**Novel Combination BMP7 and HGF Gene Therapy Instigates Selective Myofibroblast Apoptosis and Reduces Corneal Haze In Vivo**

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**PURPOSE.** We tested the potential of bone morphogenetic protein 7 (BMP7) and hepatocyte growth factor (HGF) combination gene therapy to treat preformed corneal fibrosis using established rabbit in vivo and human in vitro models.

**METHODS.** Eighteen New Zealand White rabbits were used. Corneal fibrosis was produced by alkali injury. Twenty-four hours after scar formation, cornea received topically either balanced salt solution (BSS; n = 6), polyethyleneimine-conjugated gold nanoparticle (PEI2-GNP)-naked plasmid (n = 6) or PEI2-GNP plasmids expressing BMP7 and HGF genes (n = 6). Donor human corneas were used to obtain primary human corneal fibroblasts and myofibroblasts for mechanistic studies. Gene therapy effects on corneal fibrosis and ocular safety were evaluated by slit-lamp microscope, stereo microscopes, quantitative real-time PCR, immunofluorescence, TUNEL, modified MacDonald-Shadduck scoring system, and Draize tests.

**RESULTS.** PEI2-GNP-mediated BMP7þHGF gene therapy significantly decreased corneal fibrosis in live rabbits in vivo (Fantes scale was 0.6 in BMP7þHGF-treated eyes compared to 3.3 in −therapy group; P < 0.001). Corneas that received BMP7þHGF demonstrated significantly reduced mRNA levels of profibrotic genes: α-SMA (3.2-fold; P < 0.01), fibronectin (2.5-fold, P < 0.01), collagen I (2.1-fold, P < 0.01), collagen III (1.6-fold, P < 0.01), and collagen IV (1.9-fold, P < 0.01) compared to the −therapy corneas. Furthermore, BMP7þHGF-treated corneas showed significantly fewer myofibroblasts compared to the −therapy controls (83%; P < 0.001). The PEI2-GNP introduced >10⁴ gene copies per microgram DNA of BMP7 and HGF genes. The recombinant HGF rendered apoptosis in corneal myofibroblasts but not in fibroblasts. Localized topical BMP7þHGF therapy showed no ocular toxicity.

**CONCLUSIONS.** Localized topical BMP7þHGF gene therapy treats corneal fibrosis and restores transparency in vivo mitigating excessive healing and rendering selective apoptosis in myofibroblasts.

Injuries and infections of the eye compromise corneal transparency, which accounts for two-thirds of the eye’s refractive and visual loss in an estimated 1.3 million Americans annually. Worldwide, corneal disorders are the third leading cause of preventable blindness.1,2 Despite increasing knowledge about the molecular mechanisms underlying corneal scarring,3 currently available therapies4 rely primarily on steroids and other drugs that carry the risk of multiple side effects,5 hence require repeated applications, and are often ineffective in restoring vision completely. Although mitomycin C (MMC) is a commonly used topical treatment for corneal fibrosis,6 its use continues to be the subject of debate due to its long-term side effects. A limbal graft is the treatment of choice in countries where donor corneal limbus is available.7 Corneal transplantation remains the gold standard for the treatment of corneal scars and restoration of vision.8 In 2015 alone, 48,792 corneal transplantation surgeries were performed in the United States, according to the Eye Bank Association of America (http://restoresight.org/wp-content/uploads/2016/03/2015-Statistical-Report.pdf), and about 12.7 million people in the world are awaiting donor corneal tissues.9 Besides the limited availability of donor corneas, a high immunologic rejection rate is another limiting factor for restoring vision from corneal transplantation. Thus, there is an unanswerable need for the development of effective and safe nonsurgical targeted treatments for corneal fibrosis, including gene-based therapies that would provide long-term effectiveness, require minimal clinical follow up, and show minimal side effects.
Studies show that injury and infection to the cornea/eye leads to the activation of quiescent stromal keratocytes in the cornea, which then migrate to the wound site and trans-differentiate to a wound-repair phenotype, referred to as myofibroblasts. While molecular signaling and secretory functions of myofibroblasts are essential for proper corneal wound repair, their continued production and prolonged presence in the stroma lead to corneal fibrosis (haze or scarring) due to the extended activity of TGF-β signaling during corneal wound repair. The activation and differentiation of keratocytes have been shown to occur as a response to IL-1α/IL-1β, which are released by corneal epithelial cells after injury. In addition, multiple molecules, factors, ligands, and cytokines, including TGF-β released from injured epithelial cells, play a pivotal role in the induction of inflammation and corneal scarring due to excessive biological activities and fusion of extracellular matrix (ECM) and cytoskeletal proteins. TGF-β signaling is largely responsible for the trans-differentiation of keratocytes into myofibroblasts. The persistence of myofibroblasts after wound healing is known to be a major factor in the pathogenesis of corneal fibrosis and opacity, and ultimately, vision impairment. The TGF-β superfamily proteins activate downstream signaling via the Smad family of proteins. Bone morphogenetic protein (BMP) belongs to the TGF-β superfamily and plays a significant role in ECM synthesis, tissue repair, and remodeling processes during corneal wound healing. The signaling protein BMP7 was originally described to have a significant role in the development of mammalian organs such as the kidney and the eye. BMP7 binds to the type I and II receptors and regulates receptor-regulated Smads (Smad1, Smad5, and Smad8) and inhibitory Smads (Smad6 and Smad7) in a complex wound-healing signaling network. In addition to BMP7, the expression of hepatocyte growth factor (HGF) and its receptor proteins has been found in the cornea, lacrimal glands, and tears. HGF has been identified as a mitogen that functions through the c-Met receptor tyrosine kinase in the protection and regeneration of organs as well as in the modulation of corneal repair. Injury to corneal epithelium has been shown to upregulate HGF expression in keratocytes, and in addition to its intracrine and autocrine functions, HGF has been shown to function in a paracrine manner in modulating corneal wound healing. The role of HGF and the c-Met system in diabetic corneal wound healing was recently well established in organotypic human diabetic corneal cultures. Furthermore, HGF has been reported to have a role in the breakdown of ECM deposits and in the reduction of fibrosis in several nonocular tissues. Despite the important role of HGF in corneal wound healing and TGF-β profibrotic signaling, mechanistic knowledge about the crosstalk between HGF and BMP7 in the cornea remains unknown, especially during wound healing and profibrotic microenvironment.

