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Human β-Defensin 2 and 3 and Their Mouse Orthologs Induce Chemotaxis through Interaction with CCR2

Johann Röhrl,* De Yang,* Joost J. Oppenheim,* and Thomas Hehlgans‡

β-defensins play a dual role during immune response. Their direct antimicrobial properties contribute to the local innate immune response by combating microbial invasions. Furthermore, previous studies revealed the capacity of certain β-defensin family members to chemotactically recruit immature dendritic cells and CD45RO⁺ CD4⁺ T cells through chemokine receptor CCR6. However, because β-defensins also chemotactically recruit macrophages and monocytes, which do not express CCR6, efforts have been made to identify other receptors for these polypeptides. In this study, we demonstrate the capacity of human β-defensin (hBD)2 and 3 and their mouse orthologs, β-defensin 4 and 14, to interact with CCR2, a chemokine receptor expressed on monocytes, macrophages, and neutrophils. These β-defensins, fused to the Fc region of human IgG₄, showed binding to CCR2-transfected HEK293 cells, as revealed by flow cytometry. The β-defensin fusion proteins also induced CCR2-specific chemotaxis of transfected HEK293 cells, human peripheral blood mononuclear cells, and mouse peritoneal exudate cells in a dose-dependent manner. Preincubation of human monocytes with CCL2/MCP-1, the chemokine ligand for CCR2, abolished migration induced by β-defensins. Conversely, preincubation with hBD2:Ig or hBD3:Ig inhibited MCP-1 induced migration. Peritoneal exudate cells from CCR2-deficient mice failed to migrate toward these fusion proteins. In conclusion, the β-defensins used in this study contribute to the innate and adaptive immune response in their role as chemotactic agents. Our data indicate that hBD2 and hBD3, together with their mouse orthologs (β-defensin 4 and 14), are chemotactic for a broad spectrum of leukocytes in a CCR6- and CCR2-dependent manner.

Expression and purification of β-defensin fusion proteins

β-defensin fusion proteins were generated and purified as described earlier (17). In brief, cDNA encoding for the mature polypeptides of hBD2, hBD3, or mBD4 was cloned into the Signal Ig plus vector (R&D Systems, Minneapolis, MN), containing a eukaryotic cytomegalovirus promoter, using the following primers: 5′-CCC AGA TCT CTG ACC TGC CTG AAA AGC GG-3′ for hBD2-5′; 5′-CCC AGA TCT CAG AAA TAC TAC TGC CGT G-3′ for hBD3-5′; 5′-CCC AGA TCT AAT CCA ATA ACA TGC ATG-3′ for mBD4-5′; 5′-CCC AGG CTT CCA AAA TTT TTC TGC AGA-3′ for mBD14-5′; and 5′-CG CGG CCG CCA TCA TTT ACC CCG AGA G-3′ for human IgG-Fc-3′. After transfection, stable expressing Drosophila-S2 cells were selected and maintained in hygromycin (0.3 mg/ml; Invitrogen). The β-defensin fusion proteins were purified from the culture medium using HiTrap Protein G HP columns (GE Healthcare, Munich, Germany), according to the manufacturer’s instructions. Expression and purification of the fusion proteins were confirmed by Western blotting using peroxidase-conjugated donkey anti-human IgG mAb (Dianova, Hamburg, Germany) and polyclonal Abs against hBD2, hBD3, and mBD4 (Santa Cruz Biotechnology, Heidelberg, Germany).

Cell isolation and culture

HEK293 cells expressing human CCR2 variant B (hCCR2/HEK293) and HEK293 cells expressing human CXCR4 (hCXCR4/HEK293) were maintained in DMEM (Mediatech, Herndon, VA) containing 10% FCS (HyClone, Logan, UT; inactivated for 30 min at 56˚C) and 800 µg/ml G418 and were harvested when they reached 70–80% confluence.

Human peripheral blood enriched in mononuclear cells was obtained from healthy donors by leukapheresis (Transfusion Medicine Department, Clinical Center, National Institutes of Health, with approved human subject agreement). The blood was centrifuged through Histopaque-1077 (Sigma-Aldrich, St. Louis, MO), and PBMCs collected at the interface were washed with PBS. After centrifugation through an iso-osmotic Percoll (GE Healthcare, Pittsburgh, PA) gradient, the enriched monocytes (peripheral blood monocytes) were obtained from the top of the gradient.

