Relaxin inhibits effective collagen deposition by cultured hepatic stellate cells and decreases rat liver fibrosis in vivo

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Abstract

Background—Following liver injury, hepatic stellate cells (HSC) transform into myofibroblast-like cells (activation) and are the major source of type I collagen and the potent collagenase inhibitors tissue inhibitors of metalloproteinases 1 and 2 (TIMP-1 and TIMP-2) in the fibrotic liver. The reproductive hormone relaxin has been reported to reduce collagen and TIMP-1 expression by dermal and lung fibroblasts and thus has potential antifibrotic activity in liver fibrosis.

Aims—To determine the effects of relaxin on activated HSC.

Methods—Following isolation, HSC were activated by culture on plastic and exposed to relaxin (1–100 ng/ml). Collagen deposition was determined by Sirius red dye binding and radiolabelled proline incorporation. Matrix metalloproteinase (MMP) and TIMP expression were assessed by zymography and northern analysis. Transforming growth factor β1 (TGF-β1) mRNA and protein levels were quantified by northern analysis and ELISA, respectively.

Results—Expression of activated HSC to relaxin resulted in a concentration dependent decrease in both collagen synthesis and deposition. There was a parallel decrease in TIMP-1 and TIMP-2 secretion into the HSC conditioned media but no change in gelatinase expression was observed. Northern analysis demonstrated that primary HSC, continuously exposed to relaxin, had decreased TIMP-1 mRNA expression but unaltered type I collagen, collagenase (MMP-13), alpha smooth muscle actin, and TGF-β1 mRNA expression.

Conclusion—These data demonstrate that relaxin modulates effective collagen deposition by HSC, at least in part, due to changes in the pattern of matrix degradation.

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Keywords: relaxin; hepatic stellate cell; hepatic fibrosis; type I collagen

Liver fibrosis is characterised by increased hepatic deposition of extracellular matrix proteins, particularly type I collagen. An overwhelming body of evidence indicates that the hepatic stellate cell (HSC, lipocyte, fat storing, or Ito cell) is the pivotal effector cell in hepatic fibrogenesis.1–7 Following liver injury HSC become activated, proliferate, and undergo transformation to a myofibroblast-like phenotype. In addition to expressing matrix proteins, particularly type I collagen,5–7,9 activated HSC also express matrix degrading metalloproteinases and the potent metalloproteinase inhibitors tissue inhibitor of metalloproteinases 1 and 2 (TIMP-1 and TIMP-2).5–9 We and others have postulated that progression of fibrosis results from increased synthesis of extracellular matrix molecules coupled with elevated expression of TIMP-1 and TIMP-2. Antifibrotic strategies might therefore usefully be targeted towards either reducing matrix synthesis or increasing matrix degradation.

Relaxin, an insulin-like growth factor hormone, is detectable in serum during pregnancy and plays an important role in initiating the structural remodelling of the cervix and interpubic ligament in preparation for parturition.10–12 Studies suggest the effects of relaxin are mediated by both a decrease in collagen synthesis and an increase in the production of collagenase and proteoglycanase.13–14 These observations have led to an interest in the potential role of relaxin as an antifibrotic agent. Human skin fibroblasts stimulated to overproduce collagens by inflammatory cytokines respond to relaxin by decreasing their production of type I and III collagens and increasing their synthesis and secretion of pro-collagenase, and by decreasing TIMP-1 expression.15 Similarly, scleroderma fibroblast lines decrease their excessive expression of type I collagen in response to treatment with relaxin.16 These results suggest that in vitro, relaxin may modulate matrix turnover in favour of degradation in situations of collagen overexpression.

In vivo, recombinant human relaxin reduced collagen accumulation in a rodent model of dermal fibrosis and was associated with an alteration in the organisation of collagen fibrils within the interstitium.17 Subsequently, relaxin was demonstrated to significantly reduce collagen accumulation in a bleomycin induced murine model of lung fibrosis.18 Evidence from a model of renal fibrosis suggests that the antifibrotic effect of relaxin may in part be

Abbreviations used in this paper: HSC, hepatic stellate cells; TIMP, tissue inhibitor of metalloproteinases; MMP, matrix metalloproteinase; α-SMA, alpha smooth muscle actin; TGF-β1, transforming growth factor β1; DMEM, Dulbecco’s minimal essential medium; PBS, phosphate buffered saline; BSA, bovine serum albumin; TCA, trichloroacetic acid; CCl₄, carbon tetrachloride.
mediated by downregulation of transforming growth factor β1 (TGF-β1). Moreover, in some systems, TGF-β1 has been reported to, in turn, inhibit relaxin release.

