Excess Transforming Growth Factor-α Changed the Cell Properties of Corneal Epithelium and Stroma

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Purpose. This study is to investigate the corneal anomaly caused by excess transforming growth factor-α (TGF-α) during mouse development.

Methods. Bitransgenic Keraα/β/TGF-α mice, generated via cross-mating tetO-TGF-α and Keraα mice, were induced to overexpress TGF-α by doxycycline commencing at embryonic day 0 or postnatal day 0 to different developmental stages. Bitransgenic mice with doxycycline induction were defined as TGF-αECK mice (TGF-α excess expression by corneal keratocytes). Mouse eyes were examined by hematoxylin and eosin staining, immunofluorescent staining and transmission electron microscopy. Protein and RNA from mouse cornea were subjected to western blotting and real-time quantitative polymerase chain reaction.

Results. In TGF-αECK mice, TGF-α overexpression resulted in corneal opacity. Excess TGF-α initially caused corneal epithelial hyperplasia and subsequent epithelium degeneration as the mouse developed, which was accompanied by gradually diminished K12 expression from the periphery of corneal epithelium and increased K13 expression toward the corneal center. Interestingly, K14 was detected in all layers of corneal epithelium of TGF-αECK mice, whereas it was limited at basal layer of controls. Transmission electron microscopy showed desmosome loss between corneal epithelial cells of TGF-αECK mice. In TGF-αECK mice, keratocan expression was abolished; α-SMA expression was increased while expression of Col1a1, Col1a2, and Col5a1 was diminished. Cell proliferation increased in the corneal epithelium and stroma, but not in the endothelium of TGF-αECK mice.

Conclusions. Excess TGF-α had detrimental effects on corneal morphogenesis during mouse development in that it changed the cell fate of corneal epithelial cells to assume conjunctival phenotypic expression of K13, and keratocytes to myofibroblast phenotype.

Keywords: corneal epithelium, stroma, hyperplasia, TGF-α, transgenic mouse

Transforming growth factor (TGF)-α belongs to the epidermal growth factor (EGF) family. It binds to the EGF receptor (EGFR) to elicit various signaling transduction cascades by which TGF-α is associated with multiple cellular behaviors, such as proliferation, differentiation, and migration of cells during embryonic development and maintenance of homeostasis in adult.1-4 Bidirectional interaction between epithelial cells and stromal mesenchymal cells mediated via growth factors like TGF-α play essential roles in morphogenesis during development and tissue repair.3-5

The biological functions of TGF-α during cancer pathogenesis and normal tissue morphogenesis have been intensively investigated by in vitro and in vivo systems.1,5-9 TGF-α knock-out mice displayed pronounced waviness of whiskers and fur, accompanied by eye abnormalities, including a thinner corneal epithelium. A number of TGF-α knock-out mice were born with partially open eyes which became opaque and inflamed by week 5.5 In contrast, ectopic expression of TGF-α in the mouse lens led to corneal opacities, cataracts, and microphthalmia of the transgenic mouse.10 A milder phenotype was also reported in a similar mouse line when the expression level of TGF-α was not as high in which the corneal epithelium had the normal five to seven cell layers; however, in most transgenic mice overexpressing TGF-α whose corneal epithelium has reduced cell layers around two layers instead of six to seven layers.11

To better elucidate the effects of TGF-α on ocular surface development in a more controlled manner, we developed a doxycycline [Dox] inducible TGF-α over expressing TGF-αECK mouse line (bitransgenic Keraα/β/TGF-α tetO-TGF-α)
mice fed Dox chow), which derived from cross-mating tetO-TGF-α transgenic mice and KerartTA knock-in mice. Our data indicated that excess TGF-α changed cell fate of corneal epithelial cells and stromal keratocytes. TGF-αECK mice posed detrimental effects on cornea morphogenesis during mouse development.