Mouse peritoneal exudate cells (PECs) were elicited by i.p. injection of 2 ml 3% thioglycollate into 8–12-wk-old C57BL/6 wild-type or CCR2-deficient mice. After 3 d, PECs were isolated by lavage of the peritoneum (30 ml 2 ml 3% thioglycollate into 8–12-wk-old C57BL/6 wild-type or CCR2-deficient mice). After 3 d, PECs were isolated by lavage of the peritoneum.

Chemotaxis assay

Cells were suspended in chemotaxis medium (RPMI 1640 containing 1% BSA, 20 mM HEPES, 2 mM l-glutamine, 100 U/ml penicillin, and 100 µg/ml streptomycin) at 1 × 10^6 cells/ml for monocytes, 5 × 10^5 cells/ml for PECs, and 2 × 10^5 cells/ml for human CCR2*, human CCRX4*, or wild-type HEK293 cells. In certain experiments, primary cells were precultured with 10–100 ng/ml chemotractant (as indicated) for 30 min at 37˚C in humidified air containing 5% CO2. Synthetic hBD2 and hBD3 were purchased from Peprotech (Rocky Hill, NJ) and used as positive control. The migration of cells in response to chemotactants (β-defensins or control chemotactic factors) was determined using the 48-well microchemotaxis chamber assay, as previously described (30). In brief, chemotactants diluted in chemotaxis medium at various concentrations were put into the lower wells of a 48-well microchemotaxis chamber (NeuroProbe, Gaithersburg, MD), and cell suspension was added to the upper wells. The lower and upper compartments were separated by an uncoated polycarbonate filter membrane (5 µm pore size) for primary cells or a collagen-coated polycarbonate filter membrane (10 µm pore size) for human CCR2*, human CCRX4*, or wild-type HEK293 cells (both from NeuroProbe). After incubation at 37˚C for 1.5 h for primary cells and 5 h for HEK293 cells in humidified air with 5% CO2, the filters were removed, scraped, and stained. The cells that migrated across the filter were counted under a light microscope using Bioquant Image Analysis software (Bioquant Image Analysis, Nashville, TN). The results (mean ± SD of triplicate wells) are presented as the number of cells per high-power field.

Flow cytometry

All staining steps were performed for 20 min on ice in PBS containing 2% FCS and 0.05% Azide. HEK293 cells were incubated with 10% goat serum to prevent nonspecific binding. Primary cells were blocked with 10% goat or rat serum. HEK293 cells were labeled with 1 µg/ml β-defensin fusion protein. After washing, cells were stained with a PE-conjugated anti-human IgG Ab (Jackson ImmunoResearch Laboratories, West Grove, PA). Expression of CCR2 or CXCR6 on human monocytes was analyzed using an anti-human CCR2-PE Ab (clone 48607) or an anti-human CCR6-FITC Ab (clone 53103) purchased from R&D Systems (Minneapolis, MN). Detection of CCR2 or CXCR6 expression on PECs was performed using an anti-mouse CCR2 Ab from goat (GeneTex, Irvine, CA) and an anti-goat IgG-FITC secondary Ab (Jackson ImmunoResearch Laboratories) or an anti-mouse CXCR6 Ab from rat (clone 140706; R&D Systems) and an anti-rat IgG-Alexa 488 Ab (Molecular Probes, Eugene, OR). Flow cytometry was performed on a FACScan cytometer using Cell Quest software (BD Biosciences, San Jose, CA). Flow cytometry data were analyzed with FlowJo software (version 8.1.0) (Tree Star, Ashland, OR).

Statistical analysis

Statistical analyses of chemotaxis data were performed using one-way ANOVA with the Dunnett posttest (GraphPad Prism, version 4.0c; GraphPad, San Diego, CA); medium control served as the reference value. Statistical significance was considered at p < 0.01.