The cornea represents a perfect tissue for gene therapy because of its well-defined characteristics such as transparency, simple anatomy, and ease of access, allowing topical instillation of gene delivery vectors and visual monitoring of the genes packaged within vectors. In addition, therapeutic response can also be assessed noninvasively with high-resolution ocular imaging using stereo and slit-lamp biomicroscopy. Our group previously reported the advantages of tissue-specific gene delivery using a variety of modalities, including direct instillation of hybrid nanoparticle or adenovirus vectors. We found that nonviral gene delivery systems based on synthetic polycations, such as polyethylene amine (PEI), show promise as delivery systems for gene therapy. They possess DNA-binding capabilities, provide options for functionalization, and show a good safety profile. In subsequent studies, we greatly enhanced the otherwise low transfection efficiency of PEI (2 kDa) by conjugating PEI with gold nanoparticles to synthesize PEI-conjugated gold nanoparticles (PEI2-GNP). Utilizing these hybrid gold nanoparticles, we established a novel nanoparticle-based gene delivery system for the cornea that demonstrates efficient gene transfer into rabbit corneal stroma in vivo with negligible toxicity.

BMP7 and HGF are attractive targets for modulating the profibrotic signaling pathways in corneal wound healing. Our previous study of BMP7 gene therapy by PEI2-GNP in the preclinical rabbit model of corneal fibrosis revealed that PEI2-GNP-based BMP7 gene therapy led to significant inhibition of corneal fibrosis and corneal repair through counterbalancing the deleterious effects of TGF-β-induced Smad signaling. HGF gene therapy has been shown by other investigators to regulate fibrosis in various nonocular tissues, including the liver, lung, and kidney. Phase-I and phase-II clinical trials indicate HGF gene therapy is safe in humans. Furthermore, HGF gene transfer has been found to cause selective apoptosis of myofibroblasts in nonocular tissues. These reports led to an innovative postulate that PEI2-GNP-mediated tissue-targeted localized BMP7+HGF gene therapy in rabbit cornea would effectively eliminate preexisting fibrosis in vivo without producing significant toxicity. The present study tested the therapeutic potential of PEI2-GNP-delivered BMP7+HGF gene therapy for abolishing preexisting corneal fibrosis in vivo using a well-established preclinical rabbit model of corneal fibrosis.

**Materials and Methods**

**Corneal Fibrosis Induction and Treatment in Rabbits**

Eighteen New Zealand White female rabbits, weighing 2 to 3 kg (Covance Research Products, Denver, PA, USA) were used in the study. Institutional approval of the study was obtained from the Harry S. Truman Memorial Veterans’ Hospital and the Institutional Animal Care and Use Committee of the University of Missouri (both in Columbia, MO, USA). All animals were treated in accordance with the principles of the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research. The rabbits were anesthetized by a mixture of ketamine hydrochloride (50 mg/kg) and xylazine hydrochloride (10 mg/kg), given intramuscularly, for induction of corneal alkali-induced wounding, for administration of PEI2-GNP-mediated BMP7+HGF gene delivery to the corneal stroma, and for the performance of clinical slit-lamp eye examinations and ocular stereo biomicroscopy. Topical ophthalmic proparacaine hydrochloride (0.5%; Alcon, Fort Worth, TX, USA) was administered for local anesthesia prior to all procedures.

