Apolipoprotein (Apo) E Inhibits the Capacity of Monosodium Urate Crystals to Stimulate Neutrophils

Characterization of Intraarticular Apo E and Demonstration of Apo E Binding to Urate Crystals in Vivo

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Abstract

Factors that modulate the ability of monosodium urate crystals to stimulate leukocytes could regulate gouty inflammation. Lipoproteins that bear apo B-100 and apo E bind to urate crystals and suppress crystal-neutrophil interaction. In this study, we observed that urate crystals, coated with apo E of monocye origin, had a diminished ability to stimulate neutrophils. Apo E was also detected on the surface of urate crystals recovered from gout patients. Thus, we analyzed apo E in noninflammatory synovial fluid, and found it to be associated with particles of heterogeneous size and of predominantly alpha and pre-beta electrophoretic mobility. Local articular synthesis of at least a portion of synovial fluid apo E was suggested because (a) the synovial fluid/plasma concentration ratio of apo E was significantly higher than that for both apo B and apo A-I, which are not widely synthesized by extrahepatic tissues, (b) cultured rheumatoid synovial cells in first passage secreted apo E, (c) a portion of synovial fluid apo E was heavily sialylated.

We conclude that synovial fluids contain apo E that appears partly of local origin. Apo E binds to urate crystals and could modulate gouty inflammation. (J. Clin. Invest. 1991. 87:20-26.) Key words: gout • synovial fluid • synovium • apolipoprotein B • apolipoprotein A-I

Introduction

Monosodium urate crystals encounter leukocytes in the synovial joints and periarticular bursae of individuals with gout (1), resulting in the release of inflammatory mediators that can trigger acute inflammation (2-5). However, the inflammatory response associated with free urate crystals in joints and bursae is highly variable (4, 5). Thus, much effort has been directed towards the characterization of factors that suppress urate crystal-induced leukocyte stimulation.

In this regard, it has been observed that urate crystals in clinically quiescent tophaceous synovial deposits are physically associated with variable amounts of surface proteins (6) and of lipids including esterified cholesterol (1, 7). Furthermore, a number of individual plasma proteins and lipoproteins bind to the surface of urate crystals and can alter their inflammatory potential in vitro (8-10). Significantly, when urate crystals are coated with plasma their ability to stimulate neutrophils in vitro is markedly diminished (10-12), an effect caused by the binding to crystals of lipoproteins bearing apolipoprotein (apo) B-100 (LDL) and apo B-100 and apo E (VLDL, IDL) (10-12). Apo B-100 mediates the binding of LDL to urate crystals and is itself able to suppress urate crystal-induced neutrophil stimulation (13). In this study, we demonstrate that apo E is also a potent inhibitor of the ability of urate crystals to stimulate neutrophils in vitro and that apo E is bound to urate crystals recovered from patients with gout.

These observations prompted us to quantify and characterize the apo E of synovial fluids. Apo E, in distinction to other apolipoproteins, is synthesized by both the liver and by numerous extrahepatic tissues, including cells of the monocyte/macrophage lineage (14-16). The apo E newly secreted by extrahepatic cells, including macrophages, is heavily sialylated (16, 17), which helps to distinguish it from the apo E found in circulating plasma lipoproteins (16, 17), which is totally or nearly totally desialylated via unknown mechanisms (17).

Apo E functions in lipid redistribution between tissues by serving as a ligand for the low density lipoprotein (LDL) receptor (15) and for a second, postulated receptor for chylomicron remnants (18). Apo E also can function in tissue repair by modulating lipid redistribution locally (15, 19). Last, apo E also possesses a variety of functions that are not related to lipid transport (15), including suppression, by apo E-bearing lipoproteins and by apo E polypeptides, of lymphocyte activation (20-23).

In this study, we demonstrate that apo E is present in human synovial fluids at substantial concentrations and we provide evidence for the presence there of apo E of local origin. These observations suggest that apo E may also have important functions within synovial joints.