Results

Binding of β-defensins to CCR2 but not CXCR4

The β-defensins used in this study were expressed as fusion proteins, fused to the Fc region of human IgG1 (Fig. 1A), allowing easy detection of binding. Expression of fusion proteins was detected by Western blot analysis using an anti-human IgG Ab (Fig. 1B, lower panels). The β-defensin domain of the fusion proteins was detected by polyclonal Abs against hBD2, hBD3, and mBD4 (Fig 1B, upper panels). Both detection methods revealed an apparent m.w. of 37 kDa for the β-defensin fusion proteins (Fig. 1). There is no commercially available Ab against mBD14. In earlier studies, we showed that these β-defensin fusion proteins retained antimicrobial activity comparable to unfused β-defensins (17, 31).

In this study, hBD2:lg and hBD3:lg, as well as their mouse orthologs mBD4:lg and mBD14:lg, were used to examine their interaction with CCR2 and CXCR4. FACS analysis revealed that hBD2:lg, hBD3:lg, mBD4:lg, and mBD14:lg bound to human CCR2-expressing HEK293 cells independently of species.

FIGURE 1. Expression of hBD2, hBD3, and their mouse orthologs (mBD4 and mBD14) as human IgG1-Fc fusion proteins. A, Schematic representation of the β-defensin fusion protein (not to scale). B, Western blot analysis of recombinantly expressed hBD2:lg, hBD3:lg, mBD4:lg, and mBD14:lg using polyclonal Abs against hBD2, hBD3, and mBD4 (upper panels) or anti-human IgG Ab (lower panels). Data are representative of at least three independent experiments.
specificity (Fig. 2A). In contrast, no binding of β-defensin fusion proteins was observed using human CXCR4-expressing HEK293 cells (Fig. 2B). Unfused human IgG1 did not bind to wild-type or transfected HEK293 cells (Fig. 2).

The β-defensins induced chemotaxis of human CCR2-expressing HEK293 cells

Chemotaxis assays using human CCR2-expressing HEK293 cells were performed to elucidate whether the binding of hBD2:Ig and hBD3:Ig, as well as mBD4:Ig and mBD14:Ig, to human CCR2 has functional consequences. Human MCP-1 induced dose-dependent migration of human CCR2-expressing HEK293 cells in the typical bimodal manner, reaching a maximum at a concentration of 10 ng/ml (Fig. 3A). Human SDF-1α, the ligand for human CXCR4, induced dose-dependent migration of human CXCR4-expressing HEK293 cells, with maximal migration at 100 ng/ml (Fig. 3B).

The β-defensin fusion proteins induced chemotaxis of human CCR2-expressing HEK293 cells in a dose-dependent manner, which peaked at a concentration of 100 ng/ml (Fig. 3C–F). In agreement with the inability of the β-defensin fusion proteins to bind to human CCR2-expressing HEK293 cells, they also failed to induce chemotaxis of these cells (Fig. 3C–F). To exclude effects related to the human IgG1-Fc part of the fusion proteins, we tested unfused human IgG1 and showed that it did not induce cell migration (Fig. 3G).

The β-defensin fusion proteins induced chemotaxis of human peripheral blood monocytes

Human monocytes isolated from peripheral blood, which express endogenous CCR2 but not CCR6 (Fig. 4), were used to determine whether the β-defensin fusion proteins were also chemotactic for primary monocytes. Similar to the migration induced by human MCP-1 (Fig. 5A), the chemotactic effect of hBD2:Ig and hBD3:Ig, as well as mBD4:Ig and mBD14:Ig, peaked at a concentration of 10 ng/ml (Fig. 5C–F). In contrast, monocytes preincubated with 10 ng/ml human MCP-1 failed to migrate toward the β-defensin fusion proteins (Fig. 5C–F). Conversely, preincubation of monocytes with 10 ng/ml human IgG1 failed to migrate toward human SDF1-α (data not shown). Human monocytes failed to migrate toward human IgG1 (Figs. 5B, 6). The chemotactic effect of hBD2:Ig and hBD3:Ig, at concentrations of 10 ng/ml, was comparable to that of synthetic hBD (shBD)2 and shBD3 for human monocytes (Fig. 6). Pretreatment of monocytes with 10 ng/ml shBD2 inhibited hBD2:Ig-induced migration (Fig. 6A). Similarly, pretreatment with 10 ng/ml shBD3 abolished hBD3:Ig-induced chemotaxis (Fig. 6B). Synthetic mBD4 and mBD14 were not commercially available for further analysis.

