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Mast Cells Contribute to Autoimmune Inflammatory Arthritis via Their Tryptase/Heparin Complexes

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Although mast cells (MCs) often are abundant in the synovial tissues of patients with rheumatoid arthritis, the contribution of MCs to joint inflammation and cartilage loss remains poorly understood. MC-restricted tryptase/heparin complexes have proinflammatory activity, and significant amounts of human tryptase \( \beta \) (hTryptase-\( \beta \)) are present in rheumatoid arthritis synovial fluid. Mouse MC protease-6 (mMCP-6) is the ortholog of hTryptase-\( \beta \), and this serine protease is abundant in the synovium of arthritic mice. We now report that C57BL/6 (B6) mice lacking their tryptase/heparin complexes have attenuated arthritic responses, with mMCP-6 as the dominant tryptase responsible for augmenting neutrophil infiltration in the K/BxN mouse serum-transfer arthritis model. While inflammation in this experimental arthritis model was not dependent on protease-activated receptor-2, it was dependent on the chemokine receptor CXCR2. In support of the latter data, exposure of synovial fibroblasts to hTryptase-\( \beta \)/heparin or mMCP-6/heparin complexes resulted in expression of the neutrophil chemotactic factors CXCL1/KC, CXCL5/LIX, and CXCL8/IL-8. Our proteomics, histochemistry, and immunohistochemistry data also revealed substantial loss of cartilage-derived aggrecan proteoglycans in the arthritic joints of wild-type B6 mice but not mMCP-6-null B6 mice. These observations demonstrate the functional contribution of MC-restricted tryptase/heparin complexes in the K/BxN mouse arthritis model and connect our mouse findings with rheumatoid arthritis pathophysiology. *The Journal of Immunology, 2009, 182: 647–656.

Rheumatoid arthritis (RA) is an autoimmune disease whose pathophysiology involves multiple cellular lineages and inflammatory processes that conspire to produce synovial inflammation, joint pain, joint swelling, and ultimately destruction of cartilage and bone. Because the synovium in RA patients is often infiltrated with lymphocytes and macrophages, many groups have focused their attention on the roles of these immune cells in the pathology of the autoimmune disease (1). Less widely appreciated is the marked increase in mast cell (MC) numbers in the synovial sublining of RA patients (2–10). Indeed, numerous groups reported that joint tissue isolated from RA patients contains 6- to 25-fold more human tryptase \( \beta \) (hTryptase-\( \beta \))-expressing MCs than does normal synovial tissue. In some instances, MCs comprise up to 5% of the nucleated cells in the arthritic synovium. This cellular increase in the chronically inflamed joint raises the possibility that MCs have switched from a beneficial physiologic role (e.g., as sentinel cells that protect the vulnerable synovial cavity from infectious organisms) to active participants in the immunopathology of joint inflammation and/or destruction (11). Indeed, MCs tend to localize at the junction of the pannus and cartilage, as well as in areas where the pannus is invading cortical bone (2, 12, 13). Moreover, rapid degradation of cartilage-derived aggrecan proteoglycans occurs when tryptase-expressing MCs are induced to release their granule proteases in MC/chondrocyte and MC/cartilage co-cultures (14). Nevertheless, the definitive contribution of MCs and their varied mediators in human RA remains enigmatic due to the fact that no MC-specific therapeutic intervention has been developed.

Although WBB6F1-Kit\(^{W/V}\)/Kit\(^{W/V}\) (W/W) and WCB6F1-Kit\(^{W}\)/Kit\(^{W}\)-null mice possess many non-MC-dependent disorders that greatly complicate data interpretation, numerous studies conducted on these respective Kit- and Kit ligand-defective mice have implicated prominent contributions of MCs to the pathophysiology of autoimmune inflammatory arthritis. For example, van der Broek and workers (18) noted that cartilage erosion in the knee was significantly reduced in W/W mice 14–35
days after these animals were sensitized with intradermal methylated BSA in the presence of adjuvant followed by intraarticular methylated BSA challenge. Although inflammation is a prominent feature of the methylated BSA arthritis model, swelling of knee joints is minimal in contrast to other experimental arthritis models involving distal peripheral joints. In the K/BxN mouse serum-transfer model in which mice are given pathogenic autoantibodies to glucose-6-phosphate isomerase, adoptive transfer of in vitro-differentiated Kit/tryptase-expressing wild-type (WT) MCs into W/W<sup>+</sup> mice restored arthritic susceptibility (15). Further studies in this animal model revealed that MCs are activated by IgG Fc receptors and can contribute to joint inflammation by elaborating IL-1 (19). Although the latter study demonstrated that MC-derived IL-1 was important in the initiation phase of the disease, a mediator that is more restricted to MCs must contribute significantly to the pathogenesis of autoimmune arthritis to explain the perceived MC dependence that occurs in W/W<sup>+</sup> mice due to the fact that macrophages and other cell types in the joint also express IL-1.

