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Research Article

The predominant lipoxygenase products of arachidonic acid were extracted and purified from synovial fluid and sonicates of synovial tissue of patients with rheumatoid arthritis (RA), spondyloarthritis (SA), or a noninflammatory arthropathy (NIA). The concentration of 5(S),12(R)-dihydroxy-6,8,10-(trans/trans/cis)-14-cis-eicosatetraenoic acid (leukotriene B4) in synovial fluid was elevated significantly in patients with RA and a positive latex test for rheumatoid factor ($P < 0.05$, $n = 14$) and in patients with SA ($P < 0.05$, $n = 10$), compared with that of subjects with NIA ($n = 9$). The content of 5(S)-hydroxy-6,8,11,14-eicosatetraenoic acid (5-HETE), but not of leukotriene B4, was elevated significantly in synovial tissue of seven patients with RA in comparison with that of four subjects with NIA ($P < 0.05$). A single intra-articular injection of corticosteroid significantly lowered the synovial fluid level of leukotriene B4 in six patients with RA. These data suggest an involvement of the potent chemotactic factors 5-HETE and leukotriene B4 in human inflammatory disease.

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Lipoxygenation of Arachidonic Acid as a Source of Polymorphonuclear Leukocyte Chemotactic Factors in Synovial Fluid and Tissue in Rheumatoid Arthritis and Spondyloarthritis

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ABSTRACT The predominant lipoxygenase products of arachidonic acid were extracted and purified from synovial fluid and sonicates of synovial tissue of patients with rheumatoid arthritis (RA), spondyloarthritis (SA), or a noninflammatory arthropathy (NIA). The concentration of 5(S),12(R)-dihydroxy-6,8,10-(*trans/trans/cis*)-14-*cis*-eicosatetraenoic acid (leukotriene B₄) in synovial fluid was elevated significantly in patients with RA and a positive latex test for rheumatoid factor ($P < 0.05$, $n = 14$) and in patients with SA ($P < 0.05$, $n = 10$), compared with that of subjects with NIA ($n = 9$). The content of 5(S)-hydroxy-6,8,11,14-eicosatetraenoic acid (5-HETE), but not of leukotriene B₄, was elevated significantly in synovial tissue of seven patients with RA in comparison with that of four subjects with NIA ($P < 0.05$). A single intra-articular injection of corticosteroid significantly lowered the synovial fluid level of leukotriene B₄ in six patients with RA. These data suggest an involvement of the potent chemotactic factors 5-HETE and leukotriene B₄ in human inflammatory disease.

INTRODUCTION

The activation of basophils, mast cells, and polymorphonuclear (PMN)¹ and mononuclear leukocytes

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¹Abbreviations used in this paper: HETE, mono-hydroxy-eicosatetraenoic acid; NIA, noninflammatory arthropathy; PMN, polymorphonuclear; RA, rheumatoid arthritis; SA, spondyloarthritis.

by immunological and other stimuli releases arachidonic acid, which is converted in part by lipoxygenation to unstable hydroperoxy-eicosatetraenoic acids. Although hydroperoxy-eicosatetraenoic acids are transformed largely to mono-hydroxy-eicosatetraenoic acids (HETE) (1-4), 5(S)-hydroperoxy-eicosatetraenoic acid is the unique source of a family of complex HETE, termed leukotrienes, that contain additional polar substituents and three conjugated double bonds (5). 5(S)-hydroxy-6(R)-glutathionyl - 7,9 - *trans* - 11,14 - *cis* - eicosatetraenoic acid (leukotriene C₄) and 5(S)-hydroxy-6(R)-cysteinylglycine - 7,9 - *trans* - 11,14 - *cis* - eicosatetraenoic acid (leukotriene D₄) are functionally critical components of the slow-reacting substance of anaphylaxis (6, 7). 5(S),12(R)-dihydroxy-6,8,10(*trans/trans/cis*)-14-*cis*-eicosatetraenoic acid (5,12-di-HETE or leukotriene B₄) and 5(S)-hydroxy-6,8,11,14-eicosatetraenoic acid (5-HETE) elicit human neutrophil and eosinophil chemotaxis and chemokinesis, enhance the expression of PMN leukocyte C3b receptors, increase the intraleukocyte concentration of guanosine 3':5' cyclic monophosphate, and evoke a modest release of lysosomal enzymes from human neutrophils (3, 8-10). Although the tissue content of 12-L-HETE has been found to be elevated in the lesions of psoriasis compared with noninvolved epidermis of the same patients (11), the levels of the far more potent chemotactic factors 5-HETE and leukotriene B₄ have not been assessed previously in human disease states. This report documents significantly higher concentrations of leukotriene B₄ in peripheral joint synovial fluid of patients with sero-positive rheumatoid arthritis and spondyloarthritis, and a higher content of 5-HETE in synovial tissue in sero-positive