METHODS

Generation and Genotyping of KeraRT/tetO-TGF-α Bitransgenic Mice

Bitransgenic KeraRT/tetO-TGF-α mice were generated via mating transgenic tetO-TGF-α12 and KeraRT knock-in mice.13 All mice were housed at the Animal Facility of the Indiana University. Animal care and use conformed to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research. All animal procedures were approved by the Indiana University Institutional Animal Care and Use Committee.

Transgenic mice were identified by genotyping via polymerase chain reaction (PCR) using oligonucleotide primers specific for each transgene as listed in Supplementary Table S1.

Administration of Dox Chow to Prepare TGF-αECK Mice

To induce TGF-α overexpression, the bitransgenic KeraRT/tetO-TGF-α mice were subjected to systemic induction by Dox chow (1 g/kg: Custom Animal Diets, Bangor, PA) for different periods of time. To prepare fetal TGF-αECK mice at different developmental stages, pregnant dams and nursing mothers were given an intraperitoneal injection of Dox (80 μg/g body weight in PBS, pH 7.4; Clontech Laboratories, Mountain View, CA) at embryonic day 0 (E0) or postnatal day 0 (P0), respectively; then continuously fed Dox chow (ad libitum) until euthanized at embryonic days E14.5 and E18.5 and postnatal days P0, P7, P14, P21, and P30, respectively. Control animals were littersmates containing single transgene, that is, KeraRT and TGF-α (TetO-TGF-α).

Hematoxylin and Eosin Staining and Immunofluorescent Staining

Enucleated eyes were fixed overnight in 4% paraformaldehyde solution in PBS at 4°C, followed by dehydration and paraffin embedding. Deparaffinized and rehydrated tissue sections (5 μm) were stained with hematoxylin and eosin, dehydrated in a graded ethanol series and embedded in Epon 812 epoxy resin. Mouse cornea samples were fixed in 0.1 mol/L cacodylate buffer (pH 7.4) containing 3% glutaraldehyde and 2% paraformaldehyde for 2 hours at 4°C, washed in 0.1 mol/L cacodylate buffer (pH 7.4) containing 3% glutaraldehyde and 1% osmium tetraoxide for 1 hour at 4°C, after fixation in 0.1 mol/L cacodylate buffer (pH 7.4) for 10 minutes three times, and then dehydrated in a graded ethanol series and embedded in Epon 812 epoxy resin. Sections were photographed using a Zeiss microscope equipped with a camera (Axiocam Mrm). For data acquisition, we used the Axiosvision 4.6 software (Carl Zeiss, Jena, Germany).

Western Blotting Analysis

Frozen corneas isolated from experimental mice were homogenized in RIPA buffer (89900 Thermo Fisher Scientific, Waltham, MA) containing 1× protease inhibitor cocktail (P8340, Sigma, St Louis, MO). Tissue lysates (20 μg) from each sample were separated on a 4% to 20% linear gradient Tris-HCl denaturing polyacrylamide Ready Gel (Bio-Rad Laboratories Inc., Hercules, CA) and transferred to PVDF membranes (Whatman, Maidstone, UK). The membranes were incubated with primary antibody overnight at 4°C and then were probed with horse radish peroxidase-conjugated secondary antibody for an hour at room temperature. The signal was detected using an enhanced chemiluminescence assay (Supersignal West Pico, #34080; Thermo Fisher Scientific); then examined and photographed using a VersaDoc 4000MP imaging system (Bio-Rad Laboratories Inc.). Antibodies used in this study were listed in Supplementary Table S2.

Real-Time Quantitative PCR (RT-qPCR)

Total RNA (10 μg) was isolated from mouse corneal tissues using Trizol reagent (Invitrogen, Carlsbad, CA), then annealed to random primers and reverse transcribed with avian reverse transcriptase (RT) kits (Promega, Madison, WI), according to the manufacturer’s instructions. RT-qPCR was performed using the CFX96 real-time system equipped with a C1000 Thermal Cycler (Bio-Rad Laboratories Inc.). After the initial 3-minute denaturing step at 95°C, 40 subsequent cycles at 95°C lasting 15 seconds, 62°C for 15 seconds, and 72°C for 30 seconds were performed. The cycle threshold values were used to calculate the normalized expression of genes of interest against Gapdh using Q-Gene software. Primer pairs were listed in Supplementary Table S1.