In the studies reported here, we have examined the effect of recombinant human relaxin on expression of type I collagen, TIMPs, and matrix metalloproteinases (MMPs) in culture activated HSC. Our results indicate that relaxin reduces effective collagen deposition which may result directly from downregulation of type I collagen synthesis and indirectly due to a decrease in TIMP-1 synthesis by HSC, effects that are probably not mediated by endogenous TGF-β1 expression.

Methods

HSC ISOLATION AND CULTURE

HSC were isolated from normal male Sprague-Dawley rats and cultured as previously described. Cultures were exposed to recombinant human relaxin 1–100 ng/ml (Lot 63601; Connetics Corp., Palo Alto, California, USA) either continuously from isolation or from days 5–7 (activated HSC) for varying periods of time as stated. To provide confluent monolayers of activated cells for collagen synthesis assays, activated rat and human HSC (>7 days in primary culture) were passaged into multiwell plates.

QUANTIFICATION OF COLLAGEN DEPOSITION BY CULTURED HSC

Primary HSC (activated by culture on plastic in the presence of serum) were exposed to relaxin 1–100 ng/ml for 48 hours. Cells were washed and collagen deposited in the wells was stained with Picro Sirius red dye, as described by the manufacturer (Biocolor, Belfast, UK). Unbound dye was removed by washing and the bound complex dissolved in 0.5% sodium hydroxide. Collagen was quantitated by spectrophotometry at 540 nm and results were expressed as a percentage of the untreated controls.

QUANTIFICATION OF COLLAGEN PRODUCTION BY CULTURED HSC

Collagen synthesis by HSC was determined by measuring incorporation of [H] proline into collagenous proteins using modifications of the methods of Postlewaite and colleagues and Cairns and Walls. Early passaged HSC were seeded into 12 well plates and grown until approaching confluence. Cells were made serum free by washing and incubated in fresh Dulbecco’s minimal essential medium (DMEM) supplemented with antibiotics, 25 µg/ml ascorbic acid, 0.01% bovine serum albumin (BSA), and recombinant relaxin 0–100 ng/ml in a total volume of 500 µl/well. After 24 hours the media was replaced with fresh media containing the same additions as above plus 1 µCi L-[2,3,4,5-3H] proline (Amersham, Little Chalfont, UK) per well. After a further 24 hours the cell supernatants were harvested and the cells lysed in TE (10 mM Tris HCl, 1 mM EDTA, pH 7.5) for DNA quantification.

The supernatants were divided into 4×100 µl aliquots and loaded onto 96 well multiscreen filtration plates (Millipore UK Ltd, Watford, UK) and assayed in duplicate either to measure total protein production or production of proteins resistant to digestion by a highly purified bacterial collagenase (non-collagenous proteins). Collagen production per well was calculated using the following formula:

[H proline incorporation (total protein)]−[H proline incorporation (non-collagenous protein)]/DNA content (µg/well)

A total of 35 µl of 0.2 M Tris/0.3 M calcium chloride pH 7.5 and 15 µl of 50 mM N-ethylmaleimide were added to each well while in addition 50 U/well of a highly purified bacterial collagenase (Worthington Biochem Corporation, New Jersey, USA) was added to the wells assigned to determine non-collagenous protein production. The plates were incubated at 37°C for 90 minutes. Proteins were then precipitated using 50% trichloroacetic acid (TCA) and incubation on ice for one hour. The plates were washed with 10% TCA and the incorporated radioactivity per well determined by scintillation scanning.

DNA QUANTIFICATION

HSC that had been cultured in the presence or absence of varying concentrations of relaxin were lysed in phosphate buffered saline (PBS) or TE, sonicated, and the DNA content determined by addition of bisbenzamidazole (Hoechst 33258) (Millipore) or PicoGreen (Molecular Probes, Leiden, the Netherlands) and then quantified by fluorometry.

