Cultivating hepatocytes on printed arrays of HGF and BMP7 to characterize protective effects of these growth factors during in vitro alcohol injury

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Abstract

The goal of the present study was to investigate hepato-protective effects of growth factor (GF) arrays during alcohol injury. Hepatocyte growth factor (HGF) and bone morphogenetic protein (BMP)7 were mixed with collagen (I) and robotically printed onto standard glass slides to create arrays of 500 μm diameter spots. Primary rat hepatocytes were seeded on top of the arrays forming clusters corresponding in size to the underlying protein spots. Cell arrays were then injured in culture by exposure to 100 mM ethanol for 48 h. Hepatocytes residing on GF spots were found to have less apoptosis then cells cultured on collagen-only spots. Least apoptosis (0.3% as estimated by TUNEL assay) was observed on HGF/BMP7/ collagen spots whereas most apoptosis (17.3%) was seen on collagen-only arrays. Interestingly, the extent of alcohol-induced apoptosis in hepatocytes varied based on the concentration of printed GF. In addition to preventing apoptosis, printed GFs contributed to maintenance of epithelial phenotype during alcohol injury as evidenced by higher levels of E-cadherin expression in HGF-protected hepatocytes. Importantly, GF microarrays could be used to investigate heterotypic interactions in the context of liver injury. To highlight this, stellate cells – nonparenchymal liver cells involved in fibrosis e – were added to hepatocytes residing on arrays of either HGF/collagen or collagen-only spots. Exposure of these cocultures to ethanol followed by RT-PCR analysis revealed that stellate cells residing alongside HGF-protected hepatocytes were significantly less activated (less fibrotic) compared to controls. Overall, our results demonstrate that GF microarray format can be used to screen anti-fibrotic and anti-apoptotic effects of growth factors as well as to investigate how signals delivered to a specific cell type modulate heterotypic cellular interactions.

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1. Introduction

Alcohol abuse is a prevalent etiological factor of fibrosis and cirrhosis of the liver. Ethanol injury leads to apoptosis of hepatocytes and to an increase in production of fibrogenic cytokines such as transforming growth factor (TGF)-β1 [1]. The pathogenesis of alcohol injury is complex, involving several signal transduction pathways and several liver cell types. Stellate cells – nonparenchymal cells in the liver – become activated during liver injury by alcohol and initiate programs of TGF-β production and extracellular matrix (ECM) remodeling that contribute to apoptosis of hepatocytes [1–3]. TGF-β has also been shown to induce dedifferentiation and epithelial–mesenchymal transition (EMT) in hepatocytes [4,5].

Identifying TGF-β antagonists is an important strategy in preventing fibrosis of the liver as well as other tissues. Hepatocyte growth factor (HGF) is a signaling molecule central to liver development and regeneration [6] and that has been reported to antagonize TGF-β and prevent fibrosis [7–9]. BMP7 is another important growth factor that has been shown to counteract pro-fibrogenic effects of TGF-β1 [10,11].

Several studies have reported that systemic administration of HGF in laboratory animals provided protection against liver injury [12,13]. However, these studies were only able to compare two or
three concentrations of HGF because of the cost of animals and reagents. Determining optimal concentration is critical for both HGF, which has been implicated in cancer metastasis [14,15] and BMP7, which has been shown to induce apoptosis at higher concentrations [16]. While it is beneficial to develop in vitro assays to determine liver protective concentrations and combinations of GFs, traditional cell culture approaches of adding soluble GFs to culture media require large amounts of expensive growth factors and are suboptimal for high-throughput screening.

Increasingly, the attention has turned to surface immobilization and solid-phase presentation of GFs to cells [17–20]. A number of studies reported covalent immobilization of GFs on cell culture surfaces [18,21,22]. However, in vivo, GF molecules interact with components of extracellular matrix (ECM) via secondary bonds and are dynamically released upon cell-initiated proteolysis. Moreover, association with ECM components has been shown to stabilize and enhance stimulatory effects of GFs [17,23]. Recently, we described the use of HGF/ECM protein arrays for cultivating primary rat hepatocytes [24] where HGF was mixed in solution with ECM proteins without chemical modification of GF molecules and was printed in an array format onto glass slides. Primary rat hepatocytes cultured on HGF/ECM protein spots were found to maintain differentiated hepatic phenotype for ten days at levels comparable to or better than hepatocytes receiving daily doses of soluble HGF in the media [24]. Experiments with printed HGF/ECM arrays required 200 times less of the expensive growth factor.