Tetramer-forming tryptases are selectively expressed in the MCs of every examined mammal, including mice (20, 21) and humans (22–24). The two tetramer-forming tryptases in mice are mouse MC protease (mMCP)-6 and mMCP-7. Their genes reside adjacent to one another on chromosome 17A3.3 (25), and mMCP-6 and mMCP-7 form heterotypic and homotypic tetramers (26). The human ortholog of mMCP-6 is hTryptase-β, and both serine proteases are sequestered in the granules of MCs ionically bound to heparin-containing serglycin proteoglycans. Indeed, the packaging of MC neutral proteases in the cytoplasmic granules of MCs is highly dependent on serglycin proteoglycans that contain heparin glycosaminoglycans (27, 28), and the biosynthesis of heparin in these cells is controlled by <i>NDST-2</i> (29). Heparin stabilizes the tryptase tetramer unit (29, 30) and restricts the enzyme’s substrate specificity (31). These tryptase/heparin complexes are exocytosed into the surrounding microenvironment in a regulated fashion upon MC activation (reviewed in Refs. 32, 33). MC subpopulations in mice and humans differ in their expression of tryptases and other granule proteases (34). Relevant to our study, human and mouse synovial MCs store abundant amounts of hTryptase-β and mMCP-6, respectively, in their secretory granules (7, 8, 13, 35, 36). The observation that IL-33 induces MC-committed progenitors to increase their expression of hTryptase-β (37) is significant considering the recent finding that IL-33 exacerbates collagen-induced arthritis in a MC-dependent manner (38). The accumulated data raise the possibility that MC-restricted tryptases have prominent roles in RA and in experimental arthritis.

Recent studies demonstrated functional roles for MC-restricted tryptases in innate immunity. Of particular relevance to K/BxN mouse experimental arthritis that is dependent on neutrophils (39–41), mMCP-6<sup>-/-</sup>/mMCP-7<sup>-/-</sup> (6/7<sup>-/-</sup>) C57BL/6 (B6) mice demonstrated decreased neutrophil recruitment and reduced survival after injection of the peritoneal cavity with <i>Klebsiella pneumoniae</i> (42). In support of these data, administration of recombinant mMCP-6 or hTryptase-β into the peritoneal cavity or lungs of normal mice resulted in a marked influx of neutrophils (30, 31, 43, 44). The contribution of MC-restricted tryptases to the accumulation of peripheral blood neutrophils into the arthritic joint could be due to their ability to stimulate the expression and release of neutrophil-specific chemotactic factors from nearby bystander cells. In this regard, we and others showed that tryptases can induce cultured human epithelial and endothelial cells to increase their expression of CXCL8 when exposed to a partially purified preparation of synovium-derived hTryptase-β (47). Other possible mechanisms by which MC-restricted tryptases could contribute to inflammatory pathogenesis include activation of matrix metalloproteinases (48–50), cleavage of extracellular matrix components (51), stimulation of fibroblast proliferation and collagen production (52), and proteolysis of fibrinogen (53). Indeed, MC proteases have been demonstrated to contribute to cartilage degradation in vitro in the absence of the inflammatory cells present in the arthritic joint (14).

Having previously demonstrated the presence of mMCP-6 protein in the synovial MCs of healthy and diseased mice (35), we hypothesized that the elaboration of this serine protease and/or its closely related family member mMCP-7 could represent a MC-specific contribution to the proinflammatory milieu of the joint in the K/BxN mouse serum-transfer arthritis model. The creation of B6 mice that differ in their expression of mMCP-6, mMCP-7, and NDST-2 allowed us the opportunity to evaluate the potential roles of MC-restricted tryptase/heparin proteoglycan complexes in experimental arthritis. Utilizing the autoimmune K/BxN mouse serum-transfer model of experimental arthritis and our different transgenic mouse strains, we now report that MC-restricted mMCP-6/heparin proteoglycan complexes serve as important intensifiers of inflammation within the joint via enhancing the recruitment of neutrophils in a protease receptor-2 (PAR-2)-independent manner. Furthermore, we show that enzymatically active recombinant mMCP-6 induces cultured mouse FLS to increase their expression of CXCL1/KC/GRO-α and CXCL5/LIX/ENA-78, and we demonstrate that mice lacking CXCR2 develop less joint inflammation. Finally, we show that mMCP-6/heparin proteoglycan complexes contribute to joint destruction by promoting the degradation of the aggrecan proteoglycans in the diseased cartilage.