CCR2-dependent migration of PECs

Thioglycollate-elicited PECs from C57BL/6 wild-type mice are a heterogeneous cell population. The majority of PECs expressed CCR2 (Fig. 7A). CCR6-expressing cells were not detected (Fig. 7B). Mouse MCP-1 induced maximal migration at a concentration of 1 ng/ml (Fig. 8A). However, the PECs also showed chemotactic responses to mBD4:Ig, mBD14:Ig, and hBD2:Ig, peaking at a concentration of 100 ng/ml (Fig. 8C–E). Furthermore, hBD3:Ig induced maximum migration at a concentration of 10 ng/ml (Fig. 8F). The capacity to migrate toward β-defensin fusion proteins or mouse MCP-1 was abolished in PECs from CCR2-deficient C57BL/6 mice (Fig. 8A, 8C–F). Unfused human IgG1 did not induce migration of PECs (Fig. 8B).

Discussion

In earlier reports, we described the production and characterization of β-defensin fusion proteins. These fusion proteins retained their antibacterial and chemotactic activities (17, 31) and were chemotactic for CCR6-expressing HEK293 cells, as well as for monocytes that do not express CCR6. The constant human IgG1 domain of the fusion proteins facilitated efficient purification and detection in Western blot analysis, as well as flow cytometric analysis. Polyclonal Abs against hBD2, hBD3, and mBD4 allowed specific detection of the β-defensin domains of these fusion proteins. Abs against mBD14 are not commercially available.

Observations suggested that CCR2 may be another G protein-coupled receptor for hBD3 (G. Jin, personal communication).
Therefore, we investigated whether hBD2, as well as hBD3 and their mouse orthologs mBD4 and mBD14, uses CCR2 in addition to CCR6. We highlight the effects of these β-defensins, fused to the Fc region of human IgG1, with CCR2. As we reported earlier, the interaction of mBD4:Ig and mBD14:Ig with CCR6 is species specific, but hBD2:Ig and hBD3:Ig were able to bind to human and mouse CCR6. In contrast, the binding of the mBD fusion proteins, as well as their human orthologs, to CCR2 is not species specific, and they interact with human or mouse receptors. 

Recombinant human CCR2-expressing HEK293 cells showed directional migration induced by the β-defensin fusion proteins, indicating functional interaction of hBD2, hBD3, mBD4, and mBD14 with CCR2. Primary human peripheral blood monocytes expressing CCR2, but not CCR6, also demonstrated consistent migration induced by all of the β-defensin fusion proteins used in this study.

In an effort to ascertain whether the β-defensins and MCP-1 share the same receptor, their mutual desensitizing effects were investigated. Preincubation of monocytes with human MCP-1 abolished the migration induced by hBD fusion proteins. Conversely, preincubation with hBD fusion proteins abolished migration induced by human MCP-1, indicating that these proteins share the same chemotaxis-inducing receptor. In contrast, the

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**FIGURE 3.** Comparison of the chemotactic effect of β-defensin fusion proteins and human MCP-1 on human CCR2/HEK293 cells. Migration of human CCR2/HEK293 cells, human CXCR4/HEK293 cells, or wild-type HEK293 cells induced by human MCP-1 (A) or human SDF-1α (B). Chemotaxis of chemokine receptor-expressing or wild-type HEK293 cells induced by hBD2:Ig (C), hBD3:Ig (D), mBD4:Ig (E), or mBD14:Ig (F). G, Migration induced by human IgG1. Data are representative of three independent experiments. *p < 0.01, one-way ANOVA with the Dunnett posttest versus media control.