Methods

Collection and treatment of synovial fluids. Synovial fluids, sera, and plasma were collected simultaneously from patients undergoing diagnostic procedures at the San Diego Veterans Administration Medical Center. Additional synovial fluids were a generous gift of Dr. G. Firestein (UCSD Medical Center, San Diego) and Dr. P. Fowler (University of Western Ontario, London, Ontario). Only nonhemorrhagic synovial fluids were studied. Patients with osteoarthritis were verified to have noninflammatory effusions and the absence of detectable intra-articular monosodium urate or calcium pyrophosphate crystals by compensated polarized light microscopy at 400× magnification (24). Patients with acute gout were confirmed to have intra-articular urate

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1. Abbreviation used in this paper: apo, apolipoprotein.
crystals (24). Patients with Reiter’s disease and rheumatoid arthritis had active disease and met accepted diagnostic criteria.

All synovial fluids, sera, and plasmas were treated with EDTA (5 mM), phenylmethylsulfonyl fluoride (PMSF) (100 μM) and aprotinin (0.01%) and centrifuged (6,900 g for 30 min at 4°C) to remove cells and particulates. Treatment of synovial fluids with 100 μg/ml of testicular hyaluronidase (Sigma Chemical Co., St. Louis, MO) was carried out for 3 h at 37°C to reduce viscosity and enable accurate pipetting (25). It was verified that identical treatment with hyaluronidase did not alter the measurement of apolipoproteins.

Isolation of plasma lipoproteins. The major lipoprotein density fractions were isolated from plasma by standard ultracentrifugation techniques in the presence of protease inhibitors and antioxidants as described (26). LDL and HDL were of d = 1.019–1.063 g/ml and d = 1.063–1.25 g/ml, respectively.

Quantitation of cholesterol, triglycerides, and apolipoproteins E, B, and A-1. Total cholesterol was measured enzymatically using the Boehringer Mannheim Diagnostics (Indianapolis, IN) high performance cholesterol reagent, and total triglycerides measured using the enzymatic triglyceride reagent from Abbott Laboratories (Irving, TX).

Apo E was quantitated by radioimmunnoassay (27), and apo B and apo A-I by ELISA as previously described (28, 29). Protein concentrations of apo E, LDL, and HDL were measured by a modification of the Lowry method (30).

Synovial fluid/plasma concentration ratios of apolipoproteins, cholesterol and triglycerides were compared using the paired, two-tailed t test as implemented with the Statview II program (Abacus Concepts, Inc., Berkeley, CA) on a Macintosh SE computer.

Electrophoresis and identification of apolipoproteins by Western blotting. Nondenaturating polyacrylamide (31), and agarose gel electrophoreses (13) were performed as previously described. Molecular weight standards for sizing gels were thyroglobulin (669 KD), ferritin (440 KD), catalase (232 KD), lactate dehydrogenase (140 KD), albumin (67 KD). For 13% SDS-PAGE we employed, as standards: phosphorylase b (94 KD), albumin (67 KD), ovalbumin (43 KD), carbonic anhydrase (30 KD), trypsin inhibitor (20.1 KD), alpha-lactalbumin (14.4 KD), and we employed apo E as an internal standard.

Gels were transferred to nitrocellulose paper that was probed for apolipoproteins as previously described, using antibodies I.E (apo E-specific), B47 (apo B-specific), and a pool of apo A-I–specific antibody– A.I-4, A.I-7, and A.I-10 (26). Neuraminidase treatment (16) of synovial fluid before electrophoresis was performed, by incubating 0.1 ml of whole synovial fluid with 0.1 ml of 0.1 M sodium acetate (pH 4.0) containing 1.5 units of C. perfringens neuraminidase (Sigma Chemical Co.) at 37°C for 24 h. Samples were extracted with 3 ml of ethanol:ether (3:1) at ~20°C for 30 min, and the pellet was washed once with diethyl ether (16).

To study urate crystals isolated from gouty patients, we isolated the crystals by repeated sedimentation of the fluids (1,000 g for 10 min), and crystals were washed in distilled water to lyse cells (29). Proteins were eluted from the crystal surface by boiling in SDS under reducing conditions (9), and were electrophoresed in 13% polyacrylamide gels.