COLLECTION OF CONDITIONED MEDIA AND EXTRACTION OF RNA FROM HSC

HSC cultured for varying times in serum containing media were washed three times in serum free DMEM and incubated in 2.5 ml of fresh serum free media containing antibiotics and 0.01% BSA with relaxin 1–100 ng/ml for varying periods of time as described previously. The media were collected and clarified by centrifugation. Cell monolayers were lysed in 4 M guanidinium isothiocyanate after which RNA was isolated by the acid/phenol method as described previously.

ZYMOGRAPHY/REVERSE ZYMOGRAPHY

Metalloproteinases and TIMPs were detected by gelatin substrate zymography and reverse zymography, respectively, as previously described.

QUANTITATION OF COLLAGENASE

Collagenase activity was measured in conditioned media from primary or early passage HSC that had been cultured continuously with relaxin 100 ng/ml for 1–3 days, as previously described.

NORTHERN ANALYSIS

Aliquots of 10 µg of total RNA were electrophoresed and transferred to nylon membranes. Probes for TIMP-1, MMP-13, type I collagen, alpha smooth muscle actin (α-SMA), TGF-β1, and β-actin were transcribed from the relevant cDNA using the Megaprime DNA labelling kit (Amersham), and [α-32P] ATP (Amersham).
After overnight hybridisation the membranes were washed at 55°C in 0.2% sodium dodecyl sulphate/0.15 M sodium chloride, 0.015 M sodium citrate, pH 7, and exposed to x-ray film at −70°C or to a Storm phosphor imager screen (Molecular Dynamics, Sunnyvale, California, USA).

**Results**

**RELAXIN REDUCES COLLAGEN SYNTHESIS AND DEPOSITION BY CULTURE ACTIVATED HSC**

The in vitro effect of relaxin on HSC collagen secretion was quantified by Sirius red dye binding and spectrophotometry. Incubation of HSC for 48 hours with relaxin induced a modest but consistent dose dependent reduction in the amount of collagen deposited on tissue culture plates (fig 1A) (0.05>p>0.01, n=7, for relaxin 10 and 100 ng/ml compared with control by paired t test). The effect of relaxin on collagen synthesis by activated HSC was examined by radiolabelled proline incorporation into collagens present in the tissue culture supernatant. Treatment with relaxin (100 ng/ml) resulted in a 40% decrease in collagen synthesis over a 24 hour period (fig 1B) (0.05>p>0.01, n=5, for relaxin 1, 10, and 100 ng/ml compared with control by paired t test).

A confirmatory experiment assessing collagen synthesis by 3H proline incorporation was performed on passaged human HSC in which culture with relaxin (100 ng/ml) for 24 hours led to a 71% reduction in collagen production. Secreted TIMP-1 and TIMP-2 are regulated by exposure of HSC to relaxin.

**SECRETED TIMP-1 AND TIMP-2 ARE REGULATED BY EXPOSURE OF HSC TO RELAXIN**

Zymographic techniques were used to examine MMP and TIMP secretion by HSC (fig 2). After 24 hours of relaxin exposure, a concentration dependent reduction in TIMP-1 and TIMP-2 was observed. Gelatinase A and B were constitutively secreted by activated HSC (fig 2). However, relaxin did not induce a change in the secretion of either thezymogen or active forms of gelatinase A or B. No activity attributable to secreted rat interstitial collagenase MMP-13 was observed by zymography in any of the experiments performed. Moreover, no collagenase activity was detected in 14C

![Figure 1](http://www.gutjnl.com)
RELAXIN DOES NOT MEDIATE EXPRESSION OR ACTIVATION OF TGF–β1 IN HSC
As the decrease in collagen deposition and downregulation of TIMP–1 observed in relaxin treated culture activated HSC are consistent with TGF–β1 downregulation, expression and activation of TGF–β1 were determined. Northern analysis of RNA from activated primary or early passage HSC that had been continuously cultured with relaxin 100 ng/ml for between one and five days demonstrated no alteration in

TGF–β1 mRNA expression relative to control cultures (fig 4). The amount of secreted TGF–β1 present in conditioned media from cultured HSC was quantified by ELISA. Figure 5 shows representative results of one of three independent experiments using primary HSC, demonstrating that there was no change in the levels of active (fig 5A) or latent (total–active) (fig 5B) TGF–β1 in response to culture with relaxin for 24–72 hours. Similarly, no change in latent or active TGF–β1 secretion into conditioned media was seen in early passage HSC exposed to relaxin compared with control cultures (n=2). A single further experiment was performed on passaged human HSC, and again no differences in the levels of active or total TGF–β1 were seen in the conditioned media of human HSC exposed to relaxin 100 ng/ml for 24 hours.