rheumatoid arthritis than in noninflammatory arthropathies.

METHODS

[³H]-12-L-HETE was generated by incubating partially purified lipoxigenase from homogenates of 4×10^{10} washed human platelets with 2 mCi of [³H]arachidonic acid (52 Ci/mmol, New England Nuclear, Boston, Mass.) in 0.05 M Tris-HCl/0.10 M potassium phosphate buffer (pH 8.0) containing 20 μ M indomethacin for 4 h at 37°C as described (12). [³H]-5,12-di-HETE was generated by incubating 4×10^8 human neutrophils with 2 mCi of [³H]arachidonic acid in 10 ml of Hanks' solution containing 0.05 g of recrystallized ovalbumin/100 ml, 10 μ M indomethacin, and 20 μ M calcium ionophore A23187 for 60 min at 37°C (2, 3). The radiolabeled HETE were extracted and purified by sequential silicic acid column chromatography and reverse-phase high-performance liquid chromatography as described (2, 8, 12), and were identified by cochromatography with purified products that had been characterized previously by gas chromatography-mass spectrometry (2, 3, 12). Human neutrophil chemotaxis was assessed by a modified Boyden chamber technique as described (3, 9).

Patient population. Definite and classical rheumatoid arthritis (RA) were defined by the criteria of the American Rheumatism Association and patients were classified as having sero-positive RA (RA+) or sero-negative RA (RA-) according to the results of a standard latex fixation test for rheumatoid factor. Patients with spondyloarthritis (SA) and peripheral joint arthritis had diagnoses of ankylosing spondylitis ($n = 3$), enteropathic spondylitis ($n = 1$), psoriatic arthritis ($n = 1$), and Reiter's disease ($n = 5$). Patients with a noninflammatory arthropathy such as degenerative or traumatic joint disease served as the control subjects. At the time of collection of the synovial material, three of the patients were taking a total of 75–200 mg of indomethacin/d, and the others were taking aspirin in some form at a total daily dose of 2,400–6,000 mg. Some patients were also receiving hydroxychloroquine, gold, penicillamine, and/or 2.5–15 mg/d of Prednisone, but none had received intra-articular corticosteroids in either knee joint for 3 mo or more before the collection of synovial samples for the present study.

Extraction, purification, and measurement of the HETE in fluid and tissue samples. Synovial fluid was aspirated from knee joints and centrifuged at 2,000 g for 10 min at 4°C to remove cells and particulate material; the supernatant fluid was stored at -70°C until the HETE were extracted. 2-g portions of synovial tissue, which had been obtained at surgery and stored at -20°C, were minced in 2 ml of Hanks' solution at 4°C, homogenized for 3 min (Polytron; Brinkmann Instruments, Inc., Westbury, N. Y.), and sonicated for 2 min (Branson Sonic Power Co., Danbury, Conn.; model 140D, 200 W). After 10^8 dpm of [³H]-12-L-HETE and 5×10^4 dpm of [³H]-5,12-di-HETE were added to each 2–3-ml sample of synovial fluid and synovial tissue sonicate, duplicate 20- μ l aliquots were removed for protein (13) and DNA (14) assays; the samples were titrated to pH 4.0 with 2 M citric acid. Each sample was extracted three times with 4 ml of chloroform:methanol (2:1, vol/vol), the tissue sonicates were extracted three times again with 4 ml of ethyl ether, and the organic phases were pooled and dried under N₂.

