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Accessibility
Differential expression of three T lymphocyte-activating CXC chemokines by human atheroma-associated cells

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Activated T lymphocytes accumulate early in atheroma formation and persist at sites of lesion growth and rupture, suggesting that they may play an important role in the pathogenesis of atherosclerosis. Moreover, atherosclerotic lesions contain the Th1-type cytokine IFN-γ, a potentiator of atherosclerosis. The present study demonstrates the differential expression of the 3 IFN-γ-inducible CXC chemokines — IFN-inducible protein 10 (IP-10), monokine induced by IFN-γ (Mig), and IFN-inducible T-cell chemoattractant (I-TAC) — by atheroma-associated cells, as well as the expression of their receptor, CXCR3, by all T lymphocytes within human atherosclerotic lesions in situ. Atheroma-associated endothelial cells (ECs), smooth muscle cells (SMCs), and macrophages (MO) all expressed IP-10, whereas Mig and I-TAC were mainly expressed in ECs and MO, as detected by double immunofluorescence staining. ECs of microvessels within lesions also expressed abundant I-TAC. In vitro experiments supported these results and showed that IL-1β, TNF-α, and CD40 ligand potentiated IP-10 expression from IFN-γ-stimulated ECs. In addition, nitric oxide (NO) treatment decreased IFN-γ induction of IP-10. Our findings suggest that the differential expression of IP-10, Mig, and I-TAC by atheroma-associated cells plays a role in the recruitment and retention of activated T lymphocytes observed within vascular wall lesions during atherogenesis.

hybridization (17). To date, I-TAC has not been correlated with any human diseases.

Chemokines such as IL-8, monocyte chemoattractant protein-1 (MCP-1), MCP-4, and RANTES have been shown to be expressed within atherosclerotic lesions in situ and by atheroma-associated cells in vitro (28–35). In addition, in recent in vivo studies, targeted disruption of the genes for MCP-1, CCR2, and CXCR2 significantly decreased atherosclerotic lesion formation and lipid deposition when the disrupted gene was bred or transferred into mouse strains prone to develop atherosclerotic-like lesions (36, 37, 29). Furthermore, in these 3 in vivo studies, the attenuated development of vascular lesions correlated with decreased MØ accumulation in lesions, demonstrating that chemokines play a critical role in monocyte/MØ recruitment during atherogenesis.

It is likely that chemokines also play a critical role in the recruitment and retention of activated T cells in atherosclerosis. Because IFN-γ appears to have a proatherogenic effect, we have hypothesized that the IFN-γ-inducible chemokines IP-10, Mig, and I-TAC play an important role in atherosclerosis. Moreover, we investigated whether CD40 ligand (CD154), a molecule recently implicated in atherosclerosis (38, 39), and nitric oxide (NO), which has an antiatherogenic effect (40, 41), would regulate IP-10, Mig, and I-TAC expression.

Figure 1
Expression of IP-10, Mig, and I-TAC in human atherosclerotic lesions in situ. Human carotid arteries sections were stained with specific antibodies to IP-10, Mig, and I-TAC. High-power view (×100) of atherosclerotic lesions from different stages (Intimal thickening, Fatty streak, and Atheroma) revealed the expression of the chemokines (red-brown reaction product). Adjacent sections of the same atherosclerotic carotid tissue stained for IP-10 was stained with the rabbit preimmune IP-10 serum (×100). Normal human artery exhibited no expression of IP-10 (×100). The lumen of the artery is at the upper left side of each photomicrograph. Analysis of 5–7 atheroma at each stage of lesion development from different donors, and normal tissue from 4 different donors showed similar results.
Methods