**IOP Monitoring by Tonometry**

Variations in IOP, an indicator of an ocular abnormality, may result from inflammation, swelling, rigidity, abrasion, and irregularities in corneal tissues. Administration of therapeutic genes into stroma has potential for alterations in the aqueous humor or in tissues of the anterior chamber, which is a significant concern after gene therapy. Thus, IOP measurements in rabbit eyes were recorded using a tonometer (Tono-Pen AVIA; Reichert Technologies, Depew, NY, USA) at regular timed intervals on days 1, 7, 14, and 21 and before each clinical biomicroscopy evaluation as reported earlier. All IOP measurements were performed between 9 AM and 11 AM to minimize normal diurnal variations in IOP.
**In Vivo Alkali-Induced Corneal Scarring**

Corneal scarring was induced in one eye of each rabbit, and the contralateral eye served as a naive control. To induce corneal scarring, rabbits were anesthetized and an 8-mm filter paper soaked in 0.5 N sodium hydroxide solution was applied onto the central cornea for 1.0 minutes under visualization with the surgical microscope (Leica Wild Microscope MEL53; Leica, Wetzlar, Germany). The wounded corneas were immediately and copiously rinsed with sterile balanced salt solution (BSS) to remove residual material. This method triggered wound healing and produced dense corneal scarring and peak fibrosis at 3 weeks with minimal neovascularization.11

**PEI2-GNP Transfection Solution**

Thiol-modified PEI2-GNPs were synthesized as described earlier.17 The PEI2-GNP transfection solution was prepared as reported previously.17 In brief, the PEI2-GNPs were mixed with plasmid at a nitrogen-to-phosphate (N/P) ratio of 180 by stirring 37.5 μL of 150 mM PEI2-GNPs with 10 μg plasmid DNA (pTRUF11 expressing HGF or BMP7 under control of hybrid cytomegalovirus [CMV] chicken β-actin promoter), 10% glucose (wt/vol), and bringing the volume to 100 μL with BSS. The solution was incubated at 37°C for 30 minutes prior to application on the cornea.

**In Vivo Gene Delivery**

One eye of each animal was treated and the contralateral eye served as naive control. To determine the effectiveness of gene therapy for preexisting corneal fibrosis, PEI2-GNP–mediated BMP7–HGF gene therapy was delivered into rabbit stroma 24 hours after alkali injury. BSS or transfection solution was topically applied to the cornea for 5 minutes using a cloning cylinder, as previously reported.17,19 The rabbits were divided into three groups: group 1 rabbits received BSS alone (n = 6; no gene transfer naive group); group 2 rabbits received PEI2-GNP–naked plasmid without BMP7 or HGF gene (n = 6; –therapy group); and group 3 rabbits (n = 6) received PEI2-GNP plasmids expressing HGF and BMP7 genes (n = 6; +therapy group). The cloning cylinder method is known to deliver significant levels of therapeutic genes into rabbit stroma in vivo with low toxicity.17,19

**Slit-Lamp Biomicroscopy, Haze Quantification, and Fluorescein Eye Test**

Corneal defects and general ocular health were documented at baseline and after alkali wounding at various time points using a handheld slit-lamp microscope equipped with a digital-imaging system (SL-15: Kowa Optimed, Torrance, CA, USA). Using the Fantes scale, the intensity of corneal haze was graded by three independent observers (SG, AS, MK) masked to the treatment group, as reported earlier.28,42 In brief, the grading system was the following: grade 0, completely clear cornea; grade 0.5, trace haze seen with careful oblique illumination; grade 1, more obvious haze but not interfering the visualization of fine iris details; grade 2, mild obscurcation of iris details; grade 3, moderate obscurcation of the iris and lens; and grade 4, complete opacification of the stroma in the area of ablation.

Fluorescein sodium sterile ophthalmic strips (0.6 mg, Ful-Glo; Akorn, Lake Forest, IL, USA) were used to evaluate the health of the corneal epithelium and tear production. In brief, a BSS-moistened fluorescein strip was applied to the dorsal upper eyelid of rabbits, and the rabbits were allowed to blink for distributing fluorescein stain over the entire corneal and conjunctival surface. After 30 seconds, corneal epithelium was observed under blue light using wide and narrow beams of the slit-lamp microscope. Images were obtained with a stereo microscope fitted with a fluorescence filter, spot camera, and imaging software. Tear levels were measured using commercial diagnostic strips (Tear-Flow Diagnostic Strips; HUB Pharmaceuticals, Rancho Cucamonga, CA, USA).

**Ocular Irritation Tests**

The ocular health and anomalies were evaluated independently at selected times by at least two of three examiners (MF, AS, and SG) using the established Draize13 and modified MacDonald-Shadduck41 ocular scoring systems. With the Draize eye test, the severity of ocular lesions was scored in the following manner: in the cornea, by estimating the degree of opacity and area of involvement; in the iris, by examining pupillary light reflexes; and in the conjunctiva, by assessing the degree of redness, chemosis, and discharge. With the modified MacDonald-Shadduck scoring system, ocular health scores were determined based on the cumulative average scores for corneal tissue (opacity, affected area, corneal neovascularization severity, and reepithelization) and for conjunctival tissue (congestion, chemosis, swelling, and discharge).