Synovial cell isolation and culture. Synovial cells isolated from surgical specimens freshly removed from a patient undergoing joint replacement for rheumatoid arthritis, were a generous gift of Dr. Martin Lotz (Scripps Clinic). Collagenase digestion and culture were performed essentially as described (33). Cells were maintained in DMEM supplemented with l-glutamine, penicillin, streptomycin, and 10% FCS. At confluence, the cells were trypsinized, divided 1:3, and cultured at 100,000 cells/25 cm² flask in 4.5 ml of a serum-free medium that comprised DMEM supplemented with 20 mM Hepes, 1 mM Na pyruvate, 1% BSA, and 1% Nutridoma-HU (Boeringer Mannheim Diagnostics). After 7 d the supernatants were recovered by centrifugation and the medium assayed for apo E by radioimmunnoassay.

THP-1 cell culture and isolation of secreted apo E. THP-1 cells were obtained from the American Type Culture Collection (Rockville, MD) and maintained in suspension in T-150 culture flasks (Costar, Cambridge, MA) at a cell density of 0.2 to 1.0 × 10⁶/ml in 100 ml of RPMI-1640 containing 7% FCS (Rehatuin F.S., Armour Pharmaceutical, Kankakee, IL), 10 mM Hepes, 2 mM glutamine, Pen-Strep (penicillin, 200 U/ml; streptomycin, 2 mg/ml; Sigma Chemical Co.), 1 mM sodium pyruvate and 5 × 10⁻⁵ M beta-mercaptoethanol.

To stimulate apo E production the cells were washed three times in the above medium with 1% Nutridoma-HU instead of FCS. The cells were seeded in T-150 flasks (100 ml at 1 × 10⁶/ml) and stimulated with phorbol myristate acetate (100 nM). After 7 d of culture the medium was collected and the adherent cells refed with fresh medium. In this manner, medium containing apo E was collected four times from the same culture flask. The pooled conditioned media were stored at -20°C.

To purify the apo E, the medium was filtered through a 0.45-μm filter and applied to an immunoadfinity column. The column was prepared by coupling 11 mg of purified monoclonal antibody (IE) to 14 ml of Affigel HZ (Bio-Rad Laboratories, Richmond, CA) according to a protocol from the manufacturer that exhibited > 99% coupling efficiency. Conditioned medium (100 ml) was applied to the column for each isolation. Bound apo E was eluted with 5 M guanidine HCl and exhaustively dialyzed against PBS.

Neutrophil isolation and assay of urate crystal–induced alpha-mannosidase release. Human neutrophils were isolated from normal volunteers via Ficoll-Hypaque centrifugation as previously described (10). These preparations consistently contained > 98% neutrophils. We incubated neutrophils (4 × 10⁶/ml) in a buffer containing 147 mM NaCl, 5 mM KCl, 1.9 mM KH₂PO₄, 1.1 mM Na₂HPO₄, 5.5 mM glucose, 3 mM MgSO₄, 1 mM MgCl₂, 1.5 mM CaCl₂, 10 mM Hepes, and 2.5 mg/ml BSA, pH 7.4, with urate crystals (1 mg/ml) for 30 min at 37°C in a volume of 0.5 ml. The reactions were terminated by centrifugation and alpha-mannosidase release was measured at 405 nm (10).

Synthetic urate crystals were prepared and characterized as previously described (10), and were coated with apo E (or controls) at up to 10 μg protein per ml in a total volume of 0.25 ml and 10 mM PBS, pH 7.4. The buffer was saturated with urate before use to prevent crystal dissolution (9, 10) and it contained 2 mg/ml BSA as a physiologic background protein (9, 10). After incubation with proteins the urate crystals were washed three times before incubation with neutrophils.