The HETE were resolved and purified by sequential silicic acid column chromatography and reverse-phase high-performance liquid chromatography on a 4.6×250 -mm Ultrasphere ODS column (Altex Scientific Inc., Berkeley, Calif.)

that was equilibrated and developed isocratically with methanol:water:glacial acetic acid (790:210:0.1, vol/vol) at a flow rate of 1 ml/min as described (2, 8, 12). The 5,12-di-HETE (leukotriene B₄) and 5-HETE were recognized by comparing retention times with those of standards that had been purified and identified previously by gas chromatography-mass spectrometry (2, 3, 5). Each product was quantitated by optical density at the wavelength of maximum absorption, using previously established extinction coefficients (2, 5). The radioactivity in portions of the 12-L-HETE and 5,12-di-HETE peaks was determined to provide an estimate of recovery.

TABLE I
White Blood Cell Counts and Concentrations of Predominant Lipoxigenase Products in Synovial Fluid of Patients with Arthritis or NIA

Patient	Diagnosis	Synovial fluid		
		5-HETE ng/ml	5,12-di-HETE ng/ml	Leukocytes per mm ³
1	NIA	190	31	400 (25)*
2	NIA	1,330	64	1,000 (0)
3	NIA	355	70	350 (14)
4	NIA	120	119	400 (43)
5	NIA	199	38	150 (0)
6	NIA	249	58	150 (0)
7	NIA	241	84	600 (1)
8	NIA	2,198	32	100 (46)
9	NIA	410	35	250 (28)
10	SA	323	407	28,500 (84)
11	SA	136	306	10,300 (74)
12	SA	41	331	8,650 (68)
13	SA	129	229	5,900 (72)
14	SA	311	713	30,700 (91)
15	SA	209	204	3,850 (88)
16	SA	777	67	9,400 (54)
17	SA	916	21	8,850 (38)
18	SA	689	41	40,200 (91)
19	SA	107	44	26,800 (82)
20	RA+	345	156	88,000 (80)
21	RA+	570	135	12,300 (90)
22	RA+	787	458	10,200 (78)
23	RA+	616	64	7,960 (81)
24	RA+	622	39	34,720 (85)
25	RA+	311	62	38,500 (78)
26	RA+	596	132	3,500 (69)
27	RA+	1,067	518	22,500 (74)
28	RA+	1,556	126	13,600 (83)
29	RA+	1,275	301	5,700 (49)
30	RA+	387	85	14,100 (66)
31	RA+	504	88	24,800 (85)
32	RA+	86	50	9,600 (65)
33	RA+	235	64	16,000 (97)
34	RA-	244	165	430 (17)
35	RA-	403	17	8,050 (60)
36	RA-	99	52	44,800 (91)
37	RA-	1,198	31	1,340 (64)

* The numbers in parentheses represent the percent PMN leukocytes in each sample.

The addition of 5 mg of 5-methyl prednisolone acetate to one of each pair of 1-ml samples of synovial fluid from three patients did not alter the recoveries of 5,12-di-HETE or 5-HETE relative to those of the duplicate sample lacking the corticoid. The total nanograms of 5,12-di-HETE and of 5-HETE per milliliter of synovial fluid or per milligram of DNA in synovial tissue were calculated from the quantities recovered and the percent recovery of radiolabeled 5,12-di-HETE and 12-L-HETE, respectively. The reproducibility of the purification procedure was assessed by analyses of a series of replicate portions of several fluids. In each such series, the range of values was $< \pm 15\%$ of the mean.