Transmission Electron Microscopy

To conditionally overexpress TGF-α in cornea stromal cells, KeraRT mice were crossed with transgenic mouse line tetO-TGF-α to create the double transgenic mouse KeraRT/TGF-α

RESULTS

TGF-α Was Overexpressed in the Corneal Keratocytes of Dox-Induced KeraRT/TGF-α Mice

To conditionally overexpress TGF-α in cornea stromal cells, KeraRT mice were crossed with transgenic mouse line tetO-TGF-α to create the double transgenic mouse KeraRT/TGF-α.
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The Phenotype of TGF-α<sup>ECK</sup> Transgenic Mice

The TGF-α<sup>ECK</sup> mice treated by Dox administration from E0 till P21 displayed corneal opacities (Figs. 1C, 1D). This phenotype was similar to that observed previously in TGF-α<sup>ECK</sup> mice. To further investigate and characterize the phenotypical changes in the cornea of TGF-α<sup>ECK</sup> mice, Dox induction was performed from E0 until various developmental time points including E14.5, P0, P7, P14, and P21. At each time point, experimental mice were euthanized to collect eyeballs for analysis. Hematoxylin and eosin staining showed that central corneal epithelium in the littermate controls consisted of one or two cell layers at E14.5, P0, and P7 (Figs. 2A, 2C, 2E). Then corneal epithelium had three to four cell layers at P14 (Fig. 2G) and five to six layers at P21 owing to epithelial cell stratification (Fig. 2I). In contrast, TGF-α overexpression resulted in three to four epithelial cell layers at E14.5 (Fig. 2B) and more than 10 cell layers formed by P0 (Fig. 2D). Interestingly, corneal epithelium then gradually showed signs of degeneration and a progressive decrease in cell layers starting at P7 (Figs. 2F, 2H, 2J), which was characterized by the presence of intercellular cavities within the corneal epithelium (Figs. 2D, 2F, 2H) of the TGF-α<sup>ECK</sup> mice with prolonged Dox induction. In addition, the cell density of corneal stromal cells in littermate controls increased up to P0, and then decreased after that time point until P14; however, overexpression of TGF-α increased corneal stromal cells at E14.5 and remained high after P14 (Fig. 2K).

Excess TGF-α Altered Cell Phenotype of Corneal Epithelial Cells

Considering TGF-α as a potent mitogen, the EGFR signaling pathway triggered by TGF-α have profound effects on cell differentiation of corneal epithelium and stroma. To this end, the expression of corneal epithelial differentiation markers was analyzed by immunofluorescent staining. K12, a specific corneal epithelium differentiation marker, was expressed in corneal epithelium after E14.5 in littermate controls (Figs. 3A, 3C, 3E, 3G). Interestingly, K12 expression (green) was detected at E14.5 in the corneal epithelium of TGF-α<sup>ECK</sup> mice (Fig. 3B), but its expression was caused upregulation of EGFR expression in corneal epithelium after E14.5 in littermate controls (Figs. 3A, 3C, 3E, 3G). Furthermore, TGF-α caused upregulation of EGFR expression in corneal epithelium of both littermate controls and TGF-α<sup>ECK</sup> mice at all developing stages examined as shown above (Supplementary Fig. S1A–S1H).
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FIGURE 2. Overexpression of TGF-α in corneal keratocytes resulted in changed phenotypes in corneal epithelium and stroma. (A–J) Histologic hematoxylin and eosin staining of central cornea sections of TGF-αECK mice (B, D, F, H, J) and littermate controls (A, C, E, G, I) with Dox induction at different stage. Excess TGF-α caused corneal epithelial hyperplasia (B, D) followed by degeneration (F, H, J). (K) Statistical analysis of the corneal stromal cell numbers presented in the central corneas under each microscopic field of view (*n = 6). The cell density of corneal keratocytes in littermate controls increased at around P0 and then decreased later after that until P14; however, overexpression of TGF-α increased corneal stromal cells at E14.5 and did not decrease after that. En, endothelium; Epi, epithelium; Le, lens; lid, eyelid; str, stroma.

