COL8A2 Regulates the Fate of Corneal Endothelial Cells

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PURPOSE. To investigate the effect of COL8A2 repression on corneal endothelial cells (CECs) in vitro and in vivo.

METHODS. Cultured human CECs (hCECs) were transfected with COL8A2 siRNA (siCOL8A2), and the cell viability and proliferation rate were measured. The expression of cell proliferation–associated molecules was evaluated by Western blotting and real-time reverse transcription PCR. Cell shape, Wingless-INT (WNT) signaling, and mitochondrial oxidative stress were also measured. For in vivo experiments, siCOL8A2 was transfected into rat CECs (rCECs), and corneal opacity and corneal endothelium were evaluated.

RESULTS. After transfection with siCOL8A2, COL8A2 expression was reduced (80%). Cell viability, cell proliferation rate, cyclin D1 expression, and the number of cells in the S-phase were reduced in siCOL8A2-treated cells. The cell attained a fibroblast-like shape, and SNAI1, pSMAD2, and β-catenin expression, along with mitochondrial mass and oxidative stress levels, were altered. Corneal opacity increased, and the CECs were changed in rats in the siCOL8A2 group.

CONCLUSIONS. COL8A2 is required to maintain normal wound healing and CEC function.

Keywords: COL8A2, human corneal endothelial cells, wound healing process

The posterior surface of the cornea is lined by a monolayer of corneal endothelial cells (CECs) that function as a water pump for the corneal stroma via ion pumps and channels. If the corneal endothelium is diseased, the cornea is not dehydrated and becomes edematous. Corneal edema not only causes blindness, but also creates painful blisters in the corneal epithelium. Persistent corneal edema requires corneal transplantation and can result from trauma, surgery, corneal endothelitis, Fuchs’ endothelial corneal dystrophy (FECD), or congenital hereditary endothelial dystrophy. A variety of mechanisms have been studied in the context of CEC disease, including oxidative stress, endoplasmic reticulum stress, aberrant DNA methylation, inflammation, apoptosis, and autophagy. The fundamental mechanism underlying CEC disease is limited regeneration after injury. CEC diseases caused by changes in the basement membrane of CECs, such as FECD, are well known. Changes in basement membrane composition may inhibit CEC regeneration.

Type VII collagen, a nonfibrillar collagen protein, is expressed by corneal and vascular endothelial cells. It is a heterodimer composed of two distinct polypeptide chains, α1(VII) and α2(VII) (genes: COL8A1 and COL8A2). COL8A1 is the main component of Descemet’s membrane (DM), the basement membrane of the corneal endothelium. COL8A2 has a potential role in the maintenance of CEC integrity and structure. COL8A2 is coexpressed with the activating enhancer-binding protein 2 (TFAP2, AP-2) during corneal development, and this has ultimately led to the onset and progression of FECD.

Mutations in COL8A2 are one of the causes of early-type FECD, a disease in which guttae are formed in the DM, progressively causing CEC dysfunction and edema, and ultimately requiring corneal transplantation. FECD associated with COL8A2 mutations exhibits early disease manifestation, especially in younger patients. COL8A2 mutations affect DM stiffness, causing guttae to form and contributing to the onset and progression of FECD. Homozygous knock-in mice with this mutation (Col8a2(Q455K/Q455K)) exhibited endoplasmic reticulum (ER) stress that led to hCEC death. Nevertheless, the role of COL8A2 in CECs is not thoroughly understood. In this study, we investigated the role of COL8A2 in CECs by inhibiting COL8A2 expression in vitro and in vivo.

METHODS

In vivo Transfection of Corneal Endothelium in Rats

This study was approved by the Institutional Animal Care and Use Committee (IACUC) of Hallym University Medical Center. All procedures were performed according to the Association for Research in Vision and Ophthalmology Statement for the Use of Animals in Ophthalmic and Vision Research. Six-week-old Sprague-Dawley (SD) rats, purchased from Nara Biotech (Pyeongtaek, Korea), were used for this procedure. Five rats were included in each group. The rats were maintained in a colony room with a
12-hour light/dark cycle at 25°C for seven days before initiating experiments.