**Materials and Methods**

**Mice**

WT mMCP-6<sup>-/-</sup>/mMCP-7<sup>-/-</sup> (6/7<sup>-/-</sup>) (54) B6 mice and CXCR2<sup>-/-</sup> BALB/c mice (55) were obtained from The Jackson Laboratory. PAR-2<sup>-/-</sup>/B6 mice (56) were a kind gift from Dr. Shaun Coughlin (University of California at San Francisco, San Francisco, CA). These mice and NDST-2<sup>-/-</sup> (10-generation backcrosses with B6 mice) (27), 6/7<sup>-/-</sup> (generated in B6 embryonic stem cells followed by 8-generation backcrosses with B6 mice) (42), and mMCP-6<sup>-/-</sup>/mMCP-7<sup>-/-</sup> (6/7<sup>-/-</sup>) (10-generation backcrosses with B6 mice) (57) mice were maintained under specific pathogen-free conditions. Although most examined WT mice express mMCP-6 and mMCP-7, the MCs in WT B6 mice constitutively lack mMCP-7 (58) due to a splice-site joint mutation in this gene that causes rapid catalolism of the defective transcript by the “nonsense” posttranscriptional pathway (54). K/BxN mice were maintained as described (35). All mice were sex and gender matched, and experiments were conducted using animal protocols approved by the Dana-Farber Cancer Institute/B Brigham and Women’s Hospital Animal Care and Use Committee.

**K/BxN mouse serum-transfer arthritis**

Arthritogenic K/BxN mouse serum containing autoantibodies to glucose-6-phosphate isomerase was administered to recipient mice using a modification of a previously described experimental approach (59). To induce arthritis, mice received two 150-μl i.p. injections of diluted (50 μl serum + 100 μl sterile PBS) serum on experimental days 0 and 2. Clinical indices were recorded at 24- to 48-h intervals, and indices were graded on a scale of 0 to 12 as described (15). Ankle thickness was measured using a spring-loaded dial thickness gauge (Long Island Indicator Service). Measurements were made axially across the ankle joint, with the ankle in full dorsiflexion.

**Histochemistry, enzyme cytochemistry, and immunohistochemistry of the arthritic joint**

Ankles were harvested as previously described (35). Briefly, the skin around each ankle was gently removed, and tissue from the distal one-third
of the tibia to the midpaw was collected. After fixation with 4% paraformaldehyde, ankles were decalcified in Kristen’s solution (60), dehydrated, and embedded in paraffin. Tissue was cut into 4 µm sections, and midsagittal ankle sections were stained with H&E. Histological grading of synovial inflammation and cartilage and bone erosion in the ankle sections midsagittal ankle sections were stained with H&E. Histological grading of synovial inflammation and cartilage and bone erosion in the ankle sections 

Mass spectrometry analysis of abundant proteins in arthritic joints

Detergent lysates of joint tissues from hindpaws of experimental mice were prepared by mechanical disruption, as previously described (40). Twenty-five micrograms of protein extract obtained from each mouse ankle joint was fractionated on a 4–10% Tris-HCl polyacrylamide gradient gel (Bio-Rad) run for 90 min at 130 V with Tris-glycine-SDS buffer (Bio-Rad). All gels were washed and stained with Coomassie blue G250 (Bio-Rad) for 20 min and then destained in deionized water. A total of 12 gel slices (5 mm²) were excised from each lane in the gel. Individual gel slices were placed in 0.6-ml microfuge tubes (Axygen Scientific) and incubated at room temperature for 10 min in 25 mM NH₄HCO₃, and then in 50% acetonitrile/25 mM NH₄HCO₃. This cycle was repeated twice. After the acetonitrile solution was removed, the dried gel slices were covered with 10 mM DTT and incubated at 56°C for 45 min and then at room temperature for an additional 20 min. DTT was discarded and alkylation was performed by adding 55 mM iodoacetamide (MP Biomedicals) to cover the gel slice. All slices were incubated with iodoacetamide for 45 min at room temperature in the dark. Iodoacetamide was discarded and the gel slices were washed twice with 25 mM NH₄HCO₃ followed by 50% acetonitrile/25 mM NH₄HCO₃ as described above. Trypsin (0.150 µg; Promega) was added, along with sufficient 50 mM NH₄HCO₃ to cover the slice, and the sample was incubated overnight at 37°C to generate the tryptic fragments of the unknown protein present in the arthritic joints. Formic acid was added to a final concentration of 1% to stop the reaction. The aqueous solution (peptide fraction) was transferred to a clean microtube. To the gel pieces, 50% acetonitrile/5% formic acid was added; the gel slices were vortexed 30 min and then sonicated for 5 min. The aqueous solution was transferred to the original peptide solution and the above step repeated. The pooled peptide sample was speed vacuum dried to reduce the volume to 10 µl.