RELAXIN REDUCES COLLAGEN DEPOSITION IN AN IN VIVO MODEL OF HEPATIC FIBROSIS
The in vivo effect of relaxin on the development of hepatic fibrosis was examined using the rat CCl4 model and continuous subcutaneous infusion of recombinant relaxin. The effective delivery of relaxin was confirmed by measurement of serum relaxin in the treatment group at the end of the experimental period. The mean serum relaxin level was 18.8 ng/ml
48

and 10 µg aliquots were subjected to northern blotting and probed for TGF-

hours). Total RNA was extracted from day 5 primary HSC (A) or passaged P1 HSC (B) hepatic stellate cells (HSC) cultured in the presence or absence of relaxin (100 ng/ml for 24

Figure 4 Northern blot analysis for transforming growth factor

Figure 5 Transforming growth factor β1 (TGF-β1) levels in the conditioned media of cultured HSC. Primary (day 5) hepatic stellate cells cultured in serum free conditions were incubated in the presence (+) or absence (−) of relaxin 100 ng/ml for 24–72 hours and the total amount of active and latent TGF-β1 determined as described. Levels of active TGF-β1 are represented in (A) and latent (total−active) TGF-β1 in (B). Results shown are representative of one of three independent experiments.

(range 0.7–41.7). Relaxin is undetectable in the serum of untreated male rats. After 28 days of CCl4 intoxication, relaxin treatment resulted in a significant decrease in liver weight relative to control livers (expressed as mean (SEM), relaxin treated 17.8 (0.57) g vs controls 20.1 (0.56) g; p<0.05 by Mann-Whitney; n=6 for each group). Hepatic hydroxyproline levels were reduced in the relaxin treated group relative to controls (relaxin treated 1.29 (0.06) µmol/g wet weight v control 1.85 (0.12) µmol/g wet weight). In a further assay, the hydroxyproline content of normal rat liver was determined as 0.95 relaxin µmol/g wet weight (n=3). The total liver hydroxyproline content of the relaxin treated livers was significantly decreased relative to controls (relaxin treated 22.9 (1.7) µM; p<0.01 by Mann-Whitney test; n=6 for each group)

Discussion

We have presented data demonstrating that recombinant human relaxin reduces the effective deposition of interstitial collagen in both tissue culture and in an in vivo model of hepatic fibrosis. In vitro, we examined the effects of relaxin on HSC, as the HSC has been clearly identified as having a major role in matrix homeostasis in hepatic fibrogenesis. HSC cultured on plastic assume an activated phenotype, characterised by elevated type I collagen, TIMP-1, and TGF-β1 expression. Our results demonstrate that culture activated HSC treated with relaxin show a consistent dose dependent decrease in the deposition of collagen compared with control cultures. We endeavoured to determine the mechanisms underlying these changes and so examined the protein biosynthetic levels of type I collagen, MMP-13, TIMP-1, TIMP-2, and TGF-β1.

We have also described the effect of relaxin on levels of transcripts of type I collagen, MMP-13, TIMP-1, and α-SMA. Overall, our results suggest that in this model relaxin mediates a reduction in collagen deposition directly by reducing type I collagen protein synthesis, and indirectly by decreasing TIMP-1 and TIMP-2 expression, thereby potentially enhancing matrix degradation. Moreover, our data indicate that this effect is not mediated via downregulation of TGF-β1 expression. Finally, relaxin was tested in a well established and reproducible animal model of hepatic fibrosis to examine its effect on the development of liver fibrosis in vivo. Consistent with our cell culture data, our results in this pilot study demonstrated that treatment with relaxin led to a decrease in liver size in response to fibrotic injury associated with a significant decrease in total liver hydroxyproline content.