Taken together, our data indicated that excess TGF-α altered proliferation and differentiation during embryonic corneal morphogenesis.

Excess TGF-α Altered the Ultrastructure of Corneal Epithelium

Next, we took a closer look at the ultrastructure of corneal epithelium with transmission electron microscopy. The littermate control mice had one to two corneal epithelial cell layers at E14.5, E18.5, and P7 (Figs. 5A, 5C, 5E). In contrast, there were multiple corneal epithelial cell layers in TGF-αECK mice at the same developmental stages, respectively (Figs. 5B, 5D, 5F). Furthermore, transmission electron microscopy showed that desmosomes between corneal epithelial cells were first observed in the TGF-αECK corneal epithelium at E18.5 (Fig. 5D). Although desmosomes were present at P7 in corneal epithelium of littermate controls (Fig. 5E), they were absent in the superficial layers of the TGF-αECK corneal epithelium at P7 (Fig. 5F). This dynamic change indicated that TGF-α initially promoted desmosome formation in the corneal epithelium, but it caused a loss of desmosomes at later developing stages. Accompanying desmosomes reduction; condensed and fragmented chromatins (arrowhead in Fig. 5F) were observed in some of these superficial cells. Furthermore, gaps were present between cells (asterisks in Fig. 5F). Taken together, these data suggested that excess TGF-α initially caused corneal epithelium hyperplasia, but subsequent progressive degeneration possibly owing to cells undergoing apoptosis.

Excess TGF-α Caused Phenotypic Changes of Corneal Keratocytes

The stromal extracellular matrix is critical to corneal function by establishing the appropriate mechanical stability required for maintaining corneal shape and curvature. To examine the effects of excess TGF-α on the corneal keratocytes, keratocan (Kera) expression was examined by immunofluorescent staining. To our surprise, Kera expression was decreased significantly at E18.5 and totally abolished at P0 (Figs. 6B, 6D, 6F, 6H), whereas strong Kera expression was maintained in the corneal stroma of littermate controls (Figs. 6A, 6C, 6E, 6G). Moreover, RT-qPCR data showed that expression of Col1a1, Col1a2, and Col5a1 dramatically decreased in TGF-αECK stroma when compared with that in littermate controls (Figs. 7A, 7B, 7C). However, the expression of alpha-smooth muscle actin (α-SMA) was significantly increased in Dox-induced TGF-αECK stroma, as revealed by RT-qPCR and Western blotting (Figs. 7D, 7E). These data suggested that changes of keratan sulfate proteoglycans and extracellular matrix assembly might have some influence on the corneal transparency and lead to corneal haze as observed in TGF-αECK mice.

TGF-α Promoted Corneal Epithelium and Stromal Cell Proliferation

Multiple cell layers in the corneal epithelium and higher stromal cell density implicated that cell proliferation was upregulated in the Dox-induced TGF-αECK mice. To address this hypothesis, the expression of proliferating cell nuclear
FIGURE 3. Expression patterns of Krt12 and Krt13 were altered in TGF-α ECK mice. (A, B) Immunofluorescent staining showed that TGF-α overexpression promoted K12 expression (green) in the cornea at E14.5. (C–H) K12 expression was gradually lost from the peripheral corneal regions at P0 and P7 (D, F) and almost completely absent at P14 (H). By contrast, Krt13 expression (red) was progressively extended to the central cornea in KeraRT/TGF-α mice (D, F) and almost completely covered the cornea at P14 (H). Nuclei were counterstained with DAPI (blue). En, endothelium; Epi, epithelium; Le, lens; lid, eyelid; Re, retina; str, stroma.