The tandem mass spectroscopy (MS/MS) data for the unknown protein of interest in the individual gel slices were obtained using an LTQ mass spectrometer (Thermo Fischer Scientific). The tryptic digests were first run on a pre-column (C18, PepMap100) and separated with a reverse-phase C18 column using mobile phase A (0.1% formic acid in water) and B (84% acetonitrile, 0.1% formic acid in water) with a linear gradient of 2% per min, starting with 100% of A. The peptides were then injected at a flow rate of 300 nl/min. MS/MS spectra were then acquired by the Finnigan LTQ.
The resulting raw data were analyzed using Mascot (version 2.1.03, Matrix Science) and searched against the NCBI (National Center for Biotechnology Information) mouse protein database. Searches were performed with fixed carbamidomethylation of cysteines and variable oxidation of methionine residues. A fragment tolerance of 0.4 Da was used in searches.

CXCL1, CXCL5, and CXCL8 expression in tryptase-treated FLS

Human and mouse FLS were generated as described (63, 64). In brief, human synovial tissue from RA patients was obtained as discarded specimens from joint arthroplasty surgery with approval of the Partners Institutional Review Board. Synoviocyte cell suspensions were prepared from synovial tissues by mincing, followed by gentle rocking for 1–2 h at 37°C in DMEM medium (Invitrogen) containing 1 mg/ml type IV collagenase (Worthington Biochemical) and 0.1 mg/ml DNase (Sigma-Aldrich). Mouse FLS isolation was performed similarly under procedures approved by the Dana-Farber Cancer Institute/Brigham and Women’s Hospital Animal Care and Use Committee. After euthanasia, mouse ankle joints were isolated by disarticulation distal to the tibia and at the tarsal/metatarsal junction, taking care not to violate the marrow space. The tissue was infiltrated with collagenase in DMEM and incubated with gentle rocking for 2 h, as described above. The resultant cell suspension was passed through a 100-μm cell strainer (BD Biosciences) and placed in a tissue culture flask (BD Labware) containing DMEM (Invitrogen) supplemented with 10% heat-inactivated FBS (HyClone), 2 mM l-glutamine, 100 U/ml penicillin, 100 μg/ml streptomycin, 50 μM 2-ME, and 0.1 mM nonessential amino acids (Invitrogen). Since no bone marrow-derived lineage cells were detected after passage 3 (63, 64), cultured FLS were used in our experiments between passages 4 and 8. For the in vitro stimulation studies, FLS were cultured to 80–90% confluence in 12-well plates (Corning) for 48–72 h. The resulting cells were exposed to recombinant hTryptase-β-heparin complexes (Promega) or recombinant pro- and mature mMCP-6/heparin complexes (31) in culture medium containing 1% FCS for 16 h, whereupon supernatants were collected for analysis. As a positive control for chemokine induction, replicate FLS were exposed to TNF-α. Commercial ELISA kits (R&D Systems) were used to quantify the levels of human CXCL8, mouse CXCL1, and mouse CXCL5 in the supernatants.

Statistical analysis

Data were analyzed using Prism software, version 4.00 (GraphPad Software). Cell numbers of respective lineages were described as the means ± SE. Student’s t test was performed to compare changes in cell numbers, histological analysis, and ELISA values. Curves of both indices plotted from arthritis experiments were compared using the two-way ANOVA test. p values <0.05 were considered statistically significant.

Results

Impaired autoimmune inflammatory arthritis in heparin-deficient B6 mice

Since our previous studies demonstrated that the synovium of the arthritic mouse is populated by MCs that presumably contain heparin proteoglycans due to their histochemical properties, we began our assessment of the contribution of MC-restricted tryptase/heparin proteoglycan complexes to joint inflammation in the K/BxN arthritic mouse is populated by MCs that presumably contain heparin proteoglycans due to their histochemical properties, we began our assessment of the contribution of MC-restricted tryptase/heparin proteoglycan complexes to joint inflammation in the K/BxN mouse CXCL1, and mouse CXCL5 in the supernatants. As assessed by the Limulus amebocyte lysate kit (Associates of Cape Cod), the tryptase/heparin preparations used in our study contained <0.25 EU LPS/ml.

FIGURE 2. Decreased arthritis in treated mMCP-6-deficient B6 mice. A, Immunohistochemical staining for mMCP-6 and mMCP-7 expression in the synovial tissue MCs of WT 6/7 and transgenic 6/7 and 6/7’ B6 mice. Bar, 10 μm. B and C, Arthritis severity in WT 6/7 and transgenic 6/7 and 6/7’ B6 mice. Arrows indicate when the mice received arthritogenic K/BxN mouse serum. Data were pooled from three independent experiments (n = 15 mice/group, p < 0.01). D, Histomorphometric quantification of articular tissue. The data are the mean values ± SEM (n = 15/group).