Reagents. Affinity-purified rabbit anti-human IP-10 polyclonal antibody was generated as described previously (42). Rabbit anti-human Mig polyclonal antibody was obtained from PeproTech Inc. (Rocky Hill, New Jersey, USA). Affinity-purified rabbit anti-human I-TAC polyclonal antibody was generated as described (18). Mouse anti-human CXCR3 (1C6) mAb was a gift from LeukoSite (Cambridge, Massachusetts, USA). The following human recombinant cytokines were obtained from Endogen Inc. (Cambridge, Massachusetts, USA): IFN-γ, IL-1β, and TNF-α. Human recombinant CD40 ligand (rCD40L) was a gift from P. Graber (Ares Serono, Geneva, Switzerland) and was generated as described previously (43). The NO donors S-nitroso-N-acetylpenicillamine (SNAP) and S-nitrosoglutathione (GSNO) were purchased from Sigma Chemical Co. (St. Louis, Missouri, USA).

Cell isolation and culture. Human vascular ECs were isolated from saphenous veins by collagenase treatment (1 mg/mL; Worthington Biochemical Corp., Freehold, New Jersey, USA) and were cultured in dishes coated with fibronectin (1.5 μg/cm²; New York Blood Center Reagents, New York, New York, USA) as described elsewhere (44). Cells were maintained in M199 medium (BioWhittaker Inc., Walkersville, Maryland, USA) supplemented with 1% penicillin/streptomycin (BioWhittaker Inc.), 5% FCS (Atlanta Biologicals, Norcross, Georgia, USA), 100 μg/mL heparin (Sigma Chemical Co.), and 50 μg/mL EC growth factor (ECGF; Pel-Freez Biologicals, Rogers, Arkansas, USA). Human vascular SMCs were isolated from human saphenous veins and carotid arteries by explant outgrowth (38) and were cultured in DMEM (BioWhittaker Inc.) supplemented with 1% L-glutamine (BioWhittaker Inc.), 1% penicillin/streptomycin, and 10% FCS. Both cell types were subcultured after trypsinization (0.5% trypsin [Worthington Biochemicals] and 0.2% EDTA [EM Science, Gibbstown, New Jersey, USA]) in P100 culture dishes (Becton Dickinson and Co., Franklin Lakes, New Jersey, USA) and were used throughout passages 2–4. Culture media and FCS contained less than 40 pg LPS/mL, as determined by chromogenic Limulus amoebocyte-lysozome assay analysis (QLC-1000; BioWhittaker Inc.). ECs and SMCs were characterized by immunostaining with anti-von Willebrand factor and anti–SMC α-actin antibodies (DAKO Corp., Carpinteria, California, USA), respectively. Both cell types were cultured 12 hours before the experiment in media lacking FCS; ECs were cultured in M199 supplemented with 0.1% HSA and SMCs cultured in insulin/transferrin (I/T) medium with 0.1% BSA as described elsewhere (45).

Monocytes were isolated by adherence from PBMCs after Ficoll-Hypaque gradient and were cultured in RPMI-1640 medium (BioWhittaker Inc.) containing 10% FCS (Sigma Chemical Co.). Monocyte-derived MØ

Figure 2

Colocalization of IP-10, Mig, and I-TAC with ECs, SMCs, and MØ in human atherosclerotic lesions. High-power views (×400) of human carotid sections showed specific staining for IP-10, Mig, and I-TAC (Texas red staining) within the atherosclerotic lesions. Cell types were characterized by immunofluorescence staining with anti-CD31 mAb for ECs, anti–α-actin mAb for SMCs, and anti-CD68 mAb for MØ (FITC; green staining). The lumen of the artery is at the top of each photomicrograph. Analysis of atheroma from 3 different donors showed similar results.
were serum starved 16 hours before experiments, and stimulated with IFN-γ in RPMI-1640 medium supplemented with 0.1% BSA.

