Hyaluronan production in human rheumatoid fibroblastic synovial lining cells is increased by interleukin 1 β but inhibited by transforming growth factor β 1

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Abstract

Objectives—To investigate the regulatory roles of interleukin 1 β (IL1 β), tumour necrosis factor α (TNF α), interferon γ (IFN γ) or transforming growth factor β 1 (TGF β 1) on hyaluronan (HA) synthesis by human fibroblastic synovial lining cells.

Methods—Concentrations of HA in culture supernatants of fibroblastic synovial lining cell line (RAMAK-1 cell line) with or without stimulation by IL1 β , TNF α , IFN γ or TGF β 1 were measured by sandwich binding protein assay. Levels of HA synthase mRNA of the cells with or without stimulation were detected by reverse transcribed polymerase chain reaction. Molecular weights of HA in the culture supernatants of the cells with or without stimulation were measured using high performance gel permeation liquid chromatography.

Results—HA synthesis by the cells was not significantly augmented by TNF α or by IFN γ . It was significantly stimulated by IL1 β but inhibited by TGF β 1. Molecular weights of HA in the culture supernatants of the cells were unchanged by stimulation with TNF α . They were remarkably increased by stimulation with IL1 β and IFN γ , but reduced with TGF β 1.

Conclusion—IL1 β is an up regulator of HA synthesis, while TGF β 1 is a down regulator. HA production in the synovial lining cells of inflamed joints (for example, rheumatoid arthritis) might be regulated by the balance of these cytokines.

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Hyaluronan (HA) is a linear glycosaminoglycan consisting of repeating units in which the sugars glucuronic acid and N-acetyl glucosamine alternate. It exists as the tertiary structure of a random coil with a high molecular weight. Concentrations of HA are highest in musculoskeletal tissues, skin, embryological tissues, and synovial fluid.¹ HA plays an important part in wound healing,² embryonic development,³ tumour growth,⁴ and synovial fluid viscosity.⁵

Cultured synovial fibroblasts produce HA.⁶ Immunohistochemical studies of normal synovium show that HA surrounds the lining layer cells but it is minimal in deeper layers.⁷ These

results indicate that one of the fundamental functions of fibroblastic synovial lining cells is to produce HA and release it into synovial fluid, and to maintain constitutive HA concentrations in synovial fluid.

The mechanism regulating the production of HA in synovial lining cells has not yet been fully elucidated, perhaps because the preparation of a pure synovial lining cell population from whole synovial tissues requires tedious and demanding procedures. Recently, we derived a fibroblastic synovial lining cell line (RAMAK-1 cell line) from a patient with RA.^{8,9} The cells form a cystic tumour containing lubricating fluid enclosed by a lining layer when injected subcutaneously into severe combined immunodeficient mice, which suggests that these cells produce HA in this synovial fluid-like liquid.

The purposes of this study are to determine the regulatory roles of interleukin 1 β (IL1 β), tumour necrosis factor α (TNF α), interferon γ (IFN γ) or transforming growth factor β 1 (TGF β 1) on HA production by the fibroblastic synovial lining cells.

Methods

CELL CULTURE

The fibroblastic lining cell line (RAMAK-1 cell line) was established as previously described.⁸ Cells were passaged every week in the ratio of 1:20. We used the cells at passages 20-25 in the subsequent studies.

MEASUREMENT OF HA CONCENTRATION IN THE CULTURE SUPERNATANTS

RAMAK-1 cells in subconfluent populations (5×10^4 cell/cm²) on culture dishes (Cluster 3512, Costar, Cambridge, MA) were washed with RPMI-1640 medium (Nipro, Tokyo) and further cultured with RPMI-1640 medium containing 10% fetal bovine serum (FBS; Intergen, Purchase, NY) for either 24, 48, 78 or 96 hours with or without stimulation by IL1 β (1 ng/ml; Otsuka, Tokushima, Japan), TNF α (300 units/ml; Teijin, Tokyo), IFN γ (1000 units/ml; Boehringer Mannheim Biochemica, Mannheim, Germany) or TGF β 1 (1 ng/ml; King Brewing, Kakogawa, Japan). Concentrations of HA in the supernatants of the cell culture were measured by a sandwich binding protein assay kit (Hyaluronan Plate Chugai, Chugai Pharmaceutica, Tokyo). Samples were incubated with HA binding protein coated on