For in vivo transfection, 1 nmol of negative control siRNA (siControl) or siRNA for COL8A2 (siCOL8A2) was injected into the anterior chamber of SD rats after paracentesis. Then, 7 mm Tweezertrodes (BTX, Holliston, MA, USA) were inserted onto each cornea, with the positive electrode on the siRNA-injected eye. The parameters were set at 140 V, 100 milliseconds length, 950 milliseconds interval, five pulses, and 100 V/cm². Then, a previously established corneal endothelial injury model was induced using cryoinjury two days after electroporation.15 The cornea was cryoinjured for 10 seconds in contact with a metal rod, 3 mm in diameter, that had been frozen in liquid nitrogen for 10 minutes and then irrigated with normal saline solution (day 0). Then, the corneas were evaluated.

Clinical Evaluation and Alizarin S Red Staining

Corneal opacity was observed by slit lamp biomicroscopy and photographed. Corneal opacity was graded by the Institutional Review Board/ethics committee of Hallym University Medical Center. The cells were cultured in OptiMem-I media (GIBCO/BRL Life Technologies, Grand Island, NY, USA) under a 12-hour light/dark cycle at 25°C for seven days before use and transfected with siRNA. TEER and TEPD were measured using an epithelial voltohmmeter (EVOM2; World Precision Instruments, Inc.) was used on the transwell TEER measurement.19 Cell proliferation was assessed by measurement of the TEER and TEPD. The surface of 1.12 cm², 0.4 μm pore-sized Transwell inserts (CLS3460, Corning Incorporated/Life Sciences, Tewksbury, Massachusetts, USA) were coated with FNC coating mix. Then, 1 mL of culture media was added to the bottom well of a 12-well tissue culture plate, and 200 μL was added to the inside of the collagen-coated Transwell insert. Cells were seeded onto each collagen-coated Transwell insert at a density of approximately 100,000 cells/well and cultured at 37°C, 5% CO₂ for seven days before use and transfected with siRNA. TEER and TEPD were measured using an epithelial voltohmmeter (EVOM2; World Precision Instrument, Inc., Saratoga, FL, USA). The wells were then washed twice with serum-free medium and allowed to equilibrate for one hour at 37°C, 5% CO₂ before measurement.18 TEER was measured in triplicate for each well. Each group contains four wells. STX2 chopstick electrode (4 mm wide and 1 mm thick; World Precision Instrument, Inc.) was used on the transwell TEER measurement.19 Each experiment was repeated in triplicate.

Cell Culture and Transfection

This study was performed in accordance with the tenets of the Declaration of Helsinki and was reviewed and approved by the institutional review board/ethics committee of Hallym University Medical Center. The cells were cultured in accordance with previously published methods.16,17 The corneas from a total of six donors were used.16 All the cells remained attached to the DM. The endothelial cell–DM complex was incubated for 10 minutes in 0.25% trypsin/0.02% ethylenediamine tetra-acetic acid (EDTA) solution (Invitrogen, Grand Island, NY, USA). The cells were then plated at the bottom of six-well plates coated with a fibronectin–collagen combination (FNC) coating mix (Athena Environmental Sciences, Inc., Baltimore, MD, USA). The cells were cultured in OptiMem-I media (GIBCO/BRL Life Technologies, Grand Island, NY, USA) supplemented with 8% fetal bovine serum (FBS; Cambrex Bio Science, Walkersville, MD, USA), 200 μg/mL calcium chloride (Sigma Chemical Co., St. Louis, MO, USA), 0.08% chondroitin sulfate (Sigma Chemical Co.), 20 μg/mL ascorbic acid (Sigma Chemical Co.), 100 μg/mL putituary extract (Invitrogen, Grand Island, NY), 5 ng/mL epidermal growth factor (Sigma Chemical Co., St. Louis, MO), 20 ng/mL nerve growth factor (Sigma Chemical Co., St. Louis, MO), 10 μg/mL gentamicin (Invitrogen), 100 IU/mL penicillin (Cambrex Bio Science), 100 IU/mL streptomycin (Cambrex Bio Science) and 2.5 μg/mL amphotericin (Cambrex Bio Science) under 5% CO₂. Medium was changed every two days. Cells were cultured for 10 to 14 days until conflueny and were then passaged at a ratio of 1:3 using 0.25% trypsin/0.02% EDTA solution.