FIGURE 4. Krt14 expression in corneal epithelium was changed after overexpression of TGF-α. (A–H) Immunofluorescent-stained of Krt14 (red) in the littermate controls and TGF-α ECK mice were administrated Dox from E0 to P7. Krt14 expression existed only in the basal layer of corneal epithelium in the control mice (A, C, E, G), but its expression appeared in all the layer of corneal epithelium of TGF-α ECK mice. The nuclei were counterstained with DAPI (blue). En, endothelium; Epi, epithelium; Le, lens; lid, eyelid; Re, retina; str, stroma.
FIGURE 5. The ultrastructure of corneal epithelium of littermate controls and TGF-αECK mice at different developmental time points. Corneal epithelium of littermate controls had one to two cell layers at E14.5, E18.5, and P0 (A, C, E). Desmosomes between cells were observed at P7 (E). By contrast, corneal epithelium of TGF-αECK mice had multiple cell layers at all the checked time points (B, D, F). Desmosomes were clearly and densely present in corneal epithelium at E18.5 (D). However, corneal epithelial cells lost desmosomes in the superficial layer at P7 (F). Note that dark nuclei and condensed fragmented chromatin (arrowhead) were evident in some superficial cells. Furthermore, gaps were present between epithelial cells (‘ in F). BM, basal membrane; Epi, epithelium; N, nuclei; str, stroma.

FIGURE 6. Kera expression was diminished in TGF-αECK mice. (A–F) Immunofluorescent-stained micrographs of Kera (red) showed that no positive signals were detected in KeraRT/TGF-α mice after Dox induction. (B, D, F), whereas strong signals were observed in corneal stroma of littermate controls (A, C, E). Cell nuclei were counterstained with DAPI (blue). En, endothelium; Epi, epithelium; Le, lens; lid, eyelid; Re, retina; str, stroma.

antigen (PCNA) was examined by immunofluorescent staining in mice at P0 (Dox induction beginning at E0). It showed that few of PCNA+ cells were detected in corneal epithelium, stroma, and endothelium of littermate controls at P0 (Figs. 8A, 8C). However, PCNA+ cells were plentifully present in the TGF-αECK corneal epithelium and stroma (Figs. 8B, 8D). PCNA+ cells in TGF-αECK endothelium showed no change in comparison with controls (Fig. 8B vs Figs. 8A, 8D vs Fig. 8C). Statistical analysis of PCNA immunofluorescent immunostaining data indicated that cell proliferation was increased in corneal epithelial and stromal cells of TGF-αECK mice at this developmental stage. However, endothelial cell proliferation did not change in Dox-induced TGF-αECK mice (Fig. 8E). These data indicated that TGF-α was a potent stimulator of corneal epithelial cells and stromal cells during mouse development.
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**FIGURE 7.** Corneal stromal extracellular matrix components and α-SMA expression were changed in TGF-αECK mice after Dox induction from E0 to P30. (A–C) Real-time PCR analysis showed that expression of corneal stroma extracellular matrix components including collagen type 1 α1 (A), type 1 α2 (B), and type 5 α1 (C) decreased in corneal stroma of TGF-αECK mice administered Dox from E0 to P30. (D) Real-time PCR analysis indicated that expression of α-SMA was upregulated in corneal stroma of TGF-αECK mice administered Dox from P0 to P30. Likewise, (E) Western blot results showed that α-SMA protein also increased in the stroma when compared with littermate controls.

**FIGURE 8.** TGF-α overexpression increased cell proliferation in corneal epithelium and stroma. (A–D) Immunofluorescent micrographs of the corneas probed with anti-PCNA showing a few positive PCNA (PCNA+) cells scattered at the basal corneal epithelial cells of littermate controls. In contrast, abundant PCNA+ corneal epithelial cells were present in TGF-αECK mice administrated Dox from E0 to P0 (A vs B, C vs D). Similarly, PCNA+ stromal cells were enhanced in KeraRT/TGF-α mice when compared with the littermate controls. The white and green dashed lines in each panel delineate the corneal epithelial layer and endothelial cells, respectively. Nuclei were counterstained with DAPI (blue). (E) Statistical analysis of immunofluorescent staining results showed that the percentage of PCNA positive cells in epithelium, stroma, and endothelium of littermate controls and TGF-αECK mice (n = 4). En, endothelium; Epi, epithelium; Le, lens; Str, stroma.