The present study builds on the concept of GF arrays to explore how hepatocytes residing on HGF and BMP7 arrays are impacted by exposure to ethanol—a model of acute alcohol injury. While protective effects of soluble HGF and BMP7 in the context of in vitro and in vivo liver injury have been reported [8,25,26], these effects have not been studied for solid-phase presented GFs. The microarray format has shown utility for a range of cell–surface interaction studies [27–32] and this paper highlights a new application of arrays for screening protective effects of GFs or other signaling molecules during cell injury. A combination of localized GF signaling and micropatterned cocultures described in this study may also be used to investigate heterotypic paracrine signaling underlying fibrosis.

2. Materials and methods

2.1. Chemicals and materials

Glass slides (75 × 25 mm²) were obtained from VWR (West Chester, PA), (3-Acryloxypropyl) trimethylsilanetriol was purchased from Gelest, Inc. (Morrisville, PA). Sulfuric acid, hydrogen peroxide, ethanol, collagenase, collagen from rat tail (type I), streptavidin-conjugated Alexa 546, AlexaFlour 488 anti-mouse IgG, heat shock responsive growth factor (HGF), bone morphogenetic protein 7 (BMP7), and transforming growth factor–β1 (TGF–β1) were obtained from Sigma–Aldrich (St. Louis, MO). Mouse anti-E-cadherin antibody was purchased from BD Biosciences. Concentrated phosphate-buffered saline (10X PBS) was purchased from Lonza (Walkersville, MD). Minimal essential medium (MEM), sodium pyruvate, nonessential amino acids, fetal bovine serum (FBS), Superscript III, RNaseOut (RNase inhibitor), dNTPs and bio-tinylated anti-HGF antibodies were purchased from Invitrogen (Carlsbad, CA). 384/24-well plates, 10 mm diameter spots, 384-well plate builder and 24-well plate builder were purchased from Corning (Corning, NY). 384-well plates were purchased from Lab-Path (Temecula, CA). DAPI stain mounting media was purchased from Vectorshield (Burlingame, CA).

2.2. Animals

Adult female Lewis rats weighing 125–200 g were purchased from Charles River Laboratories (Boston, MA) and fed with a commercial diet and water. All animal experiments were performed according to the National Institutes of Health (NIH) guidelines for the ethical care and use of laboratory animals and the experimental protocol was approved by the Institutional Animal Care and Use Committee (IACUC) of the University of California, Davis.

2.3. Printing collagen and GF arrays

Glass slides were modified with acryloxypropyl trimethylsilanetriol using protocols described previously [33]. Collagen was dissolved in 1X PBS – 0.005% Tween-20 at 0.2 mg/ml concentration. GFs were mixed with the ECM solution to the desired final concentration of 50, 125, 250 or 500 ng/ml and allowed to bind to the ECM protein for 30 min at room temperature prior to printing. Protein microarrays were contact-printed under ambient conditions on silane-modified 75 × 25 mm² glass slides using a hand-held MicroCaster. The pins collected protein from a 382-well plate, dispensing 20–70 nL of solution onto the glass slide and forming circular spots ~500 μm in diameter. Pins were cleaned in acetone via sonication for 5 min and subsequently washed with pin-cleaning solution (DI water and isopropanol) between each change in growth factor. Pins were dried with nitrogen before protein printing. Protein arrays were kept in a refrigerator prior to cell cultivation.

2.4. Characterization of HGF retention on collagen arrays

Immunofluorescent staining was used to determine retention of HGF on printed protein arrays. Solution containing 100, 250 and 500 ng/ml HGF and 0.2 mg/ml collagen (I) was printed onto silanized glass slides and the resultant microarrays were incubated for 2 h in cell culture media at 37 °C. Substrates with microarrays were then removed from media and incubated with 1 μg/ml (in 1X PBS) of anti-human HGF-biotin conjugate at 37 °C for 2 h followed by incubation in 10 μg/ml of streptavidin-Alexa 546 for 1 h at 37 °C. Samples were washed between each staining step with 1X PBS – 0.05% Tween-20. In order to quantify fluorescent signal emanating from the array, the laser microarray scanner (Agilent G2565BA fluorescence scanner) was employed to scan the glass slides at a spot pixel resolution of 5 μm. The fluorescence intensity of each array element was determined using GenePix Pro 6.0 data analysis software (Molecular Devices, Downingtown, PA).