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Figure 1 HA production by RAMAK-1 cells. RAMAK-1 cells were incubated for 24 or 48 hours in culture dishes with RPMI-1640 medium containing 10% FBS with or without stimulation by IL1 β (1 ng/ml), TNF α (300 units/ml), IFN γ (1000 units/ml) or TGF β 1 (1 ng/ml). At the end of culture, the concentration of HA in the supernatants was measured by sandwich binding protein assay.

microplates for one hour at room temperature. After incubation, the microplates were washed three times with wash buffer, and further incubated with 100 μ l of peroxidase labelled liquid phase HA binding protein for 30 minutes at room temperature. After incubation, the microplates were washed three times with wash buffer, and further incubated with 100 μ l of peroxidase reactant substrate for 30 minutes at room temperature in a darkroom. The reaction was stopped by the addition of 0.18 M sulphuric acid. The development of the colour was monitored at 450 nm with an Easy Reader ELISA analyser (SLT-Lab Instruments, Vienna, Austria). The HA level in each sample was calculated using the standard curve obtained with the purified HA solutions, which were included in the kit as references. According to the manufacturer's instructions, detection range of HA by the kit was 10–800 ng/ml. If a sample contained more than upper limit of the measurable range, it was diluted with reaction buffer of the kit and measured again.

EXPRESSION OF HA SYNTHASE MRNA

Total cellular RNA from 1×10^6 cells with or without stimulation for six hours by IL1 β (1 ng/ml), TNF α (300 units/ml), IFN γ (1,000 units/ml) or TGF β 1 (1 ng/ml), respectively, was isolated by a modified acid guanidinium thiocyanate/phenol/chloroform extraction method, using RNAzol B (Cinna/Biotech Laboratories, Houston, TX). The RNA was reverse transcribed and the cDNA amplified by a polymerase chain reaction (RT-PCR), using a GeneAmp RNA PCR kit (Perkin-Elmer Cetus, Norwalk, CT). The primers of HA synthase (HA synthase 1), 5'-TGG GGC GGC AAG CGC GAG GTC ATG TAC ACA GC-3' (sense) and 5'-CAC CAG AGC GCG TTG TAC AGC CAC TCA CGG AAG TA-3' (antisense) were used.^{10–12} The primers of glyceraldehyde-3-phosphate dehydrogenase (GAPDH), 5'-GTC TTC ACC ACC ATG

GAG AAG GCT-3' (sense) and 5'-CAT GCC AGT GAG CTT CCC GTT CA-3' (anti-sense), were used for detecting the positive control cDNA. Each reaction contained 2 μ l of cDNA, 8 μ l of dNTP mixture, 10 μ l of $10 \times$ Taq DNA polymerase buffer, 8 μ l of 25 mM MgCl₂, 0.5 μ l AmpliTaq DNA polymerase (Perkin-Elmer Cetus), 1 μ l of each primer, and water to 100 μ l. The samples were overlaid with mineral oil (Sigma, St Louis, MO) and cycled through 30 cycles of denaturation (95°C for one minute), annealing (50°C for two minutes), and extension (72°C for three minutes) in a programmable thermal controller (Mini-Cycler, MJ Research, MA). The PCR products were subjected to electrophoresis on 2% agarose gels (Agarose MP, Boehringer Mannheim Biochemica, Mannheim, Germany) and stained with ethidium bromide. The primer set of HA synthase amplifies a 548-bp sequence from cDNA transcribed from HA synthase mRNA.

MEASUREMENT OF HA MOLECULAR WEIGHT

The culture supernatants of the cells cultured for 96 hours with or without stimulation by IL1 β , TNF α , IFN γ or TGF β 1 were collected. The molecular weight of HA in each supernatant was measured at Seikagaku Kogyo Co (Tokyo) using high performance gel-permeation liquid chromatography (HPLC-GPC). Briefly, supernatants were separated on a TSKgel G6000PWXL column equipped with an HLC-8020 system (Toso, Tokyo), with 0.2 M NaCl as the elution buffer. The molecular weight of HA in the samples was determined by the duration times through the column, reflected by those of HA standards. For calibration, purified HA standards (molecular weight 2130 kDa, 460 kDa or 104 kDa) were used.