Impaired arthritis in tryptase-deficient B6 mice

Considering the profound defect in protease expression in the synovial MCs of NDST-2/−/− B6 mice (Fig. 1A), we proceeded to assess the contributions of mMCP-6 and mMCP-7 in the arthritis model using WT 6/7 B6 mice and newly created transgenic 6/7’ and 6/7 B6 mice. As noted in Fig. 1B, we found an ~50% decrease in clinical measurements of arthritis in these transgenic mice. The day 10 Δ ankle thickness was 0.67 ± 0.05 mm and 0.38 ± 0.06 mm in diseased WT and NDST-2/−/− B6 mice, respectively (p = 0.0008; Fig. 1B). Histomorphometric quantification of the arthritic changes in the joint tissues confirmed the clinical assessment, with significant decreases in inflammation and bone erosion scores in the NDST-2/−/− B6 mice (Fig. 1, C and D).
Histomorphometric quantification of the tissue arthritic response again confirmed our clinical measurements.

Because the neutrophil accumulation that occurs in the peritoneal cavity of *K. pneumoniae*-infected mice depends on mMCP-6, we next attempted to define the potential contribution of the tryptase to joint inflammation in the K/BxN mouse serum-transfer model. To address this issue, we compared 6/H11002 and 6/H11002/7 B6 mice. Interestingly, we found that the arthritis severity in 6/H11002/7 B6 mice was indistinguishable from that in 6/H11002 B6 mice (Fig. 2, B and C). The synovium of 6/H11001, 6/H11002, and 6/H11002/7 B6 mice contained 47.2 ± 3.1, 51.6 ± 4.7, and 48.9 ± 3.6 MCs/mm², respectively. Because no significant difference in synovial MC density was observed among the three populations of B6 mice, the reduced arthritis seen in both mMCP-6-deficient mouse strains was not a consequence of fewer MCs. The accumulated data suggest that mMCP-6 is the dominant proinflammatory tryptase in the K/BxN mouse serum-transfer arthritis model.

Characteristics of the cellular infiltrate in the synovium induced by tryptase/heparin complexes

Because large numbers of neutrophils are recruited into the joints of WT mice given proarthritic serum, we examined neutrophil recruitment as a potential mechanism by which MC-restricted mMCP-6/heparin complexes promote synovial inflammation. We quantitated synovial leukocyte populations in the arthritic joints of mMCP-6-deficient and mMCP-6-sufficient B6 mice at day 10 (Fig. 3). These histomorphometric studies revealed that the density of neutrophils in the diseased synovium of NDST-2-null or the tryptase-deficient 6/7 and 6/7 B6 mice was substantially reduced relative to that in WT B6 mice (Fig. 3, A and C). Since mMCP-6 also can regulate eosinophil infiltration in the skeletal muscle of *Trichinella spiralis*-infected mice, we next quantified the density of eosinophils in the diseased synovium. We found that the arthritic joints of WT, heparin-deficient, and tryptase-deficient mice contained similar numbers of eosinophils (Fig. 3B). In all arthritic mice, the number of neutrophils greatly exceeded the number of eosinophils. Thus, in the arthritic joint, mMCP-6/heparin complexes preferentially induce the recruitment of neutrophils.
Development of arthritis is dependent on the chemokine receptor CXCR2 but not on PAR-2/F2RL1

Based on earlier in vitro studies, numerous tryptase substrates of potential relevance to synovial inflammation have been identified. Among these potential substrates, PAR-2 (65) has been most extensively investigated. Although conflicting data have been reported pertaining to the role of this protease-activated receptor in experimental arthritis (66 – 68), a recent study concluded that PAR-2-null mice developed arthritis when given a high-dose of K/BxN mouse serum (68). Consistent with the findings of Busso and coworkers (68), we found no difference in arthritic responses when PAR-2-deficient and PAR-2-sufficient B6 mice were given lesser amounts of the proarthritic serum (Fig. 4, B and C). These data imply that mMCP-6/heparin complexes regulate neutrophil accumulation in the arthritic joints of our experimental model in a PAR-2-independent manner.

A number of related Glu-Lys-Arg CXCL chemokines (e.g., CXCL1, CXCL2/GRO-, CXCL3/GRO-/H9252, CXCL5, CXCL7/NAP-2, and CXCL8) regulate neutrophil accumulation in tissues via the chemokine receptor CXCR2. Because mMCP-6/heparin proteoglycan complexes contribute to the accumulation of neutrophils in the arthritic joint (Fig. 3), we next evaluated arthritis in CXCR2 mice. As noted in Fig. 4A, inflammation was markedly reduced in these mice, suggesting that Glu-Lys-Arg CXCL chemokines and their receptors contribute to neutrophil extravasation in this experimental arthritis model.