Results

Apo E binds to urate crystals and inhibits urate crystal–induced neutrophil stimulation. Apo E–containing plasma lipoproteins have been previously observed to bind to urate crystals exposed to human serum and plasma in vitro (10). In this study, we observed that apo E bound to the surface of washed synthetic urate crystals that had been incubated with gouty joint fluid (Fig. 1, lane 2). Thus, we studied urate crystals aspirated from patients with gout and detected bound apo E on washed urate crystals in four of four patients studied, including crystals from the knee joint and olecranon bursa of two representative pa-

Figure 1. Apo E is bound to monosodium urate crystals in vivo and in vitro. Samples were separated by SDS-PAGE in a 13% gel and probed for apo E after transfer to nitrocellulose by Western blotting with an apo E–specific antibody as described in Methods: apo E of plasma VLDL (lane 1) served as an internal standard; protein eluted from 10 mg washed synthetic urate crystals that were previously coated with 0.1 ml gouty synovial fluid (lane 2); protein eluted from washed urate crystals (20 mg) aspirated from an asymptomatic gouty knee joint (lane 3); and an asymptomatic gouty olecranon bursa (lane 4).
representative of three separate patients studied. HDL, VLDL/IDL, and THP-1 cell supernatants were prepared as described in Methods.

tients (Fig. 1, lanes 3 and 4). Therefore, apo E binds to urate crystals in vitro and in vivo.

Lipoproteins containing only apo B-100 (LDL), and both apo B-100 and apo E (VLDL and IDL), suppress neutrophil activation by urate crystals (10). To determine if apo E could modulate this function we studied particles that contained only apo E. To do so, apo E was isolated via immunoaffinity chromatography of the conditioned media of human myelomonocytic leukemia cells (THP-1 cells) grown in serum-free medium. These particles were verified to be free of both apo B and apo A-I (Fig. 2) but to contain phospholipid (27) and apo E (Fig. 2).

Coating of urate crystals with THP-1 cell-derived apo E (at up to 10 μg protein/ml) resulted in progressive inhibition of urate crystal–induced release of the neutrophil azurophil granule constituent, alpha-mannosidase (Fig. 3). THP-1 cell apo E, added directly to cells at these concentrations, did not inhibit alpha-mannosidase release from cytochalasin B-treated neutrophils exposed to 1 μM N-formyl-met-leu-phe (not shown). Thus, the inhibition of the ability of apo E–coated urate crystals to stimulate neutrophils was not due to direct effects of apo E on the functional integrity of neutrophils.

The inhibitory capacity of apo E for urate crystal–induced neutrophil stimulation was comparable with that of human plasma LDL, which was utilized as a positive control (10) (Fig. 3). Furthermore, we confirmed (10) that the potent inhibitory activities of apo E, and of LDL, were not shared by apo E–poor HDL (Fig. 3), which also binds to urate crystals (10). In addition, the fraction of THP-1 cell conditioned media that was not retained by the apo E immunoabsorbant (and which contained < 0.1 μg/ml apo E) was essentially noninhibitory for urate crystal–induced neutrophil activation at identical total protein concentrations (Fig. 3).

Quantification of apo E in synovial fluid. Because low concentrations of apo E had marked effects on urate crystal–induced neutrophil stimulation, we looked for evidence that apo E was present in synovial fluids. We first assessed the concentration of apo E in synovial fluids relative to the plasma concentration of apo E in normal volunteers and we compared this ratio to that for apo B and apo A-I.

The ratio of the concentration of synovial fluid apo E to a mean normal plasma concentration (obtained from a panel of 20 unmatched normal donors) was significantly greater than the same ratio for apo B and for apo A-I in both 21 noninflammatory synovial fluid samples (Table II). Samples were electrophoresed in 5–30% polyacrylamide gels (Pharmacia Fine Chemicals, Piscataway, NJ) under nondenaturing conditions and stained with Coomassie Blue for protein (left) or probed, after electrophoretic transfer to nitrocellulose, with antibodies specific for apo E, apo B-100, and apo A-I as described in Methods. Synovial fluid (SF) and matching, simultaneously drawn serum was from a patient with osteoarthritis (Table II, patient No. 3) and was
Table I. Synovial Fluid (SF)/Plasma Ratio of Apo B-100, Apo A-I, and Apo E