**Euthanasia and Tissue Collection**

Rabbits were humanely euthanized with pentobarbital (150 mg/kg) while they were under general anesthesia. Corneas were harvested, immediately placed in 15 × 5 × 5-mm molds (Fisher Scientific, Pittsburgh, PA, USA) containing optical cutting temperature (OCT) compound, and snap frozen by immersion in a cryo-cup containing 2-methylbutane sitting in liquid nitrogen. Frozen tissue blocks were preserved at –80°C. The corneas were cut into two equal halves; one half was used for histologic studies, and other half was used for molecular studies. For histologic studies, serial corneal sections (8 μm) were prepared with a cryostat (HM525 RX UV; Microm GmbH, Walldorf, Germany), placed on glass microscopic slides (Superfrost Plus; Fisher Scientific, Pittsburgh, PA, USA), and kept at –80°C until analysis. For molecular studies, corneas were cut into small pieces, immediately immersed in a cryo-cup placed in liquid nitrogen, and subsequently ground and processed for obtaining genomic DNA, mRNA, and cDNA following vendor’s protocols (Qiagen, Germantown, MD, USA).

**Hematoxylin and Eosin, Masson’s Trichome, Immunofluorescence and TUNEL Staining**

Hematoxylin-eosin (H&E) and Masson’s trichome staining were performed using the standard procedure for visualizing morphologic details, as reported earlier.40,45 Immunofluorescence staining was performed following reported methods17 to measure myofibroblasts in the corneas using antibody-specific α-smooth muscle actin (α-SMA), a marker for myofibroblasts. Briefly, corneal sections were blocked with 2% bovine serum albumin at room temperature for 30 minutes, followed by incubation with α-SMA mouse monoclonal primary antibody (1:200 dilution, M0851; Dako, Carpantaria, CA, USA) for 90 minutes and then incubated with Alexa-Fluor 488 goat anti-mouse IgG secondary antibody (1:1000 dilution, A11001; Invitrogen, Carlsbad, CA, USA) for 1 hour at room temperature. Appropriate positive and negative controls were included in each immunostaining. Quantification of α-SMA-positive cells was performed in six randomly selected, nonoverlapping, full-thickness central corneal columns, extending from the anterior stromal surface to the posterior stromal surface at 200× and 400× magnification fields.
The toxicity of PEI2-GNPs and BMP7/HGF gene therapy was determined by performing a TUNEL assay (ApopTag; Millipore, Temecula, CA, USA). Corneal sections were fixed in acetone at -20°C for 10 minutes, and a TUNEL assay was performed per the manufacturer's instructions, including suitable positive and negative controls. Rhodamine-conjugated apoptotic cells (red) and 4',6-diamidine-2'-phenylindole dihydrochloride (DAPI)-stained nuclei (blue) were viewed and photographed with a fluorescence microscope (Leica) fitted with a digital camera system (SpotCamRT KE; Diagnostic Instruments, Sterling Heights, MI, USA). DAPI-stained nuclei and TUNEL-positive cells in untreated and treated tissues were quantified at 200× and 400× magnification in six randomly selected nonoverlapping areas, as previously reported.17,28

RNA Extraction, cDNA Synthesis, and Quantitative Real-Time PCR

Total RNA from tissues was extracted with an RNeasy kit (Qiagen, Valencia, CA, USA) and reverse transcribed to cDNA following reported methods.17,28 Real-time PCR was performed using the StepOne Plus PCR system (Applied Biosystems, Carlsbad, CA, USA). A 20-μL reaction mixture contained 2 μL cDNA, 2 μL forward and reverse primers (200 nM each), and 10 μL 2X All-In-One quantitative PCR (qPCR) mix (GeneCopoeia, Rockville, MD, USA) and was run at a universal cycle (95°C for 10 minutes, 40 cycles at 95°C for 15 seconds, and 60°C for 60 seconds), as previously reported.16 The primer sequences of genes were the following: z-SMA—forward TGG GTG ACG AAC AGA AGC GA and reverse CTT CAG GGG CAA CAC GAA GC; fibronectin—forward CGG AGC TTC GAG ATC GTG C and reverse TCG ACG GGA TCA CCT TCA CA; collagen type I—forward AGA ACA GCG AAC GCT GTG AGA CTA and reverse CCA AGC TCC GCA CAA ATT TCT GA; collagen type IV—forward TAT CGA ACA AGC AAG GGC TGT GAG A and reverse GCC CAA GCT CCA CAC CAA ATT CT; and β-actin—forward CCG GTA CAG CTT CAC CA and reverse CGG GCA CGT CGT TCA GTC TCC. The β-actin was used for the normalization of qPCR data, and showed no appreciable relative fold change at various tested points and groups.

Quantification of Nanoparticle-Delivered Gene Copies

The frozen corneal tissues were pulverized in liquid nitrogen, and genomic DNA was isolated (DNA Easy kit; Qiagen, Valencia, CA, USA). Quantitative PCR was performed to determine PEI2-GNP–delivered gene copies of HGF and BMP7 in corneal tissues.17 A 10-fold serial dilution of plasmid having test gene (10⁴–10⁹/mg DNA) was used for standard curves. The qPCR settings were 95°C for 10 minutes, 40 cycles at 95°C for 15 seconds, and 60°C for 60 seconds.