Tryptase-induced chemokine production in FLS

Tryptase is not directly chemotactic for neutrophils (31), and exocytosed mMCP-6/heparin proteoglycan complexes are retained in connective tissues for hours due to their large size (69). mMCP-6 exocytosed from synovial MCs therefore cannot come in direct contact with peripheral blood neutrophils to induce their extravasation into the diseased joint. Because our experimental arthritis model is partly dependent on CXCR2 (Fig. 4A), we postulated that mMCP-6 and its human ortholog induce granulocyte accumulation in an indirect manner, probably by inducing the expression of chemokines that bind to CXCR2. FLS are the primary cells of mesenchymal lineage in synovial tissue, and these cells predominate in rheumatoid pannus tissue. Indeed, stimulation of mouse FLS in vitro with recombinant mMCP-6/heparin complexes resulted in increased expression of the CXCR2 ligands CXCL1 (Fig. 5A) and CXCL5 (Fig. 5B) to levels comparable to those obtained when

FIGURE 5. Tryptase/heparin complexes induce human and mouse FLS to express the chemokines CXCL1 (mouse), CXCL5 (mouse), and CXCL8 (human). Primary mouse (A and B) or human (C) FLS (passage 4 – 8) were exposed to enzymatically active recombinant hTryptase-β-heparin or mMCP-6/heparin complexes. The resulting conditioned media were assayed for their levels of the chemokines via ELISA. The data are the mean values ± SEM. Results shown are representative of three independent experiments. As negative controls, replicate human FLS were exposed to boiled hTryptase-β and mouse FLS to pro-mMCP-6. As positive controls, both populations of FLS were exposed to TNF-α.

FIGURE 6. Identification of a soluble fragment of aggrecan proteoglycan in the arthritic joints of WT B6 mice. Mouse aggrecan (GenBank accession no. AAC37670) is initially translated as a 2132-mer protein that contains a 19-mer signal peptide (77). The proteoglycan’s N-terminal globular domains bind to HA. The HA-binding region in normal cartilage that can exceed 1 billion Da. Aggrecan’s HA-binding region is susceptible to numerous proteases. For example, the Glu392Ala393 peptide bond that is cleaved by aggrecanase-1/ADAMTS4 and aggrecanase-2/ADAMTS5 is highlighted in italic boldface. Highlighted in boldface are the locations of the three MS/MS identified peptides at residues 595–604, 2003–2026, and 2066–2075 in the isolated aggrecan fragment from the arthritic joints of WT B6 mice at days 8 and 15 after transfer of K/BxN mouse serum.
these mesenchymal cells encountered TNF-α. Interestingly, mMCP-6/heparin complexes elicited CXCL1 production in PAR-2/H11002/mouse FLS as well (data not shown). To extend our animal work into a human system, we next assessed the ability of recombinant hTryptase-heparin complexes to regulate the expression of CXCL8 in cultured human FLS. As noted in Fig. 5 C, we found that this neutral protease induced FLS to express significant amounts of CXCL8. The inability of recombinant pro-mMCP-6 (Fig. 5, A and B) or boiled hTryptase-heparin (Fig. 5 C) to induce the expression of significant amounts of CXCL1, CXCL5, or CXCL8 suggests that these serine proteases must be enzymatically active to optimally affect the joint. Given the fact that LPS is heat stable, the chemokine-inducing effects of the hTryptase-heparin preparations used in our study were not the result of LPS contamination, thereby supporting our data obtained with the inactive zymogens.

**Tryptase-dependent proteolysis and loss of aggrecan proteoglycans in experimental arthritis**

A proteomics approach was next used to identify longitudinal changes in abundant protein constituents between healthy joints and the arthritic joints in the K/BxN mouse experimental model. In these assays, the proteins were extracted from the joints of replicate WT B6 mice at baseline and on days 8 and 15 after arthritis induction. The resulting extracts were subjected to SDS-PAGE and gel slices containing differentially expressed proteins/peptides were subjected to MS/MS.

In this proteomics assay, a large-sized protein species was identified in the day 8 and day 15 arthritic samples of WT B6 mice that was not present in the control day 0 sample (data not shown). MS/MS analysis of three trypsin-generated peptides from the unknown protein revealed that it was aggrecan proteoglycan that had lost its protease-susceptible N-terminal hyaluronan (HA)-binding domain. The amino acid sequence of mouse aggrecan and the location of the three MS/MS identified peptides in the isolated aggrecan fragment are noted in Fig. 6. Based on our proteomics data, the proteolytic liberation of soluble aggrecan and concomitant loss of aggrecan proteoglycans from joint cartilage are major features of the K/BxN mouse model of arthritis, as occurs in RA patients. We therefore used histochemical and immunohistochemical approaches to compare amounts of aggrecan proteoglycans from the cartilage of WT B6 mice and our transgenic NDST-2−/−, 6−/7−, and 6−/7− B6 mice. As noted in Fig. 7, loss of aggrecan proteoglycans were markedly attenuated in the joints of our trypase/heparin-deficient B6 mouse strains in established arthritis.