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Apo B</th>
<th>Apo A-I</th>
<th>Apo E</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>SF/plasma ratio</td>
<td>SF/plasma ratio</td>
<td>SF/plasma ratio</td>
</tr>
<tr>
<td>Noninflammatory</td>
<td>0.18±0.04</td>
<td>0.27±0.04</td>
<td>0.58±0.04*</td>
</tr>
<tr>
<td>Inflammatory</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RA</td>
<td>0.44±0.03</td>
<td>0.56±0.07</td>
<td>0.81±0.06*</td>
</tr>
<tr>
<td>Acute gout</td>
<td>0.42±0.07</td>
<td>0.26±0.04</td>
<td>0.72±0.07*</td>
</tr>
<tr>
<td>Reiter's</td>
<td>0.38±0.08</td>
<td>0.57±0.07</td>
<td>0.84±0.06*</td>
</tr>
</tbody>
</table>

The concentrations of apolipoproteins in pathologic synovial fluids, and in the plasma of a panel of 20 unmatched, normal donors were measured by SPRIA. (Plasma values for the panel of normals for these assays were: apo B, 979±48 µg/ml; apo A-I, 1.87±0.25 mg/ml; apo E, 86±4 µg/ml). The data represents the mean ratios for apolipoproteins of [SF concentration]/(mean plasma concentration in panel of unmatched normals donors]. OA, osteoarthritis; RA, rheumatoid arthritis. * P < 0.001 vs. apo B and apo A-I.

Table II. Concentrations of Apo E, Cholesterol, Triglycerides and Other Apolipoproteins in Synovial Fluids Relative to Matched Plasmas

<table>
<thead>
<tr>
<th>Patient's diagnosis</th>
<th>Total cholesterol mg/dl</th>
<th>Triglycerides mg/dl</th>
<th>Apo B µg/ml</th>
<th>Apo A-I µg/ml</th>
<th>Apo E µg/ml</th>
</tr>
</thead>
<tbody>
<tr>
<td>OA</td>
<td>41/127 (0.32)</td>
<td>19/108 (0.18)</td>
<td>123/597 (0.21)</td>
<td>293/1,050 (0.28)</td>
<td>5.1/11.1 (0.46)</td>
</tr>
<tr>
<td>OA</td>
<td>169/232 (0.73)</td>
<td>51/154 (0.33)</td>
<td>458/829 (0.55)</td>
<td>521/1,024 (0.51)</td>
<td>7/12.4 (0.56)</td>
</tr>
<tr>
<td>OA</td>
<td>42/252 (0.17)</td>
<td>3/201 (0.01)</td>
<td>155/1,190 (0.13)</td>
<td>247/1,550 (0.16)</td>
<td>8.8/33.6 (0.26)</td>
</tr>
<tr>
<td>OA</td>
<td>64/218 (0.29)</td>
<td>39/134 (0.29)</td>
<td>&lt;1/860 (0)</td>
<td>53/1,518 (0.03)</td>
<td>6.3/16.2 (0.39)</td>
</tr>
<tr>
<td>Acute gout</td>
<td>118/183 (0.64)</td>
<td>80/147 (0.54)</td>
<td>509/937 (0.54)</td>
<td>973/1,545 (0.63)</td>
<td>19.5/20.5 (0.95)</td>
</tr>
<tr>
<td>RA</td>
<td>16/71 (0.45)</td>
<td>4/79 (0.30)</td>
<td>164/76 (0.22)</td>
<td>561/1,24 (0.37)</td>
<td>6.9/4.6 (0.68)</td>
</tr>
<tr>
<td>Reiter's</td>
<td>119/127 (0.75)</td>
<td>50/79 (0.63)</td>
<td>370/1,96 (0.34)</td>
<td>576/1,51 (0.38)</td>
<td>8.1/10.1 (0.80)</td>
</tr>
<tr>
<td>RA</td>
<td>76/137 (0.55)</td>
<td>16/71 (0.23)</td>
<td>180/654 (0.28)</td>
<td>339/839 (0.40)</td>
<td>5.3/8.7 (0.61)</td>
</tr>
<tr>
<td>RA</td>
<td>130/210 (0.62)</td>
<td>68/161 (0.42)</td>
<td>368/947 (0.41)</td>
<td>1055/1,71 (0.62)</td>
<td>11.3/16.2 (0.70)</td>
</tr>
<tr>
<td>RA</td>
<td>87/146 (0.60)</td>
<td>38/89 (0.43)</td>
<td>293/721 (0.41)</td>
<td>751/978 (0.77)</td>
<td>7.8/11.5 (0.68)</td>
</tr>
<tr>
<td>RA</td>
<td>119/187 (0.64)</td>
<td>31/73 (0.42)</td>
<td>312/755 (0.41)</td>
<td>832/1,95 (0.43)</td>
<td>15/29 (0.52)</td>
</tr>
<tr>
<td>RA</td>
<td>78/164 (0.48)</td>
<td>24/94 (0.26)</td>
<td>282/872 (0.32)</td>
<td>457/1,52 (0.30)</td>
<td>13.6/16 (0.76)</td>
</tr>
<tr>
<td>Mean ratio</td>
<td>(0.52±0.17)</td>
<td>(0.34±0.16)</td>
<td>(0.32±0.15)</td>
<td>(0.40±0.20)</td>
<td>(0.61±0.18)*</td>
</tr>
</tbody>
</table>