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**FIGURE 4.** Expression of CCR2, but not CCR6, on human peripheral blood monocytes. Monocytes, isolated from human blood, were labeled with anti-hCCR2-PE Ab (A, open graph) or anti-CCR6-FITC Ab (B, open graph) and isotype control (shaded graph). Data are representative of three independent experiments.
chemotactic response to human SDF-1α was not desensitized by preincubation with hBDs or human MCP-1 (data not shown). The hBD fusion proteins induced chemotaxis of human monocytes similarly to synthetic hBDs. Furthermore, pretreatment of monocytes with these synthetic peptides inhibited migration induced by hBD2:Ig and hBD3:Ig, demonstrating the capacity of synthetic

![Figure 5](http://www.jimmunol.org/)

**FIGURE 5.** β-defensin fusion protein-induced chemotaxis of human peripheral blood monocytes. A. Dose-dependent migration of human monocytes induced by human MCP-1. Cell migration of monocytes preincubated for 30 min with 10 ng/ml hBD2:Ig or hBD3:Ig. B. Migration of monocytes induced by unfused human IgG1. Chemotaxis of untreated monocytes or monocytes preincubated for 30 min with 10 ng/ml human MCP-1 induced by hBD2:Ig (C), hBD3:Ig (D), mBD4:Ig (E), or mBD14:Ig (F). Data are representative of three or four independent experiments. *p < 0.01, one-way ANOVA with the Dunnett posttest versus media control.

![Figure 6](http://www.jimmunol.org/)

**FIGURE 6.** Inhibition of β-defensin fusion protein-induced chemotaxis of human peripheral blood monocytes by pretreatment with synthetic β-defensins. Migration of human monocytes induced by 10 ng/ml hBD2:Ig, shBD2, or unfused human IgG (A) and 10 ng/ml hBD3:Ig, shBD3, or unfused human IgG (B). Abrogated chemotaxis of monocytes after pretreatment for 30 min with 10 ng/ml shBD2 in response to hBD2:Ig (A) and 10 ng/ml shBD3 in response to hBD3:Ig (B). Data are representative of three independent experiments. *p < 0.01, one-way ANOVA with the Dunnett posttest versus media control.

![Figure 7](http://www.jimmunol.org/)

**FIGURE 7.** Comparison of expression of CCR2 on PECs from wild-type and CCR2-deficient C57BL/6 mice and expression of CCR6. A. Thiglycollate-elicited mouse PECs from wild-type mice (solid line) or CCR2-deficient mice (dashed line) labeled with anti-mCCCR2 polyclonal Ab from goat and secondary Ab anti-goat IgG-FITC and PECs incubated with secondary Ab alone (shaded graph). B. PECs from wild-type mice labeled with anti-mCCR6 (solid line) or isotype control (shaded graph). Data are representative of three independent experiments.

![Figure 8](http://www.jimmunol.org/)

**FIGURE 8.** Comparison of migration of wild-type PECs and CCR2-deficient PECs toward β-defensin fusion proteins. A. Chemotaxis of thioglycolate-elicited mouse PECs from wild-type mice or CCR2-deficient mice induced by human MCP-1. B. Migration of wild-type PECs induced by unfused human IgG1. Chemotaxis induced by mBD4:Ig (C), mBD14:Ig (D), hBD2:Ig (E), or hBD3:Ig (F). Data are representative of three or four independent experiments. *p < 0.01, one-way ANOVA with the Dunnett posttest versus media control.
peptides desensitize the interaction of recombinantly expressed fusion proteins with CCR2. The effects of synthetic mBD4 or mBD14 could not be tested, because these peptides are not commercially available. Additionally, chemoattraction of PECs from C57BL/6 wild-type mice by human and mBD fusion proteins was abrogated using cells from CCR2-deficient C57BL/6 mice. Because the lack of CCR2 in these gene-targeted mice is specific and irreversible, we were able to evaluate CCR2-dependent chemotaxis without concern for residual receptor function and, thus, ruled out the involvement of another chemotaxis-inducing receptor. These results clearly demonstrate specific interaction of hBD2-Ig, hBD3-Ig, mBD4-Ig, and mBD14-Ig with human and mouse CCR2.

A recent study indicated a possible interaction of hBD3 with the chemokine receptor CXCR4. The investigators described the competition of hBD3 with SDF-1α binding to CXCR4 and further reported hBD3-induced internalization of the chemokine receptor. However, direct interaction with CXCR4 was not shown (22). None of the β-defensin fusion proteins used in the present study demonstrated direct binding to human CXCR4 or chemoattraction of human CXCR4-expressing HEK293 cells. This argues against any direct interaction of these β-defensin fusion proteins with CXCR4.