**Discussion**

MCs have long been recognized as potential participants in human and mouse inflammatory arthritis. We recently identified an important contribution for MC-derived IL-1 in arthritis initiation in the K/BxN mouse serum transfer model (19). However, because many cell types produce this proinflammatory cytokine, it is likely that MCs contribute to synovial inflammation via multiple pathways. We therefore sought to identify a contribution of MCs to arthritis that is more restricted to this cell lineage. Of the diverse array of effector molecules exocytosed from activated MCs, their protease/serglycin proteoglycan complexes are unique to MCs. Our demonstration of a functional role for mMCP-6/heparin complexes in the K/BxN mouse model (Figs. 1–3, 6, and 7) now provides further insight into the roles of MCs and their protease mediators in experimental arthritis.

The identification of trypase/heparin complexes as contributors to arthritis is consistent with previous functional studies of these macromolecular complexes in mice and humans. Because synovial MCs are positioned near vessels and within the synovial mesenchyme (11, 70), they are ideally situated to amplify arthritogenic stimuli via bulk release of their preformed mediators. Tryptase is
present as an active enzyme pre-packaged tetramer in the cytoplasmic granules of human and mouse synovial MCs ionically bound to heparin-containing serglycin proteoglycans (32, 33). Although the safranin$^\text{+}$ population of MCs found in the mouse synovial lining preferentially expresses mMCP-6, all human MCs express its ortholog hTryptase-$\beta$, including the MCs in the human synovium. Previous in vivo studies employing either recombinant protein or bacterial infection of tryptase-deficient mice demonstrated that mMCP-6 and hTryptase-$\beta$ elicit neutrophil accumulation in tissues (30, 31, 35, 42–44). Neutrophils are abundant in the joints of mice given arthritogenic K/BxN serum (Fig. 3) and in the synovial fluid of humans with RA, and these granulocytes contribute to the pathogenesis of arthritis (39–41). We observed a significant decrease in the number of neutrophils in the arthritic synovium of three transgenic mouse strains that lack mMCP-6 (Fig. 3). These findings are consistent with a functional contribution to synovitis via the promotion of neutrophil recruitment into the arthritic joint. Our additional finding that hTryptase-$\beta$/heparin and mMCP-6/heparin complexes induce cultured FLS to increase their expression of CXCL chemokines. In this regard, it is now appreciated that mMCP-6 is important in the accumulation of eosinophils in the skeletal muscles. In this regard, it is now appreciated that mMCP-6 is important in the accumulation of eosinophils in the skeletal muscles of T. spiralis-infected mice (57). It also has been reported that recombinant hTryptase-$\beta$ can cleave CCL11/eotaxin and CCL5/ RANTES and therefore abrogate their chemotactic activities (71). We therefore considered the possibility that large numbers of eosinophils do not accumulate in the rheumatoid joint due to an additional tryptase-mediated inactivation of an induced CCL chemokine that preferentially attracts eosinophils. However, this does not appear to be the case in our experimental arthritis model due to the fact that we did not see increased numbers of eosinophils in the diseased joints of our treated 6/7 mice (Fig. 3).

Although the safranin$^\text{+}$ MCs in most mouse strains express mMCP-7, our studies failed to uncover a significant functional contribution for this tryptase in the K/BxN mouse serum-transfer model of experimental arthritis (Fig. 2). The $\alpha$-chain of fibrinogen is a preferred substrate of mMCP-7, and the injection of this tryptase into the peritoneal cavity of WT mice preferentially results in the recruitment of eosinophils (43, 53). As WT B6 mice are genetically deficient in mMCP-7 (54, 58) but susceptible to arthritis (Figs. 1 and 2), our congenic mouse strains that differed in their mMCP-6 and mMCP-7 expression allowed us to test for possible redundancy of tryptase activity. Our observation that mMCP-7 did not restore inflammation or granulocyte influx into the synovium of the diseased mice (Fig. 2) suggests that mMCP-6 is the dominant proinflammatory tryptase in the K/BxN mouse serum-transfer model of arthritis. These observations are consistent with findings in the methylated-BSA/JL-1 (mBSA) mouse model of experimental arthritis of the knee, which also displays significant reliance on mast cell tryptase/heparin complexes (72). The latter study confirms a key role for mMCP-6 in a very different type of experimental arthritis. Nevertheless, given redundant mMCP-6 and mMCP-7 function in the mBSA model, congenic 6/7 mice might be necessary to uncover the roles for the MC’s tryptase/heparin complexes in other physiologic and/or pathologic situations.