Plasma samples were drawn from each individual at the time of arthrocentesis. OA, osteoarthritis; RA, rheumatoid arthritis. * P < 0.001 vs. triglycerides and apo B, P < 0.01 vs. apo A-I, P < 0.05 vs. cholesterol.

These results suggested that apo E was enriched relative to apo B and apo A-I in synovial fluids. Because apo B and apo A-I would be expected to enter synovial fluids exclusively by filtration from plasma (35, 36) these observations pointed to the possibility that a fraction of synovial fluid apo E was locally produced.

Further evidence for local articular synthesis of apo E. Apo E, newly secreted as a constitutive product of a number of extrahepatic tissues, including cells of the monocyte/macrophage lineage (16), is heavily sialylated and, in SDS-PAGE, migrates distinctly (17) from the apo E of circulating plasma lipoproteins, which is largely desialylated (16, 17). We observed that a fraction of apo E in osteoarthritic synovial fluids migrated with an elevated apparent molecular weight of 36–37 kD upon SDS-PAGE, comparable with that of apo E of THP-1 cell origin (Fig. 4). Treatment of whole synovial fluids with neuraminidase (16) resulted in a reduction of molecular weight of all synovial fluid apo E to 34 kD on SDS-PAGE (Fig. 4). Thus, synovial fluid apo E included a 36–37 kD fraction that was comparable to other apo Es of known extrahepatic origin, including apo E of monocyte/macrophage origin.

Monocyte/macrophage lineage cells (along with fibroblast-like cells) constitute the cells (synoviocytes) lining the inner surface of synovium (33). In addition, macrophages are normal constituents of synovial fluid (37) and are known to be present in cultures of synovial lining cells in early passages (33). Thus, cells isolated from the synovium of a patient with rheumatoid arthritis were studied in first passage in tissue culture. After 7 d the release of a small amount of apo E into the serum-free media was detectable by immunoassay (158.4 ng/100,000 cells). This further supported evidence for local intraarticular synthesis of apo E.

Qualitative analysis of apo E–bearing particles in synovial fluids. We observed that the molecular weight of synovial fluid apo E on SDS-PAGE was heterogeneous (Fig. 4), suggesting heterogeneity of the cellular origin of the apo E in synovial fluid. To evaluate if the particles bearing apo E in synovial
fluids also were heterogeneous, we utilized nondenaturing polyacrylamide gradient gel electrophoresis (31) and agarose gel electrophoresis (13) followed by Western blotting (Figs. 2 and 5). These methods were employed in preference to density gradient ultracentrifugation because artificial dissociation of apo E from lipoproteins can be induced by ultracentrifugation (38, 39). To minimize the contribution of filtered plasma lipoproteins to the content of apo E-bearing particles we studied noninflammatory synovial fluids and found apo E there to be associated with particles of a broad range of sizes (Fig. 2). These included particles, equivalent in size to proteins > 669 kD, that comigrated with apo B–lipoproteins on nondenaturing polyacrylamide sizing gels (Fig. 2) and particles that were equivalent in size to proteins of ~ 500 and 230 kD, and that comigrated with HDL particles (Fig. 2). In contrast, apo B–bearing particles in synovial fluid were universally > 669 kD, and were entirely excluded from the sizing gels (Fig. 2).