Human Corneal Fibroblast and Myofibroblast Cultures

Primary human corneal fibroblast (HCF) cultures were generated from donor human corneas purchased from an eye bank (Saving Sight, Kansas City, MO, USA) following methods described previously.30 The corneal epithelium and endothelium were removed from corneal buttons with a surgical blade and cut into small pieces, placed on culture dishes, and incubated for 3 to 5 weeks in a humidified CO₂ (5%) incubator at 37°C in MEM supplemented with 10% fetal bovine serum. Seventy percent confluent HCF cultures that underwent fewer than four passages were used in the experiments. Human corneal myofibroblasts were produced by culturing HCFs under serum-free conditions in the presence of TGF-β1 (PeproTech, Rocky Hill, NJ, USA), 5 ng/ml for 72 hours.

Lactate Dehydrogenase Assay

Lactate dehydrogenase (LDH) assay was performed to measure cytoplasm enzyme LDH released in the extracellular medium. Briefly, 4.0 × 10⁵ cells were seeded in 96-well plates. LDH titers of fibroblast and myofibroblast cultures were assessed in the presence and absence of HGF conditions. After 24 hours, the supernatants were collected from each well. Cell monolayer was then treated with a cell lysis solution for 30 minutes at room temperature. Cell lysate and supernatant were collected. LDH activity in samples was measured using a toxicology assay kit (TOX7; Sigma-Aldrich Corp., St. Louis, MO, USA) following the manufacturer's instructions. The absorbance was determined at 490 nm, and LDH activity was expressed in international units per milliliter.

Statistical Analysis

Quantification studies were performed using Student’s t-test, 1-way and 2-way ANOVA followed by Bonferroni multiple comparisons test, or the Wilcoxon rank sum test and/or Tukey's multiple comparison tests. The results are expressed as mean ± standard error of the mean (SEM). P values < 0.05 are considered to be statistically significant.

RESULTS

Effects of BMP7/HGF on Corneal Fibrosis and Corneal Transparency

PEI2-GNP-mediated BMP7 and HGF gene therapy targeted to the corneal stroma significantly dissipated preexisting corneal fibrosis and restored corneal transparency in vivo (Figs. 1A–J). The corneas of all rabbits were transparent before injury (Figs. 1A, 1F) and developed dense scar/fibrosis 24 hours after alkali wound (Figs. 1B, 1G), and gene transfer was administered at this time. Rabbit corneas that received BMP7/HGF genes (+therapy group) not only demonstrated a marked reduction in corneal fibrosis but also concurrent increased transparency on day 7 (Fig. 1H), day 14 (Fig. 1I), and day 21 (Fig. 1J) as compared to the corneas that received naked vector (–therapy group; Figs. 1C, 1D, 1E).

Figure 2 depicts quantitative corneal clinical haze scores, based on Fantes scoring, assigned by three independent observers (SG, MK, AS) masked to the treatment group after alkali wound at days 1, 7, 14, and 21. On days 14 and 21, corneas that received PEI2-GNP–mediated BMP7/HGF therapy showed markedly lower corneal haze scores as compared to BSS-treated corneas. The BMP7/HGF gene delivery rendered a statistically significant decrease in corneal fibrosis at day 14 (2.3-fold; P < 0.01) and day 21 (5.5-fold; P < 0.001) as compared to the corresponding naked vector–delivered control corneas. As expected, no significant difference in corneal haze scores was observed in the –therapy and –therapy groups on day 1 and day 7. The comparison of the +therapy versus naive eyes at day 1, 7, and 14 showed significant corneal haze, but not at day 21 (P = 0.215).
Effects of BMP7+HGF on Healing of the Corneal Stroma and Epithelium

BMP7+HGF overexpression prevented excessive scarring in corneal stroma, and normal physiological healing was observed. Figures 3A and 3B show representative slit-lamp biomicroscopy images on day 21 in the −therapy and +therapy groups. A significant reduction in corneal haze and fibrosis was observed at day 21 in corneas that received PEI2-GNP-mediated BMP7+HGF (Fig. 3B) as compared to corneas that received PEI2-GNP-naked plasmid (Fig. 3A). Haze scores were 0.6 versus 3.3 (P < 0.001) in the +therapy and −therapy groups, respectively. Moreover, at day 21 subjective ocular examination of rabbit eyes in the BMP7+HGF group appeared normal in contrast to the eyes of the vector-only group, which showed mild edema, inflammation, redness, and discharge.

Fluorescein staining revealed that BMP7+HGF gene transfer is not detrimental to re-epithelialization of the injured corneas (Fig. 4). Fluorescence stereo biomicroscopy images showed corneal ulceration and epithelial defect on day 1 after alkali injury (Figs. 4A, 4C), but by day 21 all of the treated rabbits demonstrated complete corneal re-epithelialization and negative fluorescein staining (Figs. 4B, 4D). The lacrimal lake, as observed on tear strips, appeared similar in both PEI2-GNP-naked vector and BMP7+HGF groups, suggesting that combination therapy with BMP7+HGF does not compromise tear production or invoke a dry eye condition.