2.5. Cultivation of primary hepatocytes on GF microarrays

Primary hepatocytes were isolated from adult female Lewis rats using a standard two-step collagenase perfusion procedure [34]. Typically, 100–200 million hepatocytes were obtained with viability >90% as determined by trypan blue exclusion. Primary hepatocytes were maintained in DMEM supplemented with epidermal growth factor (EGF), glucagon, hydrocortisone sodium succinate, recombinant human insulin, 200 units/ml penicillin, 200 μg/ml streptomycin and 10% FBS.

For cell seeding experiments, a glass slide containing printed arrays of ECM + GF was cut into 1 in. × 1 in. squares to fit into a 6-well plate. The design of different GF arrays tested in our studies is shown in Fig. 1. Typically GFs were printed into arrays of 500 μm diameter spots where different rows contained different concentration of GFs. The distance between rows was 250 μm and the center-to-center spacing of the spots was 1 mm. HGF, BMP7 and TGF-β1 arrays were constructed in this manner. Hepatocytes were seeded to form cellular arrays using protocols described earlier [35]. In brief, glass slides containing printed ECM/GF spots were first exposed to 3 ml of hepatocytes suspended in culture medium at a concentration of 1 × 10⁶ cells/ml. After 1 h of incubation at 37 °C, hepatocytes bound on ECM/GF domains, but did not attach on the surrounding silane-modified surface. The samples were washed twice in PBS to remove unbound hepatocytes and fresh media was added to the sample well.

2.6. Alcohol injury of primary hepatocytes

In our experiments, hepatocytes cultured on GF/collagen microarrays were compared with cells cultured on collagen (I) arrays without GF (see Fig. 1 for description of experiments). Cells were allowed to recover after isolation for 1 day and were then exposed to 100 μM ethanol (6 μL/ml). This concentration of ethanol was chosen to mimic acute alcohol liver injury [37]. Hepatocytes were kept in ethanol-containing culture media for 48 h. To analyze alcohol-induced apoptosis we employed TUNEL (TdT-mediated dUTP nick end labeling) assay using ApoTag Red in situ Apoptosis Detection kit (Chemicon, Temecula, CA) according to the manufacturer’s instructions. DAPI was used to stain cell nucleus. TUNEL-positive cells were scored as apoptotic and were evaluated with a confocal microscope.

2.7. RT-PCR analysis of apoptotic and fibrotic markers

For real-time RT-PCR experiments, 6 × 12 arrays of each GF condition were created and cultured in a separate culture well. Cells (~20,000) were collected from microarrays using trypsin for 30 min at 37 °C. Extracted cell contents were stabilized in 100 μL of lysis buffer and stored at −20 °C. Total RNA was extracted from the cell lysates using absolute total mRNA isolation microprep kit (Stratagene) according to the manufacturer’s instructions. cDNA was synthesized using Quantitect Reverse Transcription kit (Qiagen) according to the manufacturer’s instructions using 12 μL of DNase-pretreated total mRNA. Quantitative real-time PCR was performed using SYBR Green PCR Master Mix (Applied Biosystems). Primers for rat caspase-6, TGF-β1 and GAPDH genes were selected from a database http://medgen.ugent.be/rtprimerdb (Table 1). Primer (Sigma Genosys) concentrations were optimized before use. SYBR Green Master Mix (1×) was used with 1 μl of forward and reverse
primes in a total volume of 12 µl that also included 1 µl of cDNA. All PCR reactions were done in duplicate. PCR amplification was performed as follows: 95 °C for 10 min, 40 cycles of 95 °C for 30 s, 60 °C for 30 s and 72 °C for 1 min on Mastercycler Realplex (Eppendorf). The comparative Ct value method, using housekeeping gene (GAPDH) as an internal standard, was employed to determine relative levels of caspase-6 and TGF-β1.

