was associated with reduced patient survival (Chen et al., 2011). CCL18 promoted tumor cell invasiveness and metastasis, which was shown to be mediated by PITPNM3 (phosphatidylinositol transfer protein 3; Chen et al., 2011). PITPNM3 is a six-transmembrane receptor distinct from seven-transmembrane chemokine receptors (Bonecchi et al., 2011). We did not detect PITPNM3 RNA or protein expression in our CCR8-transfected 4DE4 B cell line, whereas we did detect PITPNM3 RNA and protein in the MDA-MB-231 epithelial breast cancer cell line (unpublished data). Therefore, the CCL18-induced CCR8 activation that we described in transfected cells and primary T cells was independent of PITPNM3.

Identifying CCR8 as a receptor for CCL18 offers new insights into the role of this important human chemokine identified in many chronic inflammatory diseases. In a recent transcription profiling study, CCR8 was found to be the chemokine receptor most highly expressed in induced T reg cells (Wei et al., 2010). Recruitment of CCR8–expressing T reg cells into an IL–10–rich tumor milieu may possibly explain the association of CCL18 with increased metastasis and poor prognosis. The ability of CCL18 to recruit CCR8–expressing Th2 cells may explain the association of this chemokine with many allergic diseases. Our data support this hypothesis by showing the coexpression of CCL18 and CCR8 in human diseased esophageal tissue in individuals with active EoE. This may also be of particular importance in atopic dermatitis where CCL18 is abundantly expressed and likely facilitates the recruitment of CCR8–expressing skin-homing Th2 cells, as we have seen for mCCL8 (Islam et al., 2011). Further, the attraction of Th2 cells by CCL18-secreting AAM may provide a positive-feedback loop for the maintenance of the AAM phenotype in chronic inflammation. These are a few examples in which understanding the receptor for CCL18 will allow a greater appreciation of its role in human disease. Identifying CCR8 as the CCL18 receptor and mCCL8 as its rodent functional analogue will also enable the study of the CCR8 receptor–ligand system in murine disease models to better understand CCL18-mediated human disease pathology.

**MATERIALS AND METHODS**

**Human studies.** Peripheral blood was obtained from healthy volunteers, and fresh buffy coats were obtained from the Massachusetts General Hospital blood bank. For endoscopy studies, patients were referred for upper gastrointestinal symptoms suggestive of reflux or EoE (Table S1). Normal controls had a normal appearing esophagus on endoscopy and had no histological findings of inflammation on esophageal biopsy. In accordance with the consensus guidelines, active EoE was defined by the presence of >15 eosinophils per 40× high power field (hpf) in at least one esophageal biopsy (of a total six obtained from the proximal, middle, and distal esophagus) despite a minimum 6-wk course of high dose proton pump inhibitors (Liacouras et al., 2011). EoE patients in remission were on either elimination diet or on topical steroid treatment, and esophageal biopsy samples contained fewer than five eosinophils/hpf in all six biopsies. All human subject protocols were approved by the Partners Institutional Review Board. Informed consent was obtained from all study participants.

**Mice.** C57BL/6 mice were purchased from the National Cancer Institute (Bethesda, MD). CCR8−/− mice (from S. Lira, Mount Sinai School of Medicine, New York) in the C57BL/6 background were housed under specific pathogen-free conditions. All protocols were approved by the Massachusetts General Hospital Subcommittee on Research Animal Care.

**Human and mouse CCR8 receptor–transfected cells.** Limiting dilution clones were generated from a CCL18-responsive CCR8-transfected 4DE4 cell line (provided by J. Pease, Imperial College, London, UK) grown in low concentrations of G418 (0.1–0.4 mg/ml; Tiffany et al., 1997). We found that these clones responded better to CCL18 when grown in low G418 concentrations (and, in fact, seemed to lose activity at G418 concentrations ≥1 mg/ml) and were therefore maintained in 1.0 mg/ml G418. 4DE4 cells were provided by P. Murphy (National Institutes of Health, Bethesda, MD). hCCR8-L1.2 transfected cell line was provided by M. Locati and A. Mantovani (University of Milan, Milan, Italy). Cells were treated with 5 mM butyric acid overnight in some calcium flux, binding, and chemotaxis assays to increase surface expression of CCR8. Mouse Gr8−/−transfected Baf/3 cells were generated as previously described (Islam et al., 2011).

**Generation of human monocyte-derived DGs.** DCs were generated from PBMCs isolated from fresh human buffy coats by Histopaque-1077 density gradient (Sigma-Aldrich). Cells were treated with 5 mM butyrate overnight in some calcium flux, binding, and chemotaxis assays. Generation of Th2 and induction of highly polarized Th2 cells. CD4+ T cells were isolated from Histopaque-1077 density gradient (Sigma-Aldrich). PBMCs were resuspended in C10 media (RPMI complete medium with 10% fetal calf serum) and incubated for 2–3 h at 37°C and washed gently with PBS. The remaining adherent cells were cultured in 10 ml C10 media supplemented with 50 ng/ml of GM-CSF (BioLegend) and 50 ng/ml IL-4 (PeproTech) at 37°C. Additional GM-CSF and IL-4 was added on days 2, 4, and 5. On day 6, DCs were rested for 24 h after change of media, and then incubated with C10 media containing 100 ng/ml of LPS (Sigma-Aldrich) for 24–48 h.