In vitro, culture of primary HSC on plastic is an established model which mimics the phenotypic changes that occur during the process of HSC activation following liver injury. Collagen deposition in cell monolayers was assessed using the Sirius red dye binding assay which demonstrated a statistically significant decrease in the amount of collagen laid down by primary HSC cultured in the presence of relaxin. To determine whether the deposition was related to a decrease in collagen synthesis, we examined collagen synthesis using radiolabelled proline incorporation over a limited period of time in HSC cultured with and without relaxin. Both assays gave a clear reproducible and statistically significant effect on collagen synthesis and secretion, demonstrating that relaxin decreases both the rate of collagen synthesis and total collagen deposition by culture activated HSC.

Northern analysis was used to assess type I collagen gene expression in culture activated primary HSC, to determine whether relaxin regulated type I collagen at the level of mRNA. No consistent change was found as a consequence of culture with relaxin, a result that differs from observations in human skin fibroblasts.18 Although type I collagen mRNA levels may be regulated by protein mediated
changes in stability, continuous relaxin exposure for up to seven days from the time of HSC isolation failed to elicit any changes in mRNA. These results suggest that relaxin regulates collagen deposition at the transcriptional or post-translational level in these cells. Moreover, effective deposition may also depend on changes in the rate of matrix degradation.

We went on to examine expression of metalloproteinases and TIMPs by HSC as they may directly regulate collagen levels via matrix degradation and turnover. In keeping with the hypothesis that relaxin induces a matrix degrading phenotype in HSC, a modest but consistent decrease in TIMP-1 mRNA expression and a clear dose dependent reduction in secreted TIMP-1 and TIMP-2 activity, as determined by reverse zymography, was demonstrated following relaxin exposure. This result is significant in light of the potential importance of the metalloproteinase inhibitors, particularly TIMP-1, in the progression of liver fibrosis in vivo and in vitro. A direct effect of relaxin on fibroblast expression of TIMP is already documented. However, we detected no relaxin mediated enhancement of collagenase (MMP-13) expression. These findings do not exclude critical matrix degradation occurring in a focused and local manner immediately adjacent to HSC. Urokinase plasminogen activator activity and thus plasminogen activation has been localised to the cell surface in HSC. Gelatinase A and B, which are constitutively secreted by HSC in the presence of relaxin, have been demonstrated to have weak collagenolytic activity but were not regulated by relaxin.

We explored the hypothesis that the observed antifibrotic effects of relaxin may be mediated via downregulation of endogenous TGF-β1. The relaxin and TGF-β1 systems can be viewed as being mutually antagonistic in their effects. Moreover, TGF-β1 is expressed by HSC in an autocrine manner and upregulates TIMP-1 expression. Relative to control, TGF-β1 mRNA expression was unchanged in primary and early passage HSC exposed to relaxin and also there were no differences in secretion of latent TGF-β1 in response to relaxin. This suggests that the effect of relaxin in HSC is not mediated via TGF-β1.

Finally, relaxin was tested for its effects in CCl4 induced liver fibrosis in rats. Relaxin treatment induced a significant reduction in both liver weight and overall hydroxyproline content relative to controls. These data indicate that the observed effects of relaxin on collagen and TIMP expression in tissue culture are reflected in an in vivo model of hepatic fibrosis. Serum levels of relaxin achieved were greatly in excess of physiologic levels in the human which peak at 900 pg/ml in the first trimester of pregnancy. The decrease in liver hydroxyproline content in this model is comparable with that seen in rodent models of pulmonary fibrosis treated with relaxin but less dramatic than that observed with rodent dermal fibrosis. Taken together, the data suggest that recombinant human relaxin modulates connective tissue degradation in rodent models of fibrosis although by cross-species the effect may be attenuated. Nevertheless, these studies represent proof of concept for the need to further evaluate the effectiveness of relaxin in fibrotic models. More detailed studies of the role of relaxin in modulating a pre-existing fibrotic injury are proposed.

In summary, our results are consistent with the previously reported ability of relaxin to inhibit collagen deposition and promote collagen degradation in circumstances where collagen is overexpressed. The magnitude of change that we report is similar to that observed in other rodent studies. Our studies represent a proof of concept that relaxin is worthy of further study as an agent capable of modifying fibrosis in the liver.

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