Myofibroblast and Profibrotic Gene Expression

Determination of myofibroblast after alkali-induced corneal injury indicates that BMP7+HGF gene therapy has an inhibitory effect on myofibroblast formation and profibrotic gene expression (Fig. 5). Immunohistologic staining of α-SMA, a marker for myofibroblasts, demonstrated a clinically relevant and statistically significant reduction in α-SMA-positive cells in corneas that received BMP7+HGF genes (Fig. 5B) as compared to corneas of the naive and PEI2-GNP-naked plasmids (Fig. 5A). The immunofluorescence quantification graph (Fig. 5C) showed an 83% reduction of α-SMA-positive cells (P < 0.001) in the +therapy group as compared to the number of α-SMA-positive cells in the −therapy group (Fig. 5B).

Since BMP7+HGF gene delivery into fibrotic rabbit corneas led to a quantifiable improvement in wound healing after alkali burn in our in vivo model, the expression of five prominent genes (α-SMA, fibronectin, collagen I, collagen III, and collagen IV) involved in fibrosis pathways was also examined in the corneas of rabbits that received BSS, PEI2-GNP-naked plasmid or BMP7+HGF genes (Fig. 6). As Figure 6 illustrates, alkali injury significantly increased levels of the five tested profibrotic genes in the −therapy corneas compared to naive corneas from 2.2- to 5.2-fold (P < 0.001), whereas BMP7+HGF gene transfer significantly reduced profibrotic gene expression of the α-SMA, a 3.2-fold reduction (Fig. 6A; P < 0.01); fibronectin, a 2.3-fold reduction (Fig. 6B; P < 0.01); collagen I, a 2.1-fold reduction (Fig. 6C; P < 0.01); collagen III, a 1.6-fold reduction (Fig. 6C; P < 0.01), and collagen IV, a 1.9-fold reduction (Fig. 6C; P < 0.01).

BMP7+HGF Dissipates Myofibroblasts via Apoptosis

Double immunofluorescence staining of corneal tissue sections for α-SMA (a myofibroblast marker) and TUNEL (an apoptosis marker) demonstrated that HGF delivery dissipates myofibroblasts through apoptosis in the fibrotic cornea. Figure 7 shows
FIGURE 3. Slit-lamp images at day 21 after administration of PEI2-GNP only (A) and of PEI2-GNP-mediated BMP7þHGF (B), showing persistence of haze in the –therapy eye and a clear cornea after BMP7þHGF. Scale bar: 2 mm.

FIGURE 4. Stereo fluorescence microscopy images on days 1 and 21 post injury of eyes treated with PEI2-GNP alone (A, B) or BMP7þHGF (C, D), showing that therapy does not affect reepithelialization of the injured corneas. Scale bar: 2 mm.

FIGURE 5. Immunofluorescence images showing stromal expression of α-SMA, a myofibroblast marker, in corneas 21 days post injury in rabbits that did not receive gene therapy (A) and those that received BMP7þHGF gene therapy (B). Quantification graph depicts the significant reduction of α-SMA in the +therapy group compared to the –therapy group (P < 0.001). Scale bar: 100 μm.
representative images of double immunofluorescence performed in corneal tissue collected on day 21 after injury. The anterior stromal tissue of BMP7þHGF-treated corneas was found to contain several double-stained α-SMA-positive and TUNEL-positive cells (4.7 ± 1.2/400×) and single-stained α-SMA cells (4.5 ± 1.1/400×) and TUNEL-positive cells (21.4 ± 5.7/400×) (Fig. 7B). In contrast, the anterior stromal tissue of corneas that received PEI2-GNP-naked vector showed only α-SMA-positive cells (41.7 ± 6.1/400×) (Fig. 7A). The difference between the þtherapy and the −therapy group in the disappearance of myofibroblast via apoptosis was statistically significant (P < 0.001).

Figure 6. Bar graphs showing differential mRNA expression of (A) α-SMA, (B) fibronectin (FN), and (C) collagen I (Col I), collagen III (Col III), and collagen IV (Col IV) in the corneal tissue from rabbits in the naive, PEI2-GNP (−therapy), and BMP7þHGF (+therapy) groups. Tissue was obtained 21 days after injury. BMP7þHGF treatment reduced fibrotic gene expression in injured cornea. Quantification of mRNA expression of fibrotic-related genes by real-time PCR. Graphs depict relative fold expression of α-SMA, fibronectin, collagen I, collagen III, and collagen IV (n = 6 for each group). Error bars represent ± SEM (**P < 0.001 against naive control and *P < 0.01 against BMP7þHGF group, respectively).

Number of Gene Copies Driving Therapeutic Response

Real-time PCR quantification of the number of therapeutic gene copies responsible for reducing corneal fibrosis indicated that sufficient delivery of the genes was achieved with the PEI2-GNPs administered via cloning cylinder technique (Figs. 8A, 8B). Gene delivery by the PEI2-GNPs was significant, as 4.3 × 10⁴ ± 0.2 copies of BMP7 per 1 µg DNA (Fig. 8A) and 3.2 × 10⁴ ± 0.4 copies of HGF per 1 µg DNA (Fig. 8B) were detected in treated corneas.