To evaluate the transfection efficiency, fluorescein isothiocyanate (FITC)–conjugated siRNA (Bioneer, Daejeon, Korea) were transfected to the cells. To silence COL8A2 expression, we used small-interference RNA (siRNA), siRNA for COL8A2 (1296-1; sense, 5′-GUC AAG GCC ACC AAC GUG U-3′ and antisense, 5′-AGA CGU UUG UCG CCU UGA-3′) and nonspecific control siRNA (SN-1001) used as a negative control, were purchased from Bioneer Cooperation. Summarily, primary HCECs at a density of 5 × 10⁴ cells/cm² were transfected with siRNA specific for COL8A2 at 10 nmol/L concentrations, or with negative control siRNA, using Lipofectamine RNAiMAX (Invitrogen) according to the manufacturer's instructions. The transfections were performed at 70% confluency. After incubation for 48 to 72 hours, the cells were collected for further evaluation. The cells were separated into two groups: siRNA group targeting COL8A2 (siCOL8A2), and a control group (siControl). The effect of COL8A2 silencing was confirmed by western blot analysis or RT-PCR 48 hours after transfection.

Transendothelial Electrical Resistance (TEER) and Transendothelial Electrical Potential Difference (TEPD)

Endothelial barrier function was assessed by measurement of the TEER and TEPD. The surface of 1.12 cm², 0.4 μm pore-sized Transwell inserts (CLS3460, Corning Incorporated/Life Sciences, Tewksbury, Massachusetts, USA) were coated with FNC coating mix. Then, 1 mL of culture media was added to the bottom well of a 12-well tissue culture plate, and 200 μL was added to the inside of the collagen-coated Transwell insert. Cells were seeded onto each collagen-coated Transwell insert at a density of approximately 100,000 cells/well and cultured at 37°C, 5% CO₂ for seven days before use and transfected with siRNA. TEER and TEPD were measured using an epithelial voltohmmeter (EVOM2; World Precision Instrument, Inc., Saratoga, FL, USA). The wells were then washed twice with serum-free medium and allowed to equilibrate for one hour at 37°C, 5% CO₂ before measurement.18 TEER was measured in triplicate for each well. Each group contains four wells. STX2 chopstick electrode (4 mm wide and 1 mm thick; World Precision Instrument, Inc.) was used on the transwell TEER measurement.19 Each experiment was repeated in triplicate.

Cell Proliferation

Cell proliferation rate was measured using a commercial bromodeoxyuridine (BrDU) proliferation assay kit (Roche Diagnostics, GmbH, Mannheim, Germany) according to the manufacturer's protocol. Cells (5 × 10⁴ cells/well) were placed in 96-well plates (NUNC, 167008; Thermo Fisher Scientific, Denmark) and incubated for 48 hours in a humidified atmosphere containing 5% CO₂. Cells were then transfected and labelled with BrDu at 37°C and 5% CO₂ for 72 hours. After incubating the plate in the FixDenat solution for 30 minutes at room temperature, the cells were...
incubated with anti-BrdU-POD solution for approximately 90 minutes at room temperature. Then, substrate solution was added to each well, and the plate was incubated for 20 minutes at room temperature. Thereafter, 1 mol/L H$_2$SO$_4$ was added to each well to stop the reaction. The optical density was measured at 450 nm using an ELISA reader (Synergy HTX, Biotek, Winooski, VT, USA). Proliferation rates were expressed as the percentage of controls after subtraction of the corresponding blanks.

**Cell Cycle Analysis**

Cell cycle analysis was performed using the Muse cell analyzer (Merck Millipore, Burlington, MA, USA) with propidium iodide (PI) staining according to the manufacturer's protocol. Briefly, cells were grown in six-well plates and transfected with siRNA; then harvested by trypsinization and washed twice with phosphate-buffered saline solution (PBS). The cells were fixed with 1 mL of 70% cold ethanol at −20°C for five hours and washed twice with PBS. After centrifugation at 1500 rpm for five minutes, the samples were treated with 200 μL solution including 50 μg/mL PI (Merck Millipore) and 100 μg/mL RNase A (Biosesang, Seongnam, Korea). The percentage of cells in G0/G1, S and G2/M phases was then calculated using a Muse cell analyzer.