STATISTICAL ANALYSIS

Data are expressed as mean (SD) values. Differences were analysed by the least significant difference test using a statistical program, StatMate II (Nankodo, Tokyo). Differences were regarded as statistically significant if $p \leq 0.05$.

Results

HA PRODUCTION BY RAMAK-1 CELLS

HA concentrations in the supernatants of RAMAK-1 cells without stimulation were 1210 (385) ng/ml (24 hours) and 1964 (317) ng/ml (48 hours) (fig 1). Although HA synthesis was not significantly augmented by TNF α (966 (152) ng/ml, 24 hours; 1418 (288) ng/ml, 48 hours) or by IFN γ (1260 (136) ng/ml, 24 hours; 2214 (322) ng/ml, 48 hours) it was significantly stimulated by IL1 β (2183 (541) ng/ml, 24 hours, $p < 0.05$; 4488 (328) ng/ml, 48 hours, $p < 0.01$) (fig 1). In contrast, TGF β 1 significantly ($p < 0.01$) inhibited HA production (300 (171) ng/ml, 24 hours; 357 (54) ng/ml, 48 hours) (fig 1).

The time and dose dependent inhibition of HA production by TGF β 1 was studied in another series of experiments. TGF β 1 (at concentrations of 0.1, 1, 5 or 10 ng/ml)

Figure 2 Time and dose dependent inhibition of HA production by RAMAK-1 cells cultured with RPMI-1640 medium containing 10% FBS with or without stimulation by TGFβ1. Cells were incubated for up to 96 hours in the presence of TGFβ1. Supernatants were removed for HA determinations by sandwich binding protein assay at 24, 48, 78 or 96 hours. TGFβ1 concentrations indicated were used.

Figure 3 HA synthase mRNA of RAMAK-1 cells when stimulated for six hours by IL1β, TNFα, IFNγ or TGFβ1 using RT-PCR. Lane 1, without stimulation; lane 2, with stimulation by IL1β; lane 3, with stimulation by TNFα; lane 4, with stimulation by IFNγ; lane 5, with stimulation by TGFβ1. The primer set amplifies a 548-bp sequence from cDNA transcribed from HA synthase mRNA.

Table 1 Molecular weight of HA secreted by RAMAK-1 cells with or without stimulation by IL1β, TNFα, IFNγ or TGFβ1

	Mean (SD) molecular weight (kDa)
Control	770 (160)
IL1β (1 ng/ml)	1270 (270)*
TNFα (300 units/ml)	470 (90)
IFNγ (1000 units/ml)	1220 (280)*
TGFβ1 (1 ng/ml)	170 (20)**

*p<0.05, **p<0.01, n=3.

significantly ($p<0.05$) inhibited the production of HA in this cell line compared with that of cells without stimulation (fig 2).

EXPRESSION OF HA SYNTHASE MRNA
RAMAK-1 cells expressed HA synthase mRNA in the absence of stimulation or when stimulated by IL1β, TNFα or IFNγ. In contrast, HA synthase mRNA was not detected in TGFβ1 stimulated cells (fig 3).

MOLECULAR WEIGHT OF HA BY RAMAK-1 CELLS
The molecular weight of HA in the culture supernatants of RAMAK-1 cells in the absence of stimulation was 770 (160) kDa, whereas it was significantly ($p<0.05$) increased to 1270 (270) kDa or to 1220 (280) kDa with stimulation by IL1β (1 ng/ml) or IFNγ (1000 units/ml), respectively. In contrast, it was significantly ($p<0.01$) reduced to 170 (20) kDa with stimulation by TGFβ1 (1 ng/ml) (table 1).