The receptors and/or extracellular matrix proteins in the synovium that are cleaved by enzymatically active mMCP-6 remain to be determined. PAR-2 is a member of the protease-activated receptor family expressed on the surfaces of endothelial cells and synovial fibroblasts, and this receptor is activated by trypsin. Mirza and coworkers (73) reported that recombinant hTryptase-$\beta$ can weakly activate cells via PAR-2. Nevertheless, using varied combinational approaches, it has been shown that recombinant mMCP-6 (31) and hTryptase-$\beta$ (43, 74) prefer to cleave amino acid sequences that do not resemble the trypsin-susceptible activation site in PAR-2 (65). The substrate specificity of mMCP-6 becomes even more restricted when this tryptase is ionically bound to heparin (31). Because Compton and coworkers (75) found that naturally occurring hTryptase-$\beta$ cannot activate PAR-2 if heparin is present, the importance of the proteoglycan cofactor may not have been adequately appreciated in some of the studies that concluded that mMCP-6 and/or hTryptase-$\beta$ function via PAR-2. Nevertheless, the availability of PAR-2-null mice allowed us to more definitively evaluate the role of this G protein-coupled receptor in the K/BxN mouse serum-transfer model of experimental arthritis. Our results revealed that PAR-2 is dispensable in this arthritis model (Fig. 4). Furthermore, recombinant mMCP-6 stimulated CXCL1 production in PAR-2-null mouse FLS. These observations are consistent with the findings of Busso and coworkers (68), who evaluated the role of PAR-2 in four different mouse models of arthritis.

A major feature of RA is loss of aggrecan proteoglycans due to proteolysis of the N-terminal domain that binds to HA. Using a proteomics approach, we identified a prominent soluble fragment of aggrecan proteoglycan lacking its HA-binding domain in the joints of arthritic WT B6 mice (Fig. 6). Employing histochemistry and immunohistochemistry approaches, we then discovered that the proteolytic loss of cartilage proteoglycans is markedly attenuated in our transgenic B6 mice that lack MC-restricted tryptase/heparin complexes (Fig. 7). While our data could be a consequence of fewer protease-rich neutrophils in the arthritic joints of our tryptase-deficient mice (Fig. 3), we previously noted that an undefined serine protease present in rat peritoneal MCs rapidly cleaves the N-terminal domains of rat and bovine aggrecan proteoglycans in vitro (14), thereby resulting in a large-sized fragment that cannot recognize HA as identified in Fig. 6. It is now known that the MCs used in that earlier study express the rat ortholog of mMCP-6 and hTryptase-$\beta$ (76). Because neutrophils and other cell types were not present in the Stevens et al. 1992 aggrecan-susceptibility study (14), it is possible that MC tryptases preferentially cleave the HA-binding domain of aggrecan in a direct manner. In support of this possibility, mouse aggrecan has a number of candidate tryptase-susceptible cleavage sites in its protease-susceptible N-terminal domain (e.g., the Pro-Ile-Val-Ser-Pro-Arg sequence at residues 546–551) (31, 43, 74) (Fig. 6). Alternatively, because pro-aggrecaenase-1/ADAMTS4 is activated by tryptic cleavage at a site favorable for cleavage by mMCP-6 and hTryptase-$\beta$, MC tryptases could participate in aggrecan loss in our experimental model by activating a metalloproteinase like ADAMTS4.

A number of observations suggest that the contribution to inflammatory arthritis from mouse MCs via their tryptase/heparin complexes may be pertinent in human RA. The numbers of tryptase-expressing MCs are dramatically elevated in the synovial tissue of many patients with RA. Thus, the cellular lineage, localization, and mediators are common between mouse and man. Since abundant neutrophil recruitment to the synovial fluid is a hallmark of RA, modulation of neutrophil recruitment also is an attractive
mechanism by which MC-restricted tryptase/heparin complexes could enhance disease activity. Furthermore, the contribution of the synovial mesenchyme to joint inflammation is increasingly appreciated (64). Thus, our observations that arthritis was reduced in CXCR2−/− mice (Fig. 4) and that exposure of human and mouse FLS to tryptase/heparin complexes results in the expression of multiple chemokines (Fig. 5) that regulate neutrophil recruitment and aggrecan loss, arthritis was more diminished in our NDST-2/heparin-null C57BL/6 mice than in our MC-restricted-null C57BL/6 mice. These data suggest the participation of another MC-restricted protease mediator (e.g., mMCP-5 and/or mMCP-6) in joint inflammation and/or destruction. We therefore anticipate that the role of MCs and their varied mediators in arthritis will be multifaceted, and that future investigations will provide further novel insights into their influence in the inflammatory disorder.

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Disclosures

The authors have no financial conflicts of interest.

References

TRYPTASE-DEPENDENT EXPERIMENTAL ARTHRITIS