Figure 7. Double immunofluorescence staining for α-SMA and TUNEL in tissue from injured corneas treated with (A) PEI2-GNP alone (−therapy) and (B) PEI2-GNP-delivered BMP7þHGF (+therapy). Arrows point to α-SMA; arrowbeads point to colocalization of α-SMA-positive and TUNEL-positive cells; and arrowbeads with inside tail point to TUNEL-positive cells. Scale bar: 25 µm.
In Vivo Toxicity and Safety Studies

The results of time-dependent ocular irritation studies performed in live rabbits using the Draize and modified MacDonald-Shadduck scoring systems are highlighted in the Table. As expected, an alkali wound led to a significantly increased cumulative Draize score in the PEI2-GNP-naked vector and PEI2-mediated BMP7+HGF group as compared to naive corneas. On day 7, the average Draize score was 45.0 in the –therapy group versus 0 in the naive group (P < 0.001); on day 14, 39.0 vs. 0 (P < 0.001); and on day 21, 29.8 vs. 0 (P < 0.001). BMP7+HGF therapy was associated with a significant, time-dependent reduction in the cumulative Draize score compared to the score in the –therapy group as follows: day 7, 29.1 with BMP7+HGF vs. 45.0 with –therapy (P < 0.01); day 14, 18.9 vs. 39.0 (P < 0.01); and day 21, 5.1 vs. 29.8 (P < 0.01). The modified MacDonald-Shadduck test results showed a similar pattern of scores following alkali wounding in PEI2-GNP-naked vector and BMP7+HFG groups. Injured eyes that received BMP7+HGF therapy exhibited significantly lower modified MacDonald-Shadduck scores than did the eyes of rabbits that received PEI2-GNP-naked vector (day 7: 0.9 vs. 1.8, P < 0.05; day 14: 0.6 vs. 1.3, P < 0.01; day 21: 0.3 vs. 1.1, P < 0.001). On the subjective clinical eye examinations performed independently by three examiners, significantly improved overall ocular health was observed in the eyes of rabbits that received gene therapy as compared to eyes of rabbits that received the nanoparticles alone.

The slit-lamp and stereo biomicroscopic images were scored by three independent resident observers. The naive group did not show any ocular anomaly during Draize and modified MacDonald-Shadduck scoring (P < 0.001 or P < 0.01 or P < 0.05 between the groups at different time points).

TABLE. Draize and Modified MacDonald-Shadduck Scoring Shows the Ocular Anomaly in –Therapy and +Therapy Groups

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<td>Day 14</td>
<td>Day 21</td>
<td></td>
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Role of HGF in Apoptosis of Corneal Myofibroblasts and Fibroblasts

The molecular function of HGF on corneal fibroblasts and myofibroblasts was studied in an established in vitro model of corneal fibrosis in humans. Primary stromal cultures obtained from donor human corneas and grown in the absence of TGF-β1 provided corneal fibroblasts and grown in the presence of TGF-β1 produced corneal myofibroblasts, as demonstrated by α-SMA staining and TUNEL staining. Recombinant HGF (rHGF) treatment of these cultures caused significant apoptosis, as indicated by the detection of several TUNEL-positive cells in (Fig. 9; P > 0.1) and tear levels (data not shown) in the eyes of rabbits among the three groups.

H&E staining demonstrated noteworthy morphologic alterations in corneal epithelium and stroma at day 21 post alkali injury in the rabbits that received PEI2-GNP-naked vector (Fig. 10A), corroborating the presence of the clinical opacification visualized with slit-lamp biomicroscopy. BMP7+HGF therapy markedly mitigated the adverse impact of alkali wounding on corneal epithelial and stromal tissues (Fig. 10B). In addition, no significant cellular inflammatory infiltrates were observed in the corneas of either group, as quantified with CD11b immunofluorescence (Figs. 10C, 10D; P > 0.1). The Masson’s trichrome staining demonstrated notably decreased collagen deposition in the rabbit corneas that received BMP7+HGF therapy (Fig. 10F) compared to the –therapy naked vector-delivered corneas (Fig. 10E).
We hypothesized that HGF gene transfer in combination with BMP7 locally into the opaque cornea could reverse the fibrotic events in vivo in a preclinical rabbit model of corneal fibrosis. The results of this study demonstrate that delivery of BMP7+HGF genes into stromal fibroblasts/keratocytes via nonviral vectors (PEI-GNP) significantly reduced corneal opacity by 3 weeks post injury in a preclinical rabbit model of corneal fibrosis (Figs. 1–4). Importantly, since the BMP7+HGF therapy was administered 1 day after injury, the findings suggest that resolution of corneal opacity and vision restoration is achievable even in a significantly damaged cornea in vivo. Furthermore, administration of BMP7+HGF gene therapy was associated with a significant reduction in α-SMA, a molecular marker for myofibroblasts, and a concomitant decrease in profibrotic genes (Figs. 5 and 6). To corroborate the selective apoptosis observed in rabbit myofibroblasts was driven by HGF, we performed in vitro studies of the effects of tHGF on human corneal myofibroblasts and fibroblasts, which demonstrated that tHGF induces apoptosis in human corneal myofibroblasts but not in HCFs (Figs. 11 and 12). This finding agrees with earlier reports of HGF function in pulmonary and liver fibrosis models, which suggests that HGF-induced myofibroblast cell death is a key event in the healing process. 53 TGF-β signaling has been shown to increase the expression of c-Met, the receptor for HGF,54 suggesting that in a fibrotic microenvironment the HGF activity may be more pronounced in myofibroblasts than in quiescent stromal keratocytes. The c-Met receptor, a proto-oncogene, has multiple functions, including cell survival via sequestration of the Fas receptor and inhibition of death-domain–induced signaling.55 However, in the presence of HGF, the binding of c-Met with Fas receptors would be dampened due to c-Met activation by phosphorylation, and the cells can then respond to Fas ligand–dependent death signals.56 In addition, HGF also induces increased caspase-3 activity via intracellular signaling.57 Caspase-3–dependent cleavage of the c-Met receptor converts it into a 40-kDa proapoptotic signal.58 Therefore, the selective death of myofibroblasts observed by HGF treatment could be the result of a combination of three factors: increased c-Met receptors on myofibroblasts, elevated caspase activity, and the generation of profibrotic c-Met fragments. HGF has also been shown in lung fibrosis models to increase matrix metalloproteinase expression, leading to cell death and a concomitant reduction in ECM.59 Hence, increased matrix metalloproteinase levels could be a mechanism to prevent corneal scar formation and aberrant rearrangement of the ECM in the presence of HGF.