**DISCUSSION**

In the present study, we found that excess TGF-α produced from corneal stromal cells had a profound influence on corneal development in the TGF-αECK mice. Our data indicated that TGF-α initially promoted corneal epithelial cell hyperplasia, followed by degeneration; which was accompanied by the loss of K12 expression, whereas ectopic expression of K13 resembling conjunctivalization of corneal epithelium after prolonged Dox induction, a phenotype...
characteristic of limbal stem cells deficiency.\textsuperscript{26} Interestingly, excess TGF-α caused phenotypic changes of keratocyte, such as loss of Kera expression and downregulation of extracellular matrix components, and upregulation of α-SMA, a maker of myofibroblasts. All these defects during corneal development resulted in corneal opacities in TGF-α\textsuperscript{ECK} mice. Therefore, overexpression of TGF-α in keratocytes was detrimental to normal corneal morphogenesis.

The overexpression of TGF-α specifically in corneal stroma of TGF-α\textsuperscript{ECK} mice initially increased corneal cell proliferation both in the stroma and the epithelium. This phenomenon was expected because TGF-α is a potent mitogen for epithelial cells and fibroblasts and can affect cell behavior in an autocrine, paracrine, or even juxtacrine manner.\textsuperscript{5,6} However, we noticed that corneal endothelium in the TGF-α\textsuperscript{ECK} mice were not perturbed by excess TGF-α. This finding is not consistent with previous observations; Reneker et al.\textsuperscript{10,11} reported that the overexpression of TGF-α in the ocular lens via the mouse αA-crystallin promoter caused no endothelium formation and multiple anterior segment defects in the transgenic mice. The αA-crystallin gene was expressed in mouse lens starting at E12.5 before the formation of mouse corneal endothelium at E14.5 to E15.5,\textsuperscript{18} whereas Kera gene expression in the corneal stroma commences at E14.5.\textsuperscript{19} Therefore, different mouse drivers activated TGF-α overexpression at different developmental stages may account for the variation of phenotypes observed in these mouse strains.

One interesting observation in the present studies was that excess TGF-α driven by TGF-α\textsuperscript{ECK} led to corneal epithelial hyperplasia that was followed by subsequent epithelium degeneration (Fig. 2), likely via apoptosis (shown in Fig. 5), which was accompanied by the conjunctivalization of cornea epithelium (Fig. 3). This observation was unique and distinct from phenotypic changes found in other transgenic mouse lines that overexpress TGF-α and fibroblast growth factor-7 in eyelid stroma and corneal epithelial cells, respectively.\textsuperscript{20,21} It has been reported that overexpression of EGF and/or EGFR caused apoptosis of cultured epithelial cells in vitro.\textsuperscript{22,23} Thus, it is likely that excess TGF-α promoted corneal epithelial cell proliferation during embryonic development. However, it might lead to apoptosis at postnatal corneal endothelium under pathologic conditions owing to dysregulation of the cell cycle.\textsuperscript{24,25} The conjunctivalization could be in part explained by the rapid turnover rate of corneal epithelial cells under the influence of excess TGF-α, which might exhaust the limbal stem cells resembling the phenomenon observed in limbal deficiency.\textsuperscript{26,27} This idea will be investigated using our new driver mouse strain, Krt4\textsuperscript{rtTA} in a future study in which the rtTA minigene was knocked into the mouse Krt4 locus (Zhang J, et al. IOVS 2016;57:ARVO E-Abstract 12).