Immunohistochemistry. Surgical specimens of human carotid atheroma and normal aorta were obtained by protocols approved by the Human Investigation Review Committee at the Brigham and Women’s Hospital. Serial cryostat sections (5 µm) were cut, air dried onto microscope slides (Fisher Scientific Co., Pittsburgh, Pennsylvania, USA), and fixed in acetone at –20°C for 5 minutes. Sections preincubated with PBS containing 0.3% hydrogen peroxide were then incubated for 90 minutes with primary or control antibody, diluted in PBS supplemented with 5% appropriate serum. After washing 3 times in PBS, sections were incubated with the respective biotinylated secondary antibody (for 45 minutes; Vector Laboratories, Burlingame, California, USA) followed by avidin-biotin-peroxidase complex (VECTASTAIN ABC kit; Vector Laboratories). Immunostaining was viewed using 3-amino-9-ethyl carbazole (Vector Laboratories) according to the recommendations provided by the supplier. Cell types were characterized by double immunofluorescence staining using FITC-labeled cell-specific antibody (anti-muscle actin mAb for SMCs (Enzo Diagnostics, New York, New York), anti-CD31 mAb for ECs (DAKO Corp.), anti-CD68 mAb for MO (DAKO Corp.), and anti-human CD3 for T lymphocyte (DAKO Corp.).

Western blot analysis. Normal aortas and carotid atherosclerotic tissues were obtained from human patients after surgery. Tissues were homogenized (Ultra-turrax T...
Figure 5
Expression of the chemokine-receptor CXCR3 on T lymphocytes within human atherosclerotic lesions. Adjacent sections of human carotid arteries at different stages (Intimal thickening, Fatty streak, and Atheroma) were stained with anti-CD3 or anti-CXCR3 antibody (arrows). Photomicrographs (×100) reveal CD3 and CXCR3 expression (red-brown reaction product). The lumen of the artery is at the top of each photomicrograph. Analysis of 5–7 atheroma at each stage of lesion development from different donors showed similar results.

Figure 6
Expression of CXCR3 on CD4+ T lymphocytes within human atherosclerotic lesions. Sections of human carotid arteries were stained with anti-CXCR3 antibody. (a) Low-power (×100) and (b) high-power (×400) views show CXCR3 expression (red-brown reaction product) in the shoulder region of the plaque. Colocalization of CD4+ lymphocytes (c) (FITC; green staining) with CXCR3 (d) (Texas red staining) was performed by immunofluorescence-double staining. The lumen of the artery is at the top of each photomicrograph. Analysis of atheroma from 4 different donors showed similar results.
25; IKA-Labortechnik, Wilmington, North Carolina, USA) and lysed (0.3 g tissue/mL lysis buffer) as described previously (46). Lysates were clarified (16,000 g for 15 minutes at 4°C), and protein concentration for each tissue extract was determined using a bicinchoninic acid (BCA) protein assay according to the instructions of the supplier (Pierce Chemical Co., Rockford, Illinois, USA).

Fifty microgram of tissue lysate protein per lane and supernatants (45 μL) of cultured ECs, SMCs, and monocyte-derived MØ were separated by SDS-PAGE under reducing conditions and blotted onto PVDF membranes (Millipore Corp., Bedford, Massachusetts, USA) using a semidry blotting apparatus (3.0 mA/cm² for 30 minutes; Bio-Rad Laboratories Inc., Hercules, California, USA). Membranes were blocked in 5% defatted dry milk/PBS/0.1% Tween-20 (PBST) and were then incubated with the primary antibody (rabbit anti–IP-10 1:2,000; rabbit anti-Mig 1:1,000; rabbit anti–I-TAC 1:500) for 1 hour. Blots were washed 4 times PBST, and the secondary peroxidase-conjugated antibody (Jackson ImmunoResearch Laboratories Inc., West Grove, Pennsylvania, USA) was added (1:10,000) for another hour. Finally, membranes were washed in PBST, and detection of the antigen was carried out using the enhanced chemiluminescence detection method according to the manufacturer’s recommendations (NEN Life Science Products Inc., Boston, Massachusetts, USA), and subsequent exposure of the membranes to x-ray film.