2.8. Immunofluorescent staining for E-cadherin

For immunostaining, cells were fixed in 4% formalin·0.3% Triton-X 100 in PBS for 15 min. The cells were then incubated in blocking solution (1% bovine serum albumin (BSA)·0.3% Triton-X 100 in 1× PBS) for 1 h at room temperature and exposed to 1:100 diluted anti-E-cadherin antibody (in 1%BSA·0.1% Triton-X 100 in PBS) for 2 h at 37 °C. Finally, cells were incubated in 1:200 diluted anti-mouse IgG conjugated with Alexa 488 (in 1%BSA·0.1% Triton-X 100 in PBS) for visualization. Cells were washed between each step with 1× PBS three times for 5 min. Slides were mounted onto cover slips using DAPI stain mounting media. All incubations were performed at room temperature if not specified otherwise. Stained cells were visualized and imaged using a confocal microscope (Zeiss LSM Pascal). Intensity of E-cadherin staining was quantified using ImageJ software.

2.9. Formation of micropatterned hepatocyte–stellate cell cocultures

A human stellate cell line, created by the Zern laboratory [36], was maintained in DMEM supplemented with 10% FBS, 200 units/ml penicillin, and 200 µg/ml streptomycin at 37 °C in a humidified 5% CO₂ atmosphere. Cells were cultured until 90% confluence and then passaged. Construction of the cocultures was enabled by silane treatment that rendered glass regions non-adhesive to hepatocytes and adhesive to stellate cells. Hepatocytes, seeded first, attached exclusively on printed protein spots (≥ 12 array of each protein array type) and were allowed to spread out overnight. The following day stellate cells were resuspended at 0.25 x 10⁶ cells/ml in DMEM-based media described above and were seeded on the surface containing arrays of hepatic islands. After 30 min of incubation stellate cells became adherent to the glass substrate around hepatocyte clusters. Nonadherent cells were removed from the surface and cocultures were subsequently cultured in hepatocyte medium described in the preceding section.

Table 1

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<th>Gene</th>
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<th>Conc./µM</th>
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<td>R: GACCTGGTGGTGCACCGG</td>
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<td>Rat TGF-β1</td>
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<td>1</td>
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<tr>
<td></td>
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2.10. Alcohol injury of micropatterned cocultures and analysis of stellate cell activation

The goal of these experiments was to compare how GF signaling to the hepatocytes affected activation of neighboring stellate cells in the context of alcohol injury. Ethanol (100 ms) was added for 48 h into culture medium bathing the cocultures. Activation of stellate cells was quantified by real-time RT-PCR analysis of collagen 1, tissue inhibitor of metalloproteinases (TIMP), α-smooth muscle actin (α-SMA1), and TGF-β1 gene expression (see Table 2 for primers). Housekeeping gene used in these studies was β-actin. In these experiments cocultures were trypsinized and nucleic acids were processed according to the protocols described in the preceding section. Importantly, we checked to make sure that primers used for human stellate cells were not cross-reactive with rat hepatocytes.

Micropatterned cocultures were stained for albumin to assess extent of de-differentiation of hepatocytes after alcohol insult. Albumin immunostaining was carried out using the protocol similar to E-cadherin staining described above. Cell morphology and cell motility was observed daily via brightfield microscopy.

3. Results and discussion

This study investigated hepato-protective effects of two growth factors HGF and BMP7 co-printed with collagen (I) in a microarray format. Hepatocytes exposed to ethanol while receiving bottom-up signals from GF arrays were significantly better protected against apoptosis and fibrosis compared to hepatocytes residing on collagen-only spots. Furthermore, protection of hepatocytes on GF arrays prevented activation of adjacent stellate cells in a heterotypic coculture. Our study underscores the potential of microarrays for in vitro screening of therapeutic effects of GFs and for investigating heterotypic cellular interactions underlying fibrosis.

Table 2

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<tr>
<td></td>
<td>R: TCAGACCCATGTGACACAC</td>
<td>1</td>
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<tr>
<td>Human TGF-β1</td>
<td>F: CCTCAGACACAACTATTGC</td>
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<td></td>
<td>R: CTTCCAGCCAGTCTTCTT</td>
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3.1. Printing and characterization of GF microarrays