On the other hand, BMP7 is well known to antagonize the TGF-β pathway15,17 via upregulation of the Id3 proteins50 and Smad 7 signaling.60 Coadministration of BMP7 and HGF has been found therapeutic in renal fibrosis and tubular nephropathy.61 Expression of profibrotic factors such as α-SMA, fibronectin, and collagens has been shown to decrease in response to BMP7 treatment.17 Therefore, the two therapeutic genes, BMP7 and HGF, represent two independent pathways for controlling corneal wound healing without scarring, namely by regulating TGF-β signaling and by selective apoptosis of myofibroblasts. Such a combination treatment modality, targeting the initial signaling phase (via BMP7) and the downstream myofibroblast apoptosis and resolution (via HGF), can therefore have a greater potential for disease resolution even in advanced cases of fibrosis. To the best of our knowledge, this is a first study that demonstrates elimination of established corneal scar and restoration of corneal transparency in vivo by activating multiple signaling pathways. This innovative approach is expected to lead development of the first nonsurgical gene-based therapies to cure preexisting corneal fibrosis/scarring without significant adverse effects.
Combination BMP7 and HGF gene therapy is safe and caused no adverse effects in rabbit eyes. H&E images show no alteration in corneal morphology with PEI2-GNPs alone (A) and with BMP7+HGF gene therapy (B). CD11B immunofluorescence images 21 days after corneal injury show no infiltration of leukocytes and inflammatory markers in the stromal layer in corneal tissue from PEI2-GNP-treated rabbits (−therapy) and from BMP7+HGF-treated rabbits (+therapy). Masson’s trichrome-stained corneal tissue images show decreased collagen expression in BMP7+HGF-treated eyes (F) compared to the −therapy given eyes (E). Scale bar: 100 and 200 μm.

Immunocytochemistry images of α-SMA staining (green) and TUNEL staining (pink) in cell cultures of HCFs (A) and myofibroblasts (B) after rHGF treatment, demonstrating that rHGF selectively induces apoptosis of corneal myofibroblasts. Arrows point to TUNEL-positive myofibroblast cells. Scale bar: 50 μm.
Data from decades of clinical trials in other fibrotic diseases, including renal and pulmonary fibrosis, suggest that single-targeted therapy is typically less effective in disease resolution. In the clinical setting, patients who have incurred a corneal injury generally have well-established corneal haze (fibrosis) at the time of presentation to an ophthalmologist. An efficient gene delivery method as well as a multimodal-targeted treatment approach may offer the potential for achieving better results than what is possible with currently available therapies. Our study found that PEI2-GNP-delivered BMP7+HGF genes remained in the injured rabbit cornea at high copy numbers for up to 3 weeks after instillation (Fig. 8) without concomitant adverse ocular effects such as an increase in IOP (Fig. 9) or dry eye conditions. Furthermore, Draize and modified MacDonald-Shadduck scores (Table) indicate that ocular toxicity from BMP7+HGF combination gene therapy is minimal. Nevertheless, the long-term effects of HGF+BMP7 therapy on stromal collagen fibril organization and neighboring ocular tissues remain unknown and warrant further investigation. The lack of data on long-term effects is a limitation of the present study, and our future studies will investigate the long-term effects of PEI2-GNP-delivered BMP7+HGF gene therapy.

In conclusion, this study demonstrates that tissue-targeted BMP7+HGF combination gene therapy given locally with PEI2-GNPs 1 day after alkali damage eliminates corneal fibrosis and restores corneal transparency in vivo by negating TGF-β pathologic activity producing myofibroblasts and promoting apoptosis in established myofibroblasts utilizing multiple signaling pathways.

Acknowledgments

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References