**Northern blot analysis.** Total RNA was extracted from samples using Stat-60 (Tel-Test Inc., Friendswood, Texas, USA). For Northern analysis, 20 μg total RNA was electrophoresed on a 1.2% agarose-formaldehyde gel and then capillary transferred to a GeneScreen membrane (NEN Life Science Products Inc.). After overnight prehybridization (50% formamide, 1% SDS, 4× SSC, 4× Denhardt’s solution, 0.8% glycine, and 0.17 mg/mL denatured salmon sperm DNA) at 42°C, blots were hybridized at 42°C in 50% formamide, 10% dextran, 1% SDS, 5× SSC, 1× Denhardt’s solution, and 0.17 mg/mL denatured salmon sperm DNA with 10⁶ cpm/mL [α-32P]dCTP-radiolabeled cDNA probe prepared by nick translation. The following fragments were used as probes: a 1-kb PstI fragment from hIP-10 cDNA, a 3-kb NotI fragment from hMig cDNA (kindly provided by J. Farber, National Institutes of Health, Bethesda, Maryland, USA), and a 300-bp BamHI/AvaI fragment from hI-TAC cDNA. A GAPDH cDNA probe was used as a control for RNA loading. Signal quantitation was determined using a phosphoimager (Molecular Imager System; Bio-Rad Laboratories Inc.). Levels of chemokine expression in any given sample were normalized to the GAPDH signal for that sample. Because different expo-
sure times were used for each chemokine quantification and normalized to the same GAPDH exposure, the ratio measurements can only be used to compare levels of expression within any given exposure of a blot and not between blots.

**IP-10 ELISA assay.** Release of IP-10 from cultured human vascular ECs was measured using a sandwich-type ELISA as described previously (27). Antibody binding was detected by adding p-nitrophenyl phosphate (Sigma Chemical Co.), and absorbance was measured at 405 nm in an Molecular Devices plate reader (Du Pont, Wilmington, Delaware, USA). The amount of IP-10 detected was calculated from a standard curve prepared with the recombinant protein. Samples were assayed in triplicate.

**Results**

**Human atheroma-associated cells express the chemokines IP-10, Mig, and I-TAC.** Immunohistochemical analysis of human atherosclerotic plaques showed expression of IP-10, Mig, and I-TAC within the lesion (Figure 1). Analysis of atherosclerotic lesions at different developmental stages from early intimal thickening (n = 5) through fatty streaks (n = 5) and fully developed atheroma lesions (n = 7) revealed the expression of the 3 IFN-γ-inducible chemokines. Intimal thickening showed sparse but reproducibly detectable expression of the chemokines. In contrast, fatty streaks and well-developed atheroma lesions consistently showed strong immunoreactivity for all 3 chemokines, most prominently at the luminal border and in the shoulder region of the plaque, the margin between the lesion and unaffected portion of the artery. No immunoreactivity was observed in normal vessels (n = 4) or in atherosclerotic lesions examined with control preimmune serum as shown for IP-10 (Figure 1, lower panels), confirming the specificity of the antibodies used in these experiments. To characterize further the expression of these chemokines within atherosclerotic lesions, we performed double immunofluorescence staining (n = 3) using cell-specific antibodies. ECs, SMCs, and MØ highly expressed IP-10 within human carotid atherosclerotic plaques (Figure 2). In contrast, Mig was expressed mostly on ECs and MØ, and to a lesser extent on SMCs (Figure 2). The chemokine I-TAC was expressed only on ECs and MØ (Figure 2). Furthermore, ECs within neovessel formations in atherosclerotic plaques (n = 3) expressed I-TAC but little IP-10 or Mig (Figure 3).

To support these immunohistochemical results, Western blot analyses were performed with the same chemokine antibodies used for immunohistochemistry on tissue extracts from human atherosclerotic (n = 6) and normal vessels (n = 4). The chemokines IP-10 and Mig were detected in 3 of 4 samples from atherosclerotic vessels and comigrated with the recombinant protein (Figure 4). Extracts of unaffected arteries contained neither IP-10 nor Mig.