**Western Blotting**

Radioimmunoprecipitation assay buffer (Biosesang, Seoul, Korea), containing a p8340 protease inhibitor cocktail (one tablet/10 mL; Sigma-Aldrich) and PhosSTOP phosphatase inhibitor cocktail (one tablet/10 mL; Roche, Basel, Switzerland), was used to isolate total cellular proteins. Western blotting was conducted using standard protocols. Either 5% skim milk or gelatin was used for blocking the nonspecific binding for one hour. Primary antibodies were mouse antihuman COL8A2 antibody (sc-293350, Santa Cruz biotechnology, 1:200 dilution), rabbit antihuman cyclin D1 antibody (sc-718, Santa Cruz, 1:1000 dilution), mouse antihuman SNAI1 antibody (sc-271977, Santa Cruz, 1:2000 dilution), mouse antihuman pSMAD2 antibody (LF-PA20460, Abfrontier, Seoul, 1:1000 dilution), mouse antihuman SMAD2/3 antibody (sc-133098, Santa Cruz, 1:1000 dilution), rabbit antihuman GSK3β antibody (ab32391, Abcam, 1:1000 dilution), rabbit antihuman β-catenin antibody (ab325572, Abcam, 1:1000 dilution), goat anti-SLC4A11 antibody (NBPI-46156, Novusbio, 1:1000 dilution), mouse anti-notch1 antibody (sc-376403, Santa Cruz, 1:500 dilution), or rabbit anti-GAPDH antibody (LF-PA0212, Abfrontier, 1:5000 dilution). A horseradish peroxidase (HRP) conjugated secondary antibody and a peroxidase substrate solution (one tablet/10 mL; Sigma-Aldrich) were used for detection. Data were quantified by video image analysis. Protein bands were measured by densitometry (Image J, National Institutes of Health, Bethesda, MD, USA).

**Real-Time Reverse Transcription PCR**

RNA was extracted from the cultured HCECs separately using the ReliaPrep RNA Miniprep Systems (Promega Cooperation, Madison, WI, USA) according to the manufacturer's instructions. RNA concentrations were determined by ultraviolet spectrophotometry. The first-strand complementary DNA was synthesized from 0.2 μg of total RNA with oligonucleotide primers using a commercially available kit (GoScript Reverse Transcription System; Promega Cooperation). Complementary DNA samples were aliquoted and stored at −20°C until use. Real-time quantitative RT-PCR (RT-qPCR) were performed in 20-μL volumes using the AccuPower 2X GreenStar qPCR Master Mix (Bioneer) with real-time qPCR Primer Assay for human, and the following thermocycling parameters −95°C for 10 minutes then 40 cycles at 95°C for 15 seconds and at 60°C for 60 seconds. SYBR green fluorescence was measured at the end of each cycle. The β-actin gene, a housekeeping gene, was used as a standard for normalization. Primer sequences, purchased from Bioneer, were described in the Table. Assays were performed in triplicate. Melting curve analysis was performed to ensure good-quality specific PCR products. RT-qPCR results were analyzed, using the relative standard curve method and compared and calibrated against the control group.

**Mitochondrial Oxidative Stress Evaluation**

MitoSOX Red (Invitrogen) was used to measure mitochondrial superoxide production according to the manufacturer's protocol. Cells (1 × 10⁵) were cultured in six-well plates and transfected with siRNA for 48 hours. Cells were incubated with 5 μmol/L MitoSOX reagent for 30 minutes at 37°C in the dark and then washed with PBS. Cells were harvested by trypsinization and washed twice with PBS. Fluorescence intensity in each well was measured using Muse cell analyzer at an excitation wavelength of 510 nm and an emission wavelength of 590 nm. The mean intensities of MitoSOX Red fluorescence of the three samples in each group were obtained and compared.

**MitoTracker Red Staining**

MitoTracker red FM fluorescent probe (Invitrogen) was used to measure mitochondrial mass according to manufacturer's protocol. Cells (1 × 10⁵) were cultured in six-well plates and transfected with siRNA for 48 hours. Cells were incubated with MitoTracker red FM fluorescent probe at a final concentration of 200 nmol/L for 30 minutes and then washed with PBS. Cells were harvested by trypsinization and washed twice with PBS. MitoTracker red fluorescence of the three samples in each group were obtained and compared.

**Immunofluorescence Staining**

Immunofluorescence staining was carried out. Briefly, hCECs cultured on coverslips for three days were transfected with a siRNA. At 48 hours after transfection, cells were fixed in 3.7% formaldehyde and permeabilized with 0.5% Triton X-100. Cells were washed with PBS and incubated with blocking solution (normal goat serum, Invitrogen) for 30 minutes. Cells were incubated with rabbit anti-zonula occludens-1 (ZO-1) antibody (sc-10804, diluted 1:1000) at 4°C overnight. Cells were washed with PBS and incubated with fluorescein isothiocyanate-conjugated secondary goat anti-rabbit IgG (diluted 1:100; Invitrogen) antibodies for one hour at room temperature. Samples were mounted on slides with antifade mounting medium with
4′,6-diamidino-2-phenylindole (DAPI; Vector Laboratories, Burlingame, CA, USA). For negative control, primary antibody was omitted. Slides were viewed under a fluorescence microscope (DM2000; Leica).