We have previously described forming cellular monocultures and cocultures on GF/ECM arrays [24]. The arrays were made using a hand-held contact arrayer printing 500 μm diameter protein spots. As shown in Fig. 1, the arrays consisted of 6 by 12 members with center-to-center distance between the spots of 1 mm. In a typical experiment, four different solution concentration of a given GF were printed on a 1 in. × 1 in. glass substrate. Prior to printing of protein arrays, glass substrates were treated with acrylated silane. The silanization procedure enhanced the quality of the printed microarray spots by rendering the surface somewhat hydrophobic (53 ± 2° contact angle). More importantly, the silanization created moderately cell-adhesive glass substrates that did not support attachment of primary hepatocytes and could be used to guide/confine attachment of these cells onto protein arrays. As described later in this paper, the silanized surfaces did permit attachment of more "adhesive" nonparenchymal cells (e.g. stellate cells) and allowed for sequential assembly of micropatterned hepatocyte—stellate cell cocultures on protein microarrays.

![Fig. 2. Characterization of HGF retention and cell patterning on protein microarrays.](image)

Fig. 2. Characterization of HGF retention and cell patterning on protein microarrays. (A) Col/GF arrays printed from different solution concentrations of HGF were incubated for 2 h in cell culture media at 37 °C, immunofluorescently stained and visualized using a microarray scanner. Fluorescent signal due to presence of HGF on protein spots varied as a function of HGF solution concentration. (B) Primary rat hepatocytes adhered onto printed protein spots (500 μm diameter) creating cell arrays. Minimal attachment of hepatocytes was observed on silane-modified glass regions surrounding protein islands.

![Fig. 3. Anti-apoptotic effects of HGF- and BMP7-containing protein arrays.](image)

Fig. 3. Anti-apoptotic effects of HGF- and BMP7-containing protein arrays. (A) Hepatocytes cultured on either Col or Col/GF spots were exposed to 100 mM EtOH. The extent of apoptosis was assessed using TUNEL assay (stains nucleus red), DAPI (blue) was used to stain nuclei of hepatocytes. Representative images show that alcohol exposure caused significant apoptosis in hepatocytes on Col spots (17.3%) and limited apoptosis in hepatocytes cultured on Col/HGF/BMP7 spots (0.3%). (B–C) Quantifying apoptotic hepatocytes by TUNEL assay in the absence (B) and presence (C) of alcohol. Arrays of pro-apoptotic TGF-β were used as positive control. Hepatocytes cultured on HGF or BMP7 were protected from alcohol-induced apoptosis. Anti-apoptotic effects of printed GFs could be titrated in a concentration-dependent manner. Apoptotic nuclei were counted in 6 spots per experiment (≈1000 cells) and the percentage of apoptosis was calculated as apoptotic cells to total cells in spot (#p < 0.05, *p < 0.01; n = 6). Please note, that apoptosis for 500 ng/ml TGF-beta in part (B) and 500/250 ng/ml in part (C) could not be quantified due to cell detachment. For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.
When creating GF microarrays, a solution of collagen was mixed with either HGF and/or BMP7 or TGF-β1 in a microtiter plate and then printed onto a silanized glass slide. To verify presence of HGF after printing, arrays were incubated under cell culture conditions for 2 h, stained with fluorescently-labeled anti-HGF and visualized with a fluorescence microscopy or a microarray scanner. Fig. 2A shows fluorescence images and fluorescence intensity of arrays printed from solution with HGF concentration ranging from 100 to 500 ng/ml. As seen from these data, higher HGF solution concentration translated into higher signal of surface immobilized HGF. In our previous paper we demonstrated that ECM protein microarrays concentration translated into higher signal of surface immobilized HGF. Therefore, GF molecules were expected to be available for cell stimulation during alcohol exposure experiments that typically lasted for 48 h. While immunostaining of printed arrays of BMP7 and TGF-β was not carried, cellular responses to these arrays described in the next section clearly indicate presence of functional GF molecules on the surface.

3.2. Quantification of hepatocyte apoptosis on GF arrays during alcohol injury

Hepatocytes were cultured on HGF- and BMP7-containing collagen (I) spots to characterize protective effects of bottom-up GF signaling during alcohol insult. As shown in Fig. 2B, hepatocytes selectively attached on protein domains, each 500 μm diameter spot containing ~250 cells. The number of cells adherent to the spots was defined by presence of collagen (I) molecules and was consistent on arrays with and without GF molecules. Cells were allowed to spread out overnight before ethanol treatment.