**T lymphocytes within human atherosclerotic lesions express the chemokine receptor CXCR3.** In view of the finding that atheroma-associated cells in situ expressed the 3 IFN-γ-inducible CXC chemokines — IP-10, Mig, and I-TAC — we investigated the expression of their receptor, CXCR3, in human atherosclerotic lesions. Serial sections of atherosclerotic lesions adjacent to the ones used for examining the expression of IP-10, Mig and I-TAC (Figure 1) were used to study the expression of CXCR3 (Figure 5). Immunohistochemical analysis revealed the expression of CXCR3+ cells within atherosclerotic lesions at all stages of lesion development examined (intimal thickening, fatty streak and atheroma; n = 5, 5, 7, respectively). Adjacent sections of the same atherosclerotic tissue stained for CD3 indicated that virtually all CD3+ cells expressed CXCR3 (Figure 5). A comparison with the pattern of expression seen for the chemokines (Figure 1), revealed the colocalization of CXCR3+CD3+T-cells with the 3 CXCR3 ligands at all stages of atherogenesis. Double immunofluorescence staining (n = 4) with CD4 antibody indicated that virtually all CD4+T-cells expressed CXCR3+ (Figure 6). As demonstrated in previous studies (8), we also found that the vast majority of CD3+ cells were CD4+ lymphocytes (data not shown).

**Atheroma-associated cells in vitro express the chemokines IP-10, Mig and I-TAC.** To characterize further IP-10, Mig and I-TAC expression by atheroma-associated cells, we performed in vitro experiments using human vascular ECs, SMCs and human monocyte-derived MØ. Quiescent cells lacked detectable IP-10, Mig, and I-TAC mRNA or protein expression (Figure 7). IFN-γ(1,000 U/mL for 18
hours) induced IP-10 mRNA accumulation in ECs and MØ and, to a lesser extent, in SMCs. Mig mRNA was detected in ECs and to a lesser extent in MØ and SMCs after IFN-γ stimulation. In contrast, I-TAC mRNA was highly expressed by ECs and was not detected in SMCs and MØ. IP-10, Mig and I-TAC mRNA induction by IFN-γ in ECs depended on the concentration and time of stimulation; mRNA for all 3 chemokines accumulated after stimulation with as little as 10 U/mL of IFN-γ and occurred as early as 2 hours after stimulation (data not shown). Viewing of Mig mRNA expression in SMCs required a longer exposure of the blots shown in Figure 7a. Western blot analysis for IP-10, Mig, and I-TAC extended Northern blot results to detection of all 3 proteins after IFN-γ stimulation in ECs (Figure 7b). The absence of I-TAC and Mig detection in SMCs and monocyte-derived MØ supernatants is likely due to lower concentrations compared with ECs supernatants. The control bands were loaded with 10 ng recombinant protein, enabling quantitative comparisons between levels of chemokine expression.

Stimulation of human ECs and SMCs with other cytokines implicated in atherosclerosis, such as IL-1β (10 ng/mL), TNF-α (50 ng/mL), and CD40L (5 μg/mL) each by itself, had no effect on IP-10 mRNA (data not shown) or protein expression (Figure 8a). However, IL-1β, TNF-α, and CD40L synergized with IFN-γ in inducing the secretion of IFN-10 protein in ECs as measured by an IP-10–specific sandwich ELISA (27) (compared with IFN-γ alone; 4.2-fold for IL-1β/IFN-γ; 3.7-fold for TNF-α/IFN-γ; and 2.0-fold for CD40L/IFN-γ).