The absence of free apo E (34 kD) on sizing gels (Fig. 2) suggested that apo E in synovial fluid could be associated with lipoproteins. This observation was supported by the demonstration that synovial fluid apo E floated at a density of < 1.25 g/ml when synovial fluid was subjected to density gradient ultracentrifugation (data not shown). In addition, synovial fluid apo E migrated with lipoproteins into nondenaturing agarose gels (13) (Fig. 5), where the apo E–bearing lipoproteins were seen to be predominantly alpha (comigrating with HDL) and pre-beta (intermediate in charge between HDL and LDL) (Fig. 5).

The physiological significance of local anticular apo E synthesis likely includes the role of apo E in membrane resynthesis, as illustrated in repair of neural injury (15, 19). Potent suppression by apo E of lymphocyte activation by mitogens and antigens is also well recognized (20–23), and emphasizes that apo E can possess tissue-specific paracrine functions (48). Apo E secretion from monocyte/macrophages is increased by cholesterol loading (49), suggesting the possibility that synovial apo E production could be enhanced by the exposure of these cells to cellular debris, platelets (50), and intraarticular, oxidized LDL (51). Importantly, macrophage apo E secretion is suppressed by endotoxin (52) and certain particulates (53), suggesting that modulation of apo E secretion is one mechanism for macrophages to relate their state of activation to neighboring lymphocytes (14, 22). Thus, the extent of local apo E pro-

**Figure 4.** Elevated molecular weight of a fraction of synovial fluid apo E. Samples (0.015 ml) from two different osteoarthritis patients (Table II, patients No. 3 and 4), representative of four patients studied, were electrophoresed in a 13% SDS-PAGE gel, electrophoretically transferred to nitrocellulose and probed for apo E. Apo E of THP-1 cell origin served as an internal standard (left); serum or plasma from the patients (lanes 1 and 4); synovial fluids (lanes 2 and 5); synovial fluids after neuraminidase digestion (lanes 3 and 6).

**Figure 5.** Charge distribution of apo E–bearing particles in synovial fluids. Samples from a patient with osteoarthritis (Table II, patient No. 3) were electrophoresed on agarose strips under nondenaturing conditions, developed in barbital buffer, and probed for apolipoproteins after transfer by capillary filtration to nitrocellulose as described in Methods. Arrow indicates origin. Representative of results from three separate osteoarthritis patients.
Leukocyte activation by urate crystals is important in the pathogenesis of gouty inflammation (1, 4). The results of this study suggest that the binding of apo E to intraarticular monosodium urate crystals (Fig. 1) may have functional consequences via the potent suppression of the capacity of urate crystals to stimulate neutrophils (Fig. 3). Apo B-100 also inhibits urate crystal-induced neutrophil activation by binding to the crystal surface (13). Apo E, like apo B-100, possesses domains enriched in positively charged amino acids, and some of these regions mediate high-affinity binding to polyanionic glycosaminoglycans such as heparin (42, 43, 54, 55). Because urate crystals are also polyanionic (5), we are investigating the possibility that one or more of these regions could modulate the binding of apo E to urate crystals and mask areas on the crystal surface necessary to induce neutrophil activation. In addition, we are evaluating the potential for apo E to affect leukocyte activation by other inflammatory surfaces.

In summary, human synovial fluids contain apo E that appears to be partly of local origin. The potential functions of apo E in normal and inflamed joints, and in the regulation of the inflammatory potential of the urate crystals in gouty tophaceous deposits and synovial fluids merit further investigation.

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