RESULTS

The concentration of 5,12-di-HETE in synovial fluid was significantly higher for patients with RA+ ($P < 0.05$) and patients with SA ($P < 0.05$) than for the control subjects with a noninflammatory arthropathy (NIA) or those with RA- (Table I, Fig. 1). The difference between the concentrations of 5,12-di-HETE in SA and in the NIA control patients was more significant ($P < 0.02$) when the six patients with SA and synovial fluid levels of 5,12-di-HETE above the normal range were analyzed as a separate subgroup. Although the total white blood cell counts and the percentages of PMN leukocytes were higher in the patients with RA or SA than NIA (Table I), no significant correlation was found between the white blood cell count and the 5,12-di-HETE concentration in synovial fluid of patients with RA+ ($r = 0.13$) or SA ($r = 0.44$), or of all the patients considered as a group ($r = 0.14$). There was no statistically significant difference between the synovial fluid level of 5-HETE in any of the groups

of patients and that of the control subjects. In contrast, the 5-HETE content of synovial tissues was elevated significantly in RA+ patients relative to control subjects with NIA ($P < 0.05$), whereas the 5,12-di-HETE content of the tissues was not increased (Fig. 2). Similar results were obtained when the contents of HETE were expressed in terms of the wet weight of the tissues or the protein concentration of the tissue sonicates.

To assess the relative contribution of the HETE to the overall neutrophil chemotactic activity in inflammatory synovial fluids, the 5,12-di-HETE and 5-HETE were obtained as a mixture by silicic acid chromatography of extracts of 1-ml portions of synovial fluid from four patients with RA+. After determination of the yields of HETE according to recovery of the radioactive tracers, the mixtures of HETE each were re-suspended in the appropriate fraction of 1 ml of synovial fluid from a patient with NIA. In dose-response studies using 10–200 μl , the neutrophil chemotactic activity of unextracted portions of synovial fluid from the RA+ patients reached plateau values that ranged from 30.6 to 51.3 net neutrophils/high-power field at a concentration of 50 $\mu\text{l}/\text{ml}$ of buffer, whereas 50 μl of the synovial fluid from the patient with NIA attracted only 6.9 net neutrophils/high-power field. The increment in neutrophil chemotactic activity in the NIA synovial fluid achieved by the addition of HETE purified from the RA+ synovial fluids permitted an estimation that the HETE accounted for 35–57% of the chemotactic activity in the corresponding unextracted synovial fluids.

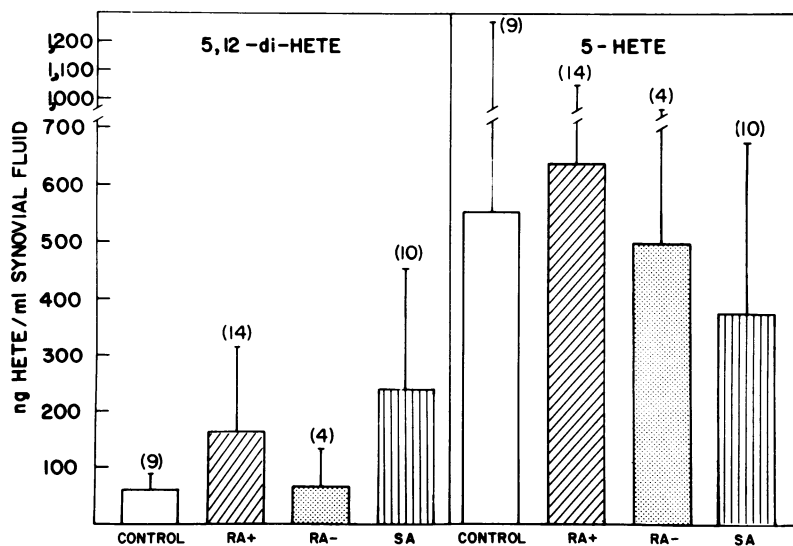


FIGURE 1 Concentrations of the predominant HETE in synovial fluid of arthritic patients. Each bar and bracket represents the mean \pm SD for the values of the number of patients shown in parentheses above the brackets. Statistical analyses employed Student's two-sample *t* test with the Welch approximation.