NO decreases IP-10, Mig, and I-TAC mRNA and protein expression by human vascular ECs. Incubation of human vascular ECs with the NO donor SNAP before IFN-γ stimulation caused a concentration-dependent decrease in IP-10, Mig, and I-TAC mRNA induction (Figure 8a). This effect was more pronounced for IP-10 and Mig than for I-TAC mRNA expression. The NO donor GSNO produced similar results (data not shown). Phosphoimager analysis showed that maximal effect (4.9-, 5.2-, and 1.5-fold decrease for IP-10, Mig, and I-TAC, respectively) was obtained with 1 mM SNAP (Figure 8b). The concentrations of NO donors described in these experiments were similar to those previously reported to act on vascular cells (47, 48), and the lowest concentration used (50 μM) still affected chemokine expression. Western blot analysis of supernatants from IFN-γ-stimulated ECs showed a similar decrease for IP-10 and Mig protein secretion after NO treatment (data not shown).

Discussion

Atherosclerotic lesions at all stages of development from fatty streaks to complicated plaques contain cells and molecules characteristic of immune-mediated processes (2). Macrophages and lymphocytes are the most numerous inflammatory cells found in atherosclerotic lesions. These cells elaborate growth factors and cytokines that mediate intimal hyperplasia and may therefore promote atherogenesis. Activated T lymphocytes accumulate early in atheroma formation and persist at sites of lesion growth and rupture, suggesting that they play an import-
eralization of these chemokines in atheroma-associated cells by NO, another agent implicated in the pathogenesis of atherosclerosis. Normal vascular ECs secrete NO in response to shear stress, whereas ECs overlying atherosclerotic plaques produce less NO (41). In addition to its vascular relaxing effects, NO plays important immunoregulatory functions, such as inhibiting nuclear factor-κB (NF-κB) activation. We found that exogenous NO limits IP-10, Mig, and, to a lesser extent, I-TAC, mRNA accumulation induced by IFN-γ in ECs. Along these lines, exogenous NO inhibits (by 25%) IL-1-induced IL-8 secretion from ECs (53), and inhibition of basal NO production upregulates EC MCP-1 mRNA and protein expression (48). These results suggest that diminished secretion of NO by ECs leads to an increase in cytokine-induced chemokine expression by these cells and, therefore, to a greater stimulus for leukocyte recruitment into atherosclerotic lesions. In addition, we observed that the CD40 pathway, recently implicated in atherogenesis, potentiated the IFN-γ-induced secretion of IP-10 from ECs. CD40 ligation induces MIP-1α, MIP-1β, RANTES, and MCP-1 in macrophages (54) and, thus, may participate in atherogenesis through the release of multiple chemokines.

Chemokines may not only influence leukocyte recruitment within atherosclerotic lesions, but they may also regulate a number of vascular cell and leukocyte functions related to the acute and chronic manifestations of atherosclerosis. In particular, IP-10 mRNA was induced in the rat carotid artery after balloon angioplasty and was shown to be a mitogenic and chemotactic factor for vascular SMCs (55), suggesting that IP-10 might be involved in vascular remodeling during atherosclerosis. In addition, IP-10 inhibits neovascularization and wound healing in vivo (56), activities that may contribute to the necrosis associated with atherosclerotic lesions. Furthermore, IP-10 augments IFN-γ production from Th1 cells (57) and thus may help establish an autocrine loop that serves to drive the inflammatory response within the diseased vessel. Thus, in addition to recruiting activated T cells, these IFN-γ-inducible chemokines may play a role in the pathogenesis of atherosclerosis by regulating SMC and EC function.

In conclusion, we have demonstrated that atheroma-associated cells in situ differentially express the 3 IFN-γ-inducible CXC chemokines IP-10, Mig, and I-TAC, and that all CD4+ T lymphocytes within the same lesions express their receptor CXCR3. The coexpression of these chemokines and their receptor within atherosclerotic lesions suggests their involvement in the regulation of lymphocyte recruitment into atherosclerotic lesions, and that neutralization of this pathway in vivo may modulate immune cell migration within vascular wall during atherogenesis.

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