Statistics

Data were expressed as mean ± standard deviation. Each experiment was repeated in triplicate. An independent t-test was used for comparison of two groups. Fold change means relative mRNA expression to control. Data were analyzed using GraphPad Prism software version 8.4 (GraphPad Software, Inc., San Diego, CA, USA).

RESULTS

In vivo Suppression of COL8A2

After transfecting siCOL8A2 into the corneal endothelium of SD rats, RT-qPCR was performed in rCECs. Expression of siCOL8A2 mRNA was reduced in the corneal endothelium (4.73% ± 1.54%; Fig. 2A). Corneal opacity increased in the siCOL8A2-transfected cornea at day 3, 1 week, and 2 weeks after in vivo transfection (P = 0.049, 0.008, and 0.001; Figs. 2B and 2C). Alizarin S red staining of the corneal endothelium revealed CEC elongation in the siCOL8A2-transfected cornea after in vivo transfection, which was similar to the change in shape of cultured hCECs after siCOL8A2 transfection (Fig. 2D).

In vitro Suppression of COL8A2

Cell Culture and Transfection. The hCECs were cultured and exhibited a mosaic pattern at P0 (Fig. 1A). Cells were transfected with FITC-conjugated siRNA to assess transfection efficiency. In transfected cells, FITC-conjugated siRNA is observed in the cytoplasm or nucleus as green fluorescence (Fig. 1B). Transfection efficiency was 81.67% ± 3.06%. The hCECs were transfected with siCOL8A2, which lead to a reduction of COL8A2 expression to 13.65% ± 8.23% compared with the siControl group, as determined by RT-qPCR. This result was also confirmed by Western blotting (15.8% ± 3.4%; Figs. 1C and 1D). The reduction in the expression of COL8A2 was sufficient for subsequent analysis.

Corneal Endothelial Function. To investigate barrier function, TEER and TEPD were measured after hCECs were cultured in the insert of a transwell chamber. TEER and TEPD reduced in the siCOL8A2 group (141.8 ± 10.5 Ω and −2.63 ± 0.06 mV) compared to the siControl group (165.8 ± 8.8 Ω and −2.83 ± 0.05 mV; 24.002 and 0.192 mV decrease; P = 0.004 and 0.007, respectively; Figs. 1E and 1F). Tight junctions are important for the barrier function, and the distribution of ZO1, a major component of tight junctions, was investigated by immunofluorescence staining. Immunofluorescence staining for ZO1 is shown in Figure 1G. ZO1 was densely localized and well-formed in the siControl group, and its formation decreased in slender hCECs transfected with siCOL8A2.

Cell Proliferation. Compared to siControl-treated cells, the cell proliferation rate (30.0% decrease; P = 0.006 and 0.044; Fig. 3A) and the percentage of cells in S-phase (52.8% decrease; P = 0.012; Figs. 3B and 3C) was reduced in siCOL8A2-treated cells. Next, molecules associated with cell cycle progression were investigated. Cyclin D1 protein expression (64.1% decrease; P = 0.047; Figs. 3D and 3E), and mRNA expression of PCNA and α-SMA (64.8% and 58.2% decrease; P < 0.001 and 0.002, respectively; Figs. 3F and 3G) decreased in the siCOL8A2 group. In contrast, mRNA expression of CDKN2A, an inhibitor of proliferation, was increased by 115.9% in the siCOL8A2 group (P < 0.019; Fig. 3H).