We investigated anti-apoptotic effects of solid-phase presented HGF and BMP7 during alcohol injury of primary rat hepatocytes in vitro. Markers for apoptosis such as internucleosome DNA degradation and caspase-6 activation were analyzed after 48 h incubation of hepatocytes in 100 mM ethanol. This ethanol concentration is physiologically relevant and has been used to mimic acute alcohol injury in vitro [37]. TUNEL staining was used to visualize apoptotic hepatocytes after ethanol exposure. Representative images shown in Fig. 3A highlight that a significant fraction of hepatocytes cultured on collagen spots underwent apoptosis whereas majority of hepatocytes cultured on GF arrays were not apoptotic. TUNEL staining of hepatocytes residing on GF arrays of varying concentrations and combinations (see Fig. 1B for experiment design) was used to quantify apoptosis as a function of the underlying substrate.

Fig. 3B,C quantify hepatic apoptosis after alcohol injury. The purpose of experiments detailed in Fig. 3B was to verify that printed GF arrays did not cause apoptosis in the absence of alcohol. These results compiled in Fig. 3B show that HGF/collagen printed in varying concentrations caused only minimal hepatic apoptosis whereas higher concentration of BMP7 did result in significantly higher number of apoptotic hepatocytes (4.0% vs. 6.4% for 250 ng/ml and 500 ng/ml respectively) compared to collagen-only spots. TGF-β1—an apoptotic and pro-fibrotic factor—was co-printed with collagen (I) to serve as a positive control. As seen from Fig. 3B, TGF-β1 signaling promoted hepatic apoptosis in concentration-dependent fashion with 250 ng/ml solution concentration of this morphogen corresponding to 19.3% apoptosis.

The next set of experiments (Fig. 3C) investigated the relationship between protein array composition and the extent of hepatic apoptosis after exposure to alcohol. These data reveal that hepatocytes cultured on collagen-only spots were susceptible to alcohol-induced apoptosis (17.3%) whereas hepatocytes on GF spots were protected from alcohol injury. Significantly, as shown in Fig. 3C, printed HGF had a dose-dependent anti-apoptotic effect with highest solution concentration of HGF (500 ng/ml) leading to lowest apoptosis (3.1% ± 0.2). Interestingly, the most protective concentration of BMP7 was 250 ng/ml (3.6% ± 1.0) whereas higher concentration (500 ng/ml) resulted in significantly more apoptosis (10.2% ± 1.8). Another significant finding was that the maximal anti-apoptotic effect was achieved with a combination of HGF and BMP7 (0.3% ± 0.5), an over 50-fold reduction in apoptosis compared to hepatocytes on collagen spots (17.3% apoptosis).

Findings described in Fig. 3 clearly demonstrate that hepatic apoptosis observed after alcohol exposure was a strong function of printed GF concentration and could be titrated up or down based on the type and concentration of GF presented on a protein array. Another finding is that with BMP7 arrays, higher concentration was not necessarily better and that an intermediate concentration was found to be most protective against apoptosis. This observation is consistent with reports of pro-apoptotic activity of BMP7 at elevated concentrations [38]. Finally, we demonstrate that lowest levels of alcohol-induced apoptosis in hepatocytes were observed by presenting both HGF and BMP7 on collagen (I) arrays. This points to benefits of combining different GF types on microarrays. All of the findings underscore the need for screening of GF-cell interactions in order to identify optimal concentrations and combinations of GPs.

To corroborate TUNEL apoptosis results, we also carried out RT-PCR analysis of caspase-6—a cysteine-aspartic protease that plays an essential role in apoptosis. As shown in Fig. 4A, ethanol treatment induced a 13.5-fold increase in caspase-6 gene expression in...
hepatocytes residing on collagen spots compared to untreated cells residing on the same surface. The caspase-6 gene expression was reduced by ~6-fold in hepatocytes residing on HGF spots during alcohol treatment. This once again supported the notion that hepatocytes cultured on top of printed GF molecules were protected against apoptosis.