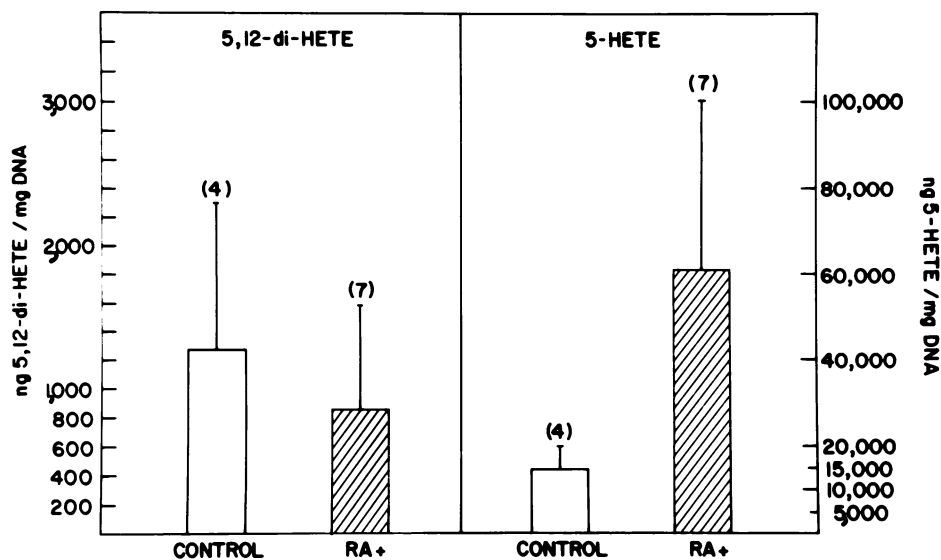


FIGURE 2 Levels of predominant HETE in synovial tissues of arthritic patients. The meaning of each bar and bracket and the statistical analyses are as in Fig. 1.

The effect of a single intra-articular injection of 40 mg of 5-methyl prednisolone acetate was examined in six RA+ patients. Before the injection of corticosteroid, the concentrations of 152 ± 157 ng (\pm SD) 5,12-di-HETE/ml and 542 ± 182 ng 5-HETE/ml were similar to those for other RA+ patients (Fig. 1). 3 d to 2 wk after the intra-articular injection, the concentration of 5,12-di-HETE had been reduced significantly to 67 ± 40 ng/ml ($p < 0.05$), whereas the concentration of 5-HETE was 526 ± 431 ng/ml and had not been altered by the corticosteroid.

DISCUSSION

That lipoxygenase products of arachidonic acid are capable of eliciting inflammatory cellular infiltrates in vivo was demonstrated initially by the rapid influx of eosinophils and the later accumulation of neutrophils in response to the intraperitoneal instillation of microgram quantities of 12-L-HETE in guinea pigs (15). The possibility that lipoxygenase products are the predominant stimuli of the cellular component of some inflammatory reactions in vivo was suggested by the capacity of the lipoxygenase inhibitor BW755C to suppress the infiltration of PMN leukocytes evoked in the rat paw by the injection of carrageenin, while the cyclooxygenase inhibitor indomethacin prevented the alterations in vascular permeability, but not the leukocyte response (16). The present data indicate that the synovial levels of the lipoxygenase-derived PMN leukocyte chemotactic factors 5,12-di-HETE or leukotriene B_4 and 5-HETE are elevated in relation to active synovitis and compared with the levels in synovia of subjects with noninflammatory arthropathies

(Figs. 1 & 2). Leukotriene B_4 is chemotactic for neutrophils and eosinophils in vitro at a concentration as low as 3 ng/ml and evokes a maximal chemotactic response at 30 ng/ml, compared with 1,000 ng/ml for 5-HETE and 10,000–20,000 ng/ml for 11-HETE and 12-HETE (10). Thus, the concentration of both leukotriene B_4 and 5-HETE in synovial fluid may be sufficient to contribute to the local inflammatory reaction (Fig. 1). The reduction in the synovial fluid level of leukotriene B_4 that followed the intra-articular injection of a corticosteroid in patients with rheumatoid arthritis is similar to the suppression of cutaneous levels of 12-HETE after topical corticosteroid therapy in psoriasis (17). The reduction in the synovial fluid leukocyte count and the transformation of the residual population from PMN to mononuclear leukocytes after the intra-articular injection of a corticosteroid (18) thus may be mediated in part by the suppression of the generation of lipoxygenase products of arachidonic acid, which are preferentially chemotactic for PMN leukocytes.

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