**Generation of Th2 cells and induction of highly polarized Th2 cells.** CD4+ T cells were isolated from Histopaque-1077 density gradient PBMCs by two negative selection columns: (1) CD4+ T cell Biotin Antibody Cocktail (Miltenyi Biotec), followed by incubation with anti-biotin microbeads (Miltenyi Biotec) to isolate CD4+ T cells; and (2) CD45RO PE (BD) staining, followed by incubation with anti-PE microbeads to isolate naive (CD45RO−) CD4+ T cells. DCs and T cells were cultured in C10 media with 50 ng/ml IL-4 and 10 µg/ml anti–IFN-γ (BioLegend) for 5 d at 37°C. At day 5, new C10 supplemented with 250 ng/ml IL-2 (Roche) was added. For repolarization of Th2 cells at day 7 to generate Th2 R2 cells, Th2 R1 cells were repolarized with irradiated human PBMCs, 1 µg/ml anti-CD3 (BioLegend), 1 µg/ml anti-CD28 (BioLegend), 50 ng/ml IL-4, and 10 µg/ml anti–IFN-γ (BioLegend). A similar procedure was used to generate Th2 R3 cells from Th2 R2 cells. Prior to assays, Th2 cells were activated with 2 µg/ml plate-bound mAb to CD3 and 1 µg/ml soluble mAb to CD28 for 24 h. Cells were then washed, replated, and rested for 4–6 h in low-IL–2 medium for migration or IL–2–free medium for calcium flux. Highly polarized and activated mouse Th2 cells were generated for chemotaxis assays in a similar manner as previously described (Islam et al., 2011).

**Generation and activation of human and mouse macrophages.** Human monocytes were isolated from peripheral blood by negative selection with the monocyte isolation kit II (Miltenyi Biotec) and differentiated into...
macrophages by culturing in the presence of 50 ng/ml m-CSF (PeproTech) in C10 media for 7 d in 24- or 12-well plates. Human AAMs were obtained by culturing these macrophages in fresh C10 supplemented with 25 ng/ml IL-4 (PeproTech), 10 ng/ml IL-10 (Miltenyi Biotec), and 20 ng/ml TNF (Miltenyi Biotec) for an additional 24–72 h. Bone marrow cells isolated from femurs and tibias were cultured in non–tissue culture–treated 100-mm bacterial Petri dishes (Thermo Fisher Scientific) in C10 media supplemented with 10 ng/ml murine M-CSF (PeproTech) and fed with fresh media on days 4 and 8. Cells were harvested and replated in 24- or 12-well plates in M-CSF-containing media and stimulated with 20 ng/ml IL-4, 10 ng/ml IL-10, and 20 ng/ml TNF and harvested after 24 or 72 h.

Quantitation of integrin LFA-1 activation in highly polarized Th2 cells. Prior to assays, human Th2 cells were activated with 2 µg/ml plate-bound mAb to CD3 and 1 µg/ml soluble mAb to CD28 for 24 h. Cells were then washed, replated, and rested for 4–5 h in low-IL-2 medium before assays. For LFA-1 activation assays, cells were washed and stained in 1% human serum/RPMI with 10 µg/ml mAb24 (Hycult Biotech) in the presence of 200 nM CCL18, 200 nM CXCL12, or 50 ng/ml PMA or in the absence of stimulation for 30 min at 37°C, washed, and then stained with 1 µg/ml secondary goat anti–mouse PE for 20 min, washed, fixed, and acquired on a FACS Calibur. As a control, CD3/CD28-activated Th2 cells were stained with 10 µg/ml mAb24 in the presence of 200 nM CCL18 at 4°C for 30 min, followed by secondary antibody staining. As an additional control, Th2 cells that were not activated with CD3/CD28 were stimulated with 200 nM CCL18 and stained with 10 µg/ml mAb24 antibody at 37°C for 30 min followed by secondary antibody staining. CCL18 did not induce m24 epitope expression on these resting Th2 cells.

Flow cytometry and intracytoplasmic cytokine staining. For CCR8 staining, cells were stained with the 433H mAb to human CCR8, which was purified from the hybridoma (American Type Culture Collection) followed by PE-conjugated anti–mouse IgG1 (SouthernBiotech; Islam et al., 2011). Intracytoplasmic staining was performed using a standard protocol (Islam et al., 2011). Briefly, cells were stained with PE-conjugated mAb to IL-4 and allopurinol-cyan-conjugated mAb to IL-5 (BD), or appropriate fluorochrome-conjugated IgG1 isotype-control (BD). Cells were acquired on a FACS Calibur and analyzed with Flowjo (9.5.2).