3.3. Maintenance of epithelial phenotype in hepatocytes cultured on HGF arrays during alcohol injury

The production of fibrogenic TGF-β molecules in the liver has historically been assigned to nonparenchymal cells (e.g. activated stellate cells), however, more recently TGF-β has been detected in primary hepatocytes cultured in vitro [4]. These recent findings are part of an emerging paradigm with hepatocytes assuming a more prominent role in initiating/promoting liver fibrosis and contributing to epithelial–mesenchymal transition (EMT) during injury [5]. Given the importance of TGF-β signaling in promoting fibrosis and mesenchymal phenotype, the hepatic expression of this cytokine after alcohol injury was assessed by RT-PCR. As shown in Fig. 4B, TGF-β gene expression of ethanol-treated hepatocytes on collagen spots was 9-fold higher compared to untreated hepatocytes. Hepatocytes residing on HGF/collagen spots during alcohol insult produced significantly less TGF-β transcripts (3.9-fold increase over untreated controls). These data corroborate previous reports of pro-fibrogenic cytokine production by injured hepatocytes [4] and also demonstrate anti-fibrogenic effects of solid-phase presented HGF in our system.

We hypothesized that attenuation of pro-fibrogenic and mesenchymal signaling on HGF arrays should occur concurrently with maintenance or promotion of epithelial phenotype. Expression of E-cadherin, a cell–cell adhesion molecule, is one marker of epithelial phenotype. As shown in Fig. 5, immunofluorescence staining revealed considerably stronger E-cadherin expression in hepatocytes residing on HGF/collagen spots compared to collagen-only spots. Importantly, strong E-cadherin expression was observed in hepatocytes cultured on HGF arrays during exposure to ethanol. Antagonizing TGF-β expression with surface-bound HGF in our experiments delayed or prevented EMT in primary rat hepatocytes.

3.4. Alcohol injury of micropatterned hepatocyte–stellate cell cocultures

Chronic liver injury, such as alcohol consumption, leads to an inflammatory response that includes activation of stellate cells — mesenchymal cells of the liver. These cells are central players in fibrosis and are responsible for secreting TGF-β, depositing collagen I and other ECM proteins in the fibrotic liver [1]. However, liver is a complex organ where multiple cell types residing in close proximity may contribute to fibrosis via paracrine signaling. We were intrigued by our data suggesting TGF-β expression in injured hepatocytes and by the possibility that injured hepatocytes can initiate fibrosis [4,5]. Therefore, we wanted to create a simplified in vitro mimic of the liver where hepatocytes and stellate cells reside in close proximity during alcohol injury. Micropatterned cocultures, described by us and others previously [35,39–41] are particularly suited for these studies because the behavior of one cell type with respect to the other is easier to monitor and characterize. Another significant advantage of the micropatterned coculture format is the possibility of delivering GF signals to a specific cell type during injury [24].

Fig. 5. Immunofluorescent staining for E-cadherin in hepatocytes residing on Col or Col/HGF spots in the absence or presence of alcohol. These images show that E-cadherin expression was significantly stronger in hepatocytes cultured on HGF-containing protein spots and was unaffected by exposure to ethanol.
As mentioned previously in this paper, printing GF/collagen arrays on silanized glass substrates created surfaces with highly cell-adhesive protein islands surrounded by moderately adhesive silane regions. When seeded first, primary rat hepatocytes selectively adhered on protein spots forming 500 µm cell islands. Subsequent seeding of stellate cells resulted in a micropatterned coculture with stellate cells adhering around the hepatic islands.

![Fig. 6.](image)

**Fig. 6.** Alcohol insult of micropatterned hepatocyte–stellate cell cocultures. (A) Left panel demonstrates clusters of hepatocytes formed on Col/HGF spots and surrounded by stellate cells. After alcohol injury hepatocytes expressed high levels of intracellular albumin (green fluorescence) while stellate cells remained on the periphery of hepatocyte islands. Right panel shows that hepatocyte clusters formed on Col spots synthesized less albumin and were invaded by stellate cells. (B) RT-PCR analysis revealed that exposure of cocultures to alcohol resulted in significant activation of stellate cells in the case where neighboring hepatocyte were cultured on Col spots. Conversely, expression of stellate cell activation markers was significantly lower when hepatocytes were cultured on Col/HGF spots during alcohol injury. Markers of stellate cell activation were tissue inhibitor of metalloproteinases (TIMP), α-smooth muscle actin (α-SMA) and transforming growth factor-β1 (TGF-β1). For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.
is surrounded by stellate cells. We compared cocultures where hepatocytes were residing either on HGF/collagen or collagen-only spots during alcohol injury. To assess hepatic phenotype, cocultures were stained for albumin—a hallmark liver product. As seen from Fig. 6A (left panel), hepatocytes receiving bottom-up HGF signaling during alcohol insult maintained cell—cell contacts within the cluster, synthesized a lot of albumin (green fluorescence signal) and kept stellate cells on the periphery. In contrast, hepatocytes residing on collagen-only spots during alcohol injury expressed considerably less albumin and the hepatic clusters were infiltrated by stellate cells (Fig. 6A, right panel). The results of Fig. 6A are in line with other results presented in this paper suggesting that HGF signaling from the bottom-up promotes E-cadherin expression in hepatocytes, and helps maintain differentiated hepatic (epithelial) phenotype.