Another interesting finding in this study was that Kera expression diminished at P0 in the corneal stroma of TGF-α\textsuperscript{ECK} mice with Dox induction from E0 (Fig. 6). This finding implied that de novo rtTA synthesis might no longer take place in the stroma of TGF-α\textsuperscript{ECK} mice at P0; however, rtTA protein might persist and maintain TGF-α expression, alternatively, which would lead to the termination of TGF-α overexpression, but the influence of excess TGF-α from E0 through P0 might reach a point of no return and irreversibly alter the cell fate of corneal epithelial cells and stromal cells of TGF-α\textsuperscript{ECK} mice.

Kera is one of the important keratan sulfate proteoglycans uniquely expressed in the corneal stroma and plays a crucial role in the development and maintenance of corneal transparency\textsuperscript{20}; therefore, the loss of Kera expression affects corneal transparency. Extracellular matrix assembly have also been implicated in corneal transparency.\textsuperscript{29–31} Collagen I and collagen V are two fibril-forming collagens synthesized by keratocytes,\textsuperscript{32,33} and our data indicated that both types of collagen decreased significantly (Figs. 7A–7C), which might contribute to the corneal haze. In contrast, our RT-qPCR and Western blot results showed that α-SMA expression was boosted in corneal stroma of TGF-α\textsuperscript{ECK} mice (Figs. 7D, 7E). Upregulation of α-SMA may be triggered by inflammatory cytokines released from corneal inflammatory cells owing to excess TGF-α in the cornea. In the double transgenic mouse strain, K12\textsubscript{rtTA}/TGF-α, corneal inflammation was reported after TGF-α overexpression in epithelial cells (Kao WW-Y, et al. IOVS 2011;52: ARVO E-Abstract 309). In the same mouse model, increased expression of Wnt5a was also found after excess TGF-α was produced in the cornea.\textsuperscript{14} Wnt5a is an inflammatory mediator that promotes inflammatory response.\textsuperscript{33} Thus, in our current TGF-α\textsuperscript{ECK} mice, TGF-α overexpression may indirectly result in corneal inflammation, which leads to the expression of α-SMA. α-SMA was known as one of the hallmarks of myofibroblast, and the presence of α-SMA may imply that some corneal stromal cells might undergo transformation. The exact mechanism of this transformation caused by TGF-α will be investigated in future studies.

In our current study, we used the novel knock-in driver mouse line Kera\textsuperscript{rtTA} to express TGF-α in corneal keratocytes, Kera\textsuperscript{rtTA} mouse line displayed strong rtTA expression in Kera expressing cells.\textsuperscript{13} Previously, we generated a transgenic mouse line called KR by randomly inserting the Kera\textsuperscript{promoter-rtTA} minigene into the mouse genome, in which expression of rtTA was driven by mouse Kera promoter region.\textsuperscript{34} The double transgenic mouse KR/TGF-α was created to express TGF-α in corneal stromal cells with Dox administration. The eyes of Dox-induced KR/TGF-α mice looked normal and transparent;\textsuperscript{29} no dramatic phenotypes were observed. This inconsistent consequence of ectopic expression of TGF-α in cornea of these two mouse strains is probably owing to different expression levels of TGF-α in their corneal stroma, which suggested that lower TGF-α level was produced in KR/TGF-α mouse strain. Reneker et al.\textsuperscript{16,17} also reported that different level of TGF-α expression caused variable phenotypes in their transgenic mouse lines. Based on our experiences, Kera\textsuperscript{rtTA} knock-in mouse line (driver gene activity mainly in the corneal stroma) was more effective and faithful in driving transgene expression mimicking Kera expression patterns limited in corneal stroma of TGF-α\textsuperscript{ECK} mice, whereas transgenic mouse line KR (driver gene activity mainly in eyelid stroma) is to modify gene expression in mouse eyelid stroma.

**Conclusions**

Results of our studies indicated that excess TGF-α synthesized by corneal stromal cells led to conjunctivalization of corneal epithelium expressing K13 resembling limbal stem cells deficiency and transition of keratocytes to assume myofibroblastic cell fate, for example, enhanced α-SMA expression.
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