Endothelial-Mesenchymal Transition (EMT). EMT was investigated in cultured hCECs. Upon evaluating cell shape, hCECs transfected with siCOL8A2 were slender, elongated, and spindle-shaped, whereas cells in the siControl group had a polygonal shape (Fig. 4A). EMT-associated molecules were next evaluated. pSMAD2 protein levels, which is an activated form of SMAD2 that is present at elevated levels during EMT, were reduced in the siCOL8A2 group compared to that in the siControl group (84.3% decrease; P < 0.001; Fig. 4B). SNAI1 mRNA expression, which has been implicated in EMT, was reduced in the siCOL8A2 group compared with that in the siControl group (80.5% decrease; P = 0.008; Fig. 4B), and this was confirmed by Western blotting (28.5% decrease; P = 0.034; Fig. 4C). The mRNA expression of SNAI1 was increased in the siCOL8A2 group compared to that in the siControl group (77.3% increase; P = 0.002; Fig. 4D). Next, EMT-associated cytoskeletal molecules were evaluated. The mRNA expression of α-SMA, vimentin, and TGF-β1 was reduced in the
siCOL8A2 group compared to that in the siControl group (41.9%, 26.3%, and 84.3% decrease; \( P = 0.007, < 0.001, \) and 0.006; Figs. 4E, 4F, 4G).

**Wingless-INT (WNT) Signaling.** The WNT signaling pathway was evaluated alongside EMT-related signaling. GSK3\( \beta \) and \( \beta \)-catenin are involved in WNT signaling.\(^{22} \) WNT signaling was evaluated in cultured hCECs. The expression of GSK3\( \beta \) was unchanged (Fig. 5A) and mRNA expression of \( \beta \)-catenin was decreased in the siCOL8A2 group compared to that in the siControl group (26.3% decrease; \( P < 0.001 \)), which was confirmed by Western blotting (Fig. 5B). The mRNA expression of SLC4A11 was decreased in the siCOL8A2 group compared to that in the siControl group (53.5% decrease; \( P < 0.001 \)), which was confirmed by Western blotting (Fig. 5C). The mRNA expression of Notch1 was increased in the siCOL8A2 group compared to that in the siControl group (307.2% increase; \( P = 0.006 \)), which was also confirmed by Western blotting (Fig. 5D). The mRNA
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FIGURE 2. Functions of corneal endothelial cells after siCOL8A2 transfection. (A) COL8A2 mRNA expression in rat corneal endothelium after COL8A2 siRNA transfection in vivo was reduced (4.73% ± 1.54%). Fold change means relative mRNA expression to control. (B and C) Corneal opacity during the experiments. (D) Alizarin S red staining showed CEC elongation, which was similar with the change in cultured cell shape after siCOL8A2 transfection. Bar: 100 μm. *Statistically significant.

expression of ZO1 and NOX4 was higher in the siCOL8A2 group compared with that in the siControl group (202.4% and 134.5% increase; P < 0.001 for both; Figs. 5E and 5F).

Mitochondrial Oxidative Stress and MitoTracker Red Fluorescence Levels. Mitochondrial oxidative stress levels were evaluated in cultured hCECs using the MitoSOX probe. Increased fluorescence intensity indicated elevated mitochondrial oxidative stress. The increase in mitochondrial oxidative stress levels (25.9% increase; P = 0.004; Figs. 6A and 6B) was confirmed by fluorescence imaging (Fig. 6C). The MitoTracker red fluorescence intensity decreased in the siCOL8A2 group compared to that in the siControl group (56.6% decrease; P < 0.001; Figs. 6D and 6E), which was also confirmed by fluorescence imaging (Fig. 6F).

DISCUSSION

COL8A2 is a major component of DM, the basement membrane of hCECs, and COL8A2 mutations have been reported to be associated with early onset of FECD. In this study, we revealed the role of COL8A2 in hCECs.

In this study, we found that corneal opacity increased in siCOL8A2-transfected rat corneas in vivo. In vivo, rCECs exhibited slender elongation, which was similar to the results of in vitro experiments. COL8A2 is involved in determining the structure and integrity of the cell shape. Mutation of COL8A2 led to biomechanical changes to the DM preceding endothelial cell loss in an early-onset murine model of FECD and inhibition of COL8A2 impaired CEC function. Although silencing the COL8A2 gene did not disrupt the structural integrity of DM, it could change the expression of proteins present and the functions of cells.