Invasion of hepatocyte clusters by the stellate cells during alcohol injury (see Fig. 6A, right panel) suggested activation of these cells. We therefore wanted to assess the extent of stellate cell activation in cocultures in the presence or absence of bottom-up HGF signals delivered to the hepatocytes. These experiments were designed to study the role hepatocytes may play in activating neighboring nonparenchymal cells during liver injury. To investigate this, micropatterned cocultures were exposed to 100 μM etanercept for 48 h, trypsinized and analyzed by RT-PCR for expression of genes associated with activated stellate cell phenotype. It should be noted that stellate cells and hepatocytes were from different species (human and rat respectively) which allowed us to design species-specific PCR assays. RT-PCR data presented in Fig. 6B offer compelling evidence that HGF protection delivered to hepatocytes significantly diminished expression of markers associated with activation of proximal stellate cells. Compared to cocultures created on collagen spots, stellate cells cultured next to HGF-protected hepatocytes had 10.4-fold lower level of collagen type I α1 transcripts, 3.1-fold lower tissue inhibitor of metalloproteinases (TIMP) and 11.7-fold lower α-smooth muscle actin1 (α-SMA1) gene expression.

Results presented in Fig. 6 demonstrate that printed HGF contributed to maintenance of hepatic phenotype in micropatterned cocultures undergoing alcohol injury. In addition, HGF signaling to the hepatocytes had dramatic effects in attenuating fibrogenic response of neighboring stellate cells. The latter result, in concert with the findings of GF-induced protection against hepatic apoptosis and fibrosis, points to an important role of injured hepatocytes in triggering/initiating fibrogenic signaling and stellate cell activation.

One should note that stellate cells as well as hepatocytes express HGF receptor c-met and that HGF, while protecting hepatocytes, has been shown to activate stellate cells[42]. Therefore, HGF introduced in soluble form to hepatocytes—stellate cell coculture would have complex and divergent effects on the two cell types. Providing HGF signal from the bottom-up to a specific cell type within the coculture offers a much cleaner way of modulating heterotypic interactions.

**4. Conclusions**

The present paper explored protective effects of GF arrays during in vitro liver injury. Primary rat hepatocytes cultured on HGF- and BMP7-containing collagen arrays during exposure to ethanol were effectively protected against apoptosis. The number of apoptotic hepatocytes decreased with an increasing concentration of printed GFs. Protein spots containing both HGF and BMP7 provided best protection to hepatocytes undergoing alcohol exposure. Additional studies pointed to decrease in TGF-β gene expression and maintenance of epithelial hepatic phenotype in HGF-protected hepatocytes. These experiments strongly support the notion that bottom-up GF signals delivered from the microarrays protect hepatocytes against alcohol-induced apoptosis and fibrosis. In addition, we employed micropatterned cocultures to demonstrate that hepatic apoptosis and fibrosis cause activation of stellate cells, likely through the release of paracrine factors such as TGF-β. Cultivation of hepatocytes on HGF arrays resulted in attenuation of pro-fibrogenic gene expression in neighboring stellate cells. The cell microarray format combined with an in vivo injury model may be used in the future for high-throughput screening of other prospective anti-fibrotic agents. In addition, GF stimulation of specific cells within micropatterned cocultures may help parse out heterotypic signaling in the context of liver injury and may represent an interesting in vitro model of cell type-targeted GF delivery in vivo.

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**Appendix**

Figures with essential colour discrimination. Figs. 1–3 and Figs. 5 and 6 in this article have parts that are difficult to interpret in black and white. The full colour image can be found in the online version, at doi:10.1016/j.biomaterials.2010.04.006.

**References**


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