Discussion
HA, isolated initially from the vitreous body of the eye, is an ubiquitous component of the extracellular matrix.¹³ It has been used for the clinical benefit of patients undergoing ocular operations¹⁴ or with arthritic joint diseases.¹⁵⁻¹⁸ In joints, HA at the synovial lining layer or in synovial fluid retains water though the synovial vessels by its high water binding capacity and

plays rheological or viscoelastic parts.¹⁵⁻¹⁹ HA concentrations are lower in rheumatoid synovium than in non-inflamed synovium.²⁰ HA synthesis by RAMAK-1 cell line is up regulated when stimulated by IL1β but is inhibited by TGFβ1 stimulation (figs 1, 2). Inhibition of HA production by TGFβ1 was also confirmed by the disappearance of its synthase mRNA—that is, HA production was inhibited by TGFβ1 at the transcriptional level (fig 3).

With respect to the concentrations of TGFβ1, TGFβ1 at concentrations of 1 ng/ml and higher, served to increase prolyl 4-hydroxylase production by dermal fibroblasts.²¹ For the stimulation of rheumatoid synovial cells, 1–5 ng/ml of TGFβ1 were often used as optimal dose.²²⁻²³ In this study, this cytokine at concentrations of 1–10 ng/ml remarkably inhibited HA production by RAMAK-1 cells.

Haubeck *et al*²² reported that IL1β and TGFβ1 were major stimulators of HA synthesis by synovial lining cells while TNFα and IFNγ were not. The findings in this study with respect to TGFβ1 are in direct contrast to those of Haubeck *et al*.²² A difference in the degree of cell differentiation or that in the behaviour of “normal” and “rheumatoid” synovial cells might be behind the contrasting observations. Haubeck *et al*²² used cells derived from normal synovial tissue, while we only used a cell line derived from synovial tissue of a patient with rheumatoid arthritis. Furthermore, they used Opti-MEM1 medium without the addition of serum to culture their synovial lining cells. Opti-MEM1 was developed to reduce the amount of added serum by more than 50%, but it still requires a serum supplement. Finally, in their paper it is not clear whether or not cell viability was sustained for 60 hours without the addition of serum. In our culture system, a 10% addition of FBS maintained good viability of the cells. In fibroblast cultures, the addition of 10% FBS in culture medium augments basal HA synthesis.²⁴ Differences in culture conditions might also have caused the discrepancy between our data with respect to TGFβ1 and those of Haubeck *et al*.²²

With respect to culture condition, we used a 10% addition of FBS throughout the study. Other culture conditions, such as no serum or reduced serum concentrations, were not examined. If cells were cultured under such conditions, basal production of HA might have decreased, so that the inhibitory effect of TGFβ1 might have been less observed.

TGFβ1 is a potent inducer/modulator of a wide range of biological processes in many different systems, such as inhibition of growth, differentiation of several kinds of cells,²⁵⁻²⁶ and suppression of inflammation.²⁷ This cytokine is also referred to as a bifunctional regulator of cell function to emphasise its qualitatively heterogeneous effects. Intra-articular injection of TGFβ1 into normal rabbit joints induces synovial inflammation and hyperplasia with a predominantly mononuclear phagocyte infiltrate that contributes locally to joint inflammation.²⁸⁻²⁹ In contrast, when administered systemically,

TGF β 1 significantly ameliorates collagen type II induced arthritis or inhibits the development of streptococcal cell wall induced polyarthritis that is immunosuppressive and consistently shows therapeutically desirable effects.^{30, 31} According to the results of this study, TGF β 1 might reduce the oedema of the soft connective tissues at the joints by inhibiting HA production. Inhibition of HA synthesis by TGF β 1 might improve the swelling of the inflamed joints, while the anti-inflammatory or analgesic effects³² of HA might lessen as HA production is reduced. The molecular weight of HA synthesised by RAMAK-1 cells with stimulation by IL1 β or IFN γ is strikingly augmented compared with that in the absence of stimulation (table 1). Interestingly, IFN γ did not augment HA production in the light of total amount but significantly affected the molecular weight of HA. Although the precise mechanism for formation of high molecular weight of HA with stimulation by IL1 β or IFN γ was not elucidated, it is suggested that HA with a higher molecular weight produced by synovial lining cells in vivo could play a part as an anti-inflammatory and an analgesic effector³² by an interaction with HA receptor(s).³³

Results of this study show that RAMAK-1 cells readily produce HA and that the production of HA is stimulated by IL1 β and inhibited by TGF β 1. Therefore, overall HA production in the synovial lining cells of RA inflamed joints is likely to be regulated by a balance of these cytokines.

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