Selective interaction of β-defensins with CCR2 and CCR6 raises the question of the underlying common binding motif. Although β-defensins and CC-chemokines, such as CCL2/MCP-1 or CCL20/MIP-3α, do not share significant sequence similarities, they do share structural similarities, such as the α helix, the abundance of cationic residues, and disulfide linkages, responsible for the distinct tertiary structure. Thus, common structural motifs and electrostatic overlap may be responsible for the CC-chemokine receptor-dependent chemotaxis-inducing activity of certain β-defensins (32). Although some chemokines also interact with multiple receptors, and because no chemokines interact with both CCR2 and CCR6, further studies are necessary to understand how β-defensins interact with several distinct chemokine receptors.

CCR2 and CCR6 were shown to be crucial for the recruitment of professional APCs to inflamed tissue (33). In particular, CCR2+ monocytes are recruited to sites of tissue inflammation, contributing to clearing pathogens and triggering adaptive immune response, whereas CCR2− monocytes infiltrate noninflamed tissues, where they may be involved in tissue homeostasis (24). Our findings support a potential role for β-defensins as mediators of enhanced immune reaction during inflammation. CCR6 is expressed by different lymphocyte subsets, such as CD45RO+ CD4+ T cells (34) and Th17 cells (35), as well as immature dendritic cells (14). Chemoattraction of these cells to sites of infection is necessary to initiate the adaptive immune response. However, CCR2 is largely expressed on myeloid cells, such as monocytes, macrophages (36, 37), and neutrophils (38), which are crucial for inflammation and phagocytosis. Thus, our data suggest that β-defensins contribute to the recruitment of a broad spectrum of leukocytes to sites of infection and inflammation.

In addition to their chemotactic and antimicrobial activity, there is evidence suggesting that β-defensins participate in the regulation of host innate and adaptive immune responses. A recent study provided data showing the capacity of hBD1 to induce the expression of maturation markers (e.g., MHC class II and CD83), costimulatory markers (e.g., CD80, CD86, and CD40), and proinflammatory cytokines (e.g., TNF, IL-6, and IL-12p70) in human monocyte-derived dendritic cells (39). Funderburg et al. (13) described TLR1- and TLR2-mediated expression of costimulatory molecules on monocytes and myeloid dendritic cells induced by hBD3. Furthermore, mBD2 is able to induce TLR4-mediated maturation of dendritic cells (12). Thus, we hypothesize that β-defensins, expressed by epithelial cells and leukocytes at sites of infection and inflammation, promote chemotactic recruitment and activation of leukocytes, such as monocytes, macrophages, and dendritic cells. Consequently, β-defensins produced at nanomolar concentrations by epithelial cells and leukocytes in response to infectious and injurious challenges form a haptotactic gradient in the tissue and along blood vessel walls to attract more APCs and monocytes to the endangered site. In addition, micro- molar concentrations of β-defensins exert their antimicrobial activity against invasive microorganisms.

Recent studies indicated that CCR2+ macrophages are crucial mediators of ulcerative colitis-associated colon carcinogenesis (40). In oral carcinoma, hBD3 was shown to be overexpressed by tumor cells, correlating with the recruitment and infiltration of macrophages (41). Furthermore, endothelial progenitor cells were reported to be recruited into tumors in a CCR2- and CCR5-dependent manner, contributing to neovascularization (42). These findings, together with our results, make it plausible for β-defensins to contribute to tumorigenesis and tumor development, as suggested by G. Jin (personal communication). Further studies are required to identify potentially different cellular recruitment patterns in tissues following infection, inflammation, or tumorigenesis, depending on the type of β-defensin expressed.

CCR2 seems to be the unidentified G protein-coupled receptor responsible for CCR6-independent β-defensin-mediated chemotaxis of monocytes. The β-defensin fusion proteins are able to recruit a broad spectrum of leukocytes in a CCR6- and CCR2-dependent manner. Based on our data, we conclude that the recruitment capabilities of β-defensins contribute to the innate and adaptive immune response in their role as chemoattractants.

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Disclosures

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