Chemotaxis assays. CCR8-transfected cells, untransfected cells, and human Th2 cells (2.5 × 10^5–10^6 per well) in RPMI containing 0.5% BSA were placed on the top of a 96-well Chemtox chemotaxis apparatus with 5-µm pores (NeuroProbe). CCL18, human CCL1, and CXCL12 (R&D Systems) were used as chemotaxis assays. CCL18 was added to the top well in varying concentrations. The apparatus was incubated at 37°C and 5% CO2 for 3–4 h, and the cells migrating at each concentration of chemoattractant were counted using an inverted microscope at the end of the experiment or after fixation with 10% buffered formalin. The number of cells migrating at each concentration of chemokine was normalized to the number of cells migrating in the presence of medium alone (chemokinesis) to calculate the chemotactic index for each leukocyte type. PTX (Sigma-Aldrich) or the 433H neutralizing mAb to human CCR8 was added to the media at 37°C. RNA extraction and RT-PCR analysis for esophageal specimens and macrophages. Total RNA was extracted from cryopreserved esophageal biopsy samples using mirVana Isolation Kit (P/N: 1560; Ambion) according to the manufacturer’s protocol. Power SYBR green qRT-PCR assay (Applied Biosystems) was used for mRNA quantification using StepOnePlus Real-Time PCR System (Applied Biosystems). The following primers were used: CCL26, 5'-CAAGACCTGTCTGTCCAA-3' and 5'-GAATTCTATAGTCTGGCAACCA-3'; GTR2H, 5'-TAAAAAAGCTCAGGGAAGGGTTAAATG-3' and 5'-ACCCGGGAACCAAGAGAGAC-3'; CCR8, 5'-GGTCTACATCTCTGGTCCCTTG-3' and 5'-CAGGCGACGTGTCATCAGG-3'; CCR3, 5'-TCCAGACCTTGGGACAGCTC-3' and 5'-GGTGCAGGAGCTGGGA-3'; CCL1, 5'-TGCAGATCATCACCACACCC-3' and 5'-TGCAGATCATCACCACACCC-3'; CCL18, 5'-TGGAATTTGAGATATGGCAAAGCTAGTTG-3' and 5'-GGTGCAGGAGCTGGGA-3'; and GAPDH, 5'-GAAGCTGAGGAGTCGTGTTCCC-3' and 5'-GAAATGGTGATGGGATGAG-3'. Using the comparative Ct method, relative gene expression was calculated as 2^ΔΔCt, where ΔΔCt = Ct (gene of interest) – Ct (normalizer = GAPDH) and the ΔΔCt = Ct (sample) – Ct (calibrator = mean ΔCt of normal). Harvested mouse and human macrophages were harvested and stored in RLT buffer and purified RNA was isolated using the RNeasy kit (Qiagen). Real-time PCR was performed as above with the mouse and human primers described below. Mouse primers: Agl, 5'-CAGAAAGAATGGAAGGTGAGCAG-3' and 5'-CAGATATGCGAGGAGTCCACC-3'; Ym1, 5'-TCACAGGTCTGGCAATTTCTCTG-3' and 1896 CCR8 is a CCL18 receptor | Islam et al.
S'-TTTGGTCTTCTAGAGGGCTTCCTCCG-3', Fizz1, 5'-GGTCCCGAGGATGAGGACATGGAGAAG-3' and 5'-CACCCTCTCTCCGAGGAGACAGTGTTGACG-3'; Ccl8, 5'-TCTAACGGACGTTCTCTTTTGCCC-3' and 5'-AAGGGGATGCTTCTGAGCCTTAATGA-3'; Ccl17, 5'-CCCATGGAACCTTTCTCCTCC-3' and 5'-CATCCCTGGACACACTCCACTC-3'; and Gapdh, 5'-GGCAATTTCAAGCGGCAACTGGT-3' and 5'-AGATGGTGTGATGCTCCCC-3'. Human primers: CCL18, 5'-GGTTGCTACCTCCTCAAAAGAAGA-3' and 5'-GGCTGATTATTGTGGAGCACCCTT-3'; and CCL1C7, 5'-CCATGCTTTTTTGACTTGTCGACG-3' and 5'-TGCGATTCTTCACTCTTTGGTG-3'.

Statistical analysis. Statistical analysis was performed using Student’s unpaired two-tailed t test for means, except where indicated in Fig. 1 e (two-way ANOVA to assess the effect of multiple concentrations of CCL18 on CCL1-induced chemotaxis), Fig. 3 c (two-way ANOVA to compare the binding displacement of 125I-CCL18 by multiple concentrations of cold CCL18 and cold CCL1 to that induced by the non-CCR8 ligand CCL3), and Fig. 5 e (Mann-Whitney U test for nonparametric comparison of means among normal controls, patients with inactive EoE, and patients with active EoE). Spearman’s correlation coefficients were used for nonparametric testing of statistical dependence between pairs of predefined EoE-associated variables. P < 0.05 was considered significant.

Online supplemental material. Table S1 shows a summary of study subjects who underwent endoscopic biopsy after being referred for suspected reflux or EoE. Online supplemental material is available at http://www.jem.org/cgi/content/full/jem.20122842/DC1.

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