CECs are not only a barrier between the corneal matrix and the anterior chamber but also play a role in dehydration of the stroma by pumping out ions. Thus the maintenance of TEER and TEPD is important for hCEC function. Clinically, reductions in TEER and TEPD can result in corneal edema, which can impair vision and require corneal transplantation. This study revealed that transfection with siCOL8A2 reduced TEER and TEPD in cultured hCECs, which represents hCEC function. TEER is a functional parameter for in vitro barrier systems. TEER is the measurement of electrical resistance across a cellular monolayer and is a very sensitive and reliable method to assess the integrity and permeability of the monolayer. TEER reflects the ionic conductance of the paracellular pathway in the epithelial or endothelial monolayer. The TEER of hCECs was reported at 20 to 120 Ω/cm², which is much lower than that of corneal epithelial cells. Furthermore, inhibition of COL8A2 (siCOL8A2) reduced TEER, indicating impaired barrier functions in hCECs. TEPD is a very sensitive index of endothelial transport generated by ionic transport. It is a manifestation of the activity of endothelial fluid pumps that
maintain the cornea at the level of hydration required for transparency. TEPD values have been reported at 1 to 3 mV. Although TEER can be applied to measure barrier function in virtually any type of cultured cell, TEER may be influenced by indeterminate regions of the cell culture outside the edges of the electrodes because it measures impedance of the culture area directly above and between the electrodes. The applied boundary conditions can be identical by applying AC current between electrodes. In this study, the boundary effect was not measured because the current and frequency were not changed. Therefore, there are limitations in measuring TEER.

This study demonstrated that COL8A2 plays a role in cultured hCEC proliferation. siCOL8A2 reduced the proliferation rate, as detected by BrdU, and expression of proliferation-promoting proteins including cyclin D1 and CDK2, and increased the expression of CDKN2A, a proliferation inhibitor. COL8A2 has been suggested to generate a matrix environment that permits or stimulates cell proliferation, and has been reported to modulate the biological effects of TGF-β1 on mesangial cells in the kidney. Lack of collagen VIII in mice ameliorates mesangial cell proliferation and fibrosis. However, our study revealed that COL8A2 inhibition in cultured hCECs suppressed EMT-related changes. COL8A2 is a key extracellular matrix protein associated with cell structure and integrity. siCOL8A2 induced changes in cell shape in vitro and in vivo, which are similarly induced during EMT or mesenchymal-endothelial transition (MET). Thus, EMT-associated proteins were evaluated in cultured hCECs. SNAI1 and pSMAD2/3 expression...
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Figure 4. Endothelial-mesenchymal transition. (A) Cell shape was changed in siCOL8A2. Bar: 100 μm. (B) SNAI1 and pSMAD2 expressions in siControl and siCOL8A2. (C) mRNA expression of SNAI1 was reduced in siCOL8A2 compared to siControl. (D) mRNA expression of SMAD1 was increased in siCOL8A2 compared to siControl. (E, F, and G) mRNA expressions of α-SMA, vimentin, and TGF-β1 in siControl and siCOL8A2. *Statistically significant.

was reduced and SMAD1 expression was increased in the siCOL8A2 group. SNAI1 interacts with SMAD signaling and controls TGF-β1-induced EMT.α,β The α-SMA, vimentin, and β-catenin expression was decreased on siCOL8A2 treatment. The α-SMA, vimentin, and β-catenin are cytoskeletal proteins that are highly expressed during EMT.α,β Vimentin is an intermediate filament of mesenchymal origin.α,β TGF-β1 plays a role in fibrogenesis.α,β...
oxidase that regulates intracellular oxidative stress levels by producing reactive oxygen species (ROS). The major source of intracellular ROS is mitochondria. This study found that COL8A2 modulated mitochondrial function; mitochondrial oxidative stress increased and MitoTracker red intensity decreased, indicating mitochondrial dysfunction. Oxidative stress has been reported to be a cause of CEC disease. ZO1 is a tight junction protein that contributes to the barrier function of the corneal endothelium. During EMT, ZO1 relocates from adhesion membrane complexes to the cytoplasm. In contrast, EMT inhibition causes ZO1 to localize to the cell membrane. ZO1 expression is affected by oxidative stress. Notch1 signaling regulates EMT and inhibition of Notch1 represses EMT. Notch1 signaling is induced by oxidative stress and Notch1 suppresses oxidative stress-induced cell death.
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Pre-existing DM in vivo may not be affected by the temporary knock-down of COL8A2, although long-term knockdown of COL8A2 distorts the structure of the DM by altering protein levels and extracellular matrix (ECM) accumulation. Temporary knock-down of COL8A2 may have more effect on cell shape and barrier function than on the structural integrity of DM. COL8A2 is not known to play a role in the ECM in normal cells. However, further study is necessary to investigate the effect on the ECM using ECM coating.

In conclusion, COL8A2 may contribute to the functions of hCECs, as well as the structural integrity of DM. COL8A2 is essential in maintaining normal CEC morphology and function, and its inhibition may play an essential role in FECD pathogenesis.

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