Bromfenac Inhibits TGF-β1–Induced Fibrotic Effects in Human Pterygium and Conjunctival Fibroblasts

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Purpose. Nonsteroidal anti-inflammatory drugs (NSAIDs) have shown antifibrotic effects on several diseases. The aims of the present in vitro study were to investigate the antifibrotic effects of bromfenac (a kind of NSAID) on primary human pterygium fibroblasts (HPFs) and primary human conjunctival fibroblasts (HConFs), as well as to explore the possible mechanisms of these effects.

Methods. The cells used in this study were primary HPFs and HConFs, and profibrotic activation was induced by transforming growth factor-beta1 (TGF-β1). Western blot, quantitative real-time PCR, and immunofluorescence (IF) assays were used to detect the effects of TGF-β1 and bromfenac on the synthesis of fibronectin (FN), type III collagen (COL3), and alpha-smooth muscle actin (α-SMA) in HPFs and HConFs; the changes of signaling pathways were detected by Western blot; cell migration ability was detected by wound healing assay; cell proliferation ability was detected by CCK-8 assay; and pharmaceutical inhibitions of the downstream signaling pathways of TGF-β1 were used to assess their possible associations with the effects of bromfenac.

Results. Bromfenac suppressed the TGF-β1–induced protein expression of FN (0.59 ± 0.07 folds, P < 0.001), COL3 (0.48 ± 0.08 folds, P = 0.001), and α-SMA (0.61 ± 0.03 folds, P = 0.008) in HPFs. Bromfenac also attenuated TGF-β1–induced cell migration (0.30 ± 0.07 folds, P < 0.001), cell proliferation (0.64 ± 0.03 folds, P = 0.002) and the expression levels of p-AKT (0.66 ± 0.08 folds, P = 0.032), p-ERK1/2 (0.69 ± 0.11 folds, P = 0.003), and p-GSK-3β-S9 (0.65 ± 0.10 folds, P = 0.002) in HPFs. PI3K/AKT inhibitor (wortmannin) and MEK/ERK inhibitor (U0126) reduced the TGF-β1–induced synthesis of FN, COL3, and α-SMA in HPFs. All the results were similar in HConFs.

Conclusions. Bromfenac protects against TGF-β1–induced synthesis of FN, α-SMA, and COL3 in HPFs and HConFs at least in part by inactivating the AKT and ERK pathways.

Keywords: pterygium, fibrosis, fibroblast, NSAIDs, Bromfenac, TGF-β1, AKT, ERK
bind to their receptors, which activate multiple intracellular signal-transmitting proteins, including Smads, PI3K/AKT, MEK/ERK, RHO-A, and JNK/P38. In the present study, TGF-β1 was used to induce fibrosis in human pterygium fibroblasts (HPFs) and human conjunctival fibroblasts (HConFs).

Nonsteroidal anti-inflammatory drugs (NSAIDs) have shown antifibrotic properties in diseases such as breast cancer and skin carcinoma. The main target of NSAIDs is cyclooxygenase-2 (COX-2). It has been reported that pterygial specimens clearly express COX-2, whereas normal conjunctiva specimens have no COX-2 expression. As far as we know, only one study has reported that the NSAID called nimesulide could suppress the proliferation of HPFs. NSAIDs have been used during the perioperative period of pterygium for its analgesic and anti-inflammatory effects, but the potential preventive effects of NSAIDs on pterygium recurrence remain unclear. Bromfenac is a kind of NSAID, which is approximately three to four times more effective at inhibiting COX-2 in vitro as compared with other ophthalmic NSAIDs amfenac, ketorolac, and diclofenac. Bromfenac is usually used to prevent macular edema and inflammation after cataract and complex ocular surgeries, which is less toxic toward corneal epithelial cells. Our study intended to explore whether bromfenac has the potential to prevent the excessive wound healing that takes place after the pterygium surgeries.

The aims of the present study were to investigate the antifibrotic effects of bromfenac on HPFs and HConFs, as well as to explore the possible mechanisms involved in these effects.

**METHODS**

**Isolation and Cultivation of Primary HPFs, HConFs, and Human Conjunctival Epithelial Cells (HConEs)**

Twenty-one different pterygium samples were obtained from 21 patients with mean age of 61.7 ± 7.4 years (mean ± SD; range, 49–76 years) who underwent pterygium surgical excision at our hospital. Parasatal pterygium tissues were obtained at the time of excision; the heads and bodies of the pterygium samples were used for cultures. All of the pterygium specimens were classified as grade T3 (fleshy) according to the grade T classification. The fleshiness of stroma is significantly associated with pterygium recurrence after excision. The heads of these pterygium tissues invaded over the corneal limbus for more than 3 mm. All patients were treated in accordance with the Declaration of Helsinki, and each of them provided informed written consent. This study protocol was approved by the institutional review board of our institute. The diagnosis of primary pterygium was entirely clinical and without pathological evidence. We excluded patients with recurrent pterygium, a history of surgery on the studied eyes, systemic contagious diseases, diabetes mellitus, deficiencies of the immune system, or other ocular diseases except for cataract.

Ten different conjunctiva samples were obtained from 10 donors of the eye bank of Second Affiliated Hospital, Zhejiang University School of Medicine (Hangzhou, China). The mean age of the 10 donors was 56.7 ± 6.9 years (mean ± SD; range, 43–66 years), and the normal conjunctiva tissues of the donors were taken to be cultivated.

After the surgeries, the pterygium and conjunctiva samples were put into culture medium (Advanced Dulbecco’s Modified Eagle Medium/Ham’s F-12 [DMEM/F-12]; Gibco Life Technologies, Karlsruhe, Germany) supplemented with 10% fetal bovine serum (FBS; Gibco Life Technologies), 100 U/mL penicillin, and 100 g/mL streptomycin (Gibco Life Technologies), and then transported on ice for less than 2 hours before processing. On a bacteria-free workbench, the samples were washed with PBS (Gibco Life Technologies) three times, then cut into small pieces (approximately 1 mm × 2 mm), and incubated in trypsin-EDTA (0.25%, Gibco Life Technologies) at 37°C for 15 minutes. Then, the solution was diluted with culture medium, filtered with 70-μm cell strainers (BD Falcon, Franklin Lakes, NJ), and centrifuged at 200g for 5 minutes. Next, the sediment was resuspended with culture medium, inoculated into 12-well cell culture plates, and incubated in a humidified atmosphere of 5% CO2 at 37°C. The medium was changed every 2 days thereafter.

Not all of the specimens were able to be isolated and to cultivate primary cells successfully, we cultivated fibroblasts from 21 pterygium and 10 conjunctiva samples, and the characteristics of the patients and donors are summarized in Supplementary Table S1. The cells were grown as communities, cells in a well usually present only one of the two types of morphologies, fusiform or multilateral. If all of the cells in a well presented an elongated and fusiform morphology, the cells were amplified and prepared to be identified as HPFs or HConFs. As long as there were cells presenting typical epithelial multilateral morphology mixed into the elongated and fusiform cells, the corresponding wells were abandoned. If all of the cells in a well presented a typical epithelial multilateral morphology, the cells were also amplified and prepared to be identified as epithelial cells. After the authentications that the fibroblasts were not contaminated with epithelial cells by means of Western blot, immunofluorescence staining assays, and morphology, the fibroblasts were amplified for experiments. Each person’s cells were not mixed with other people’s cells. The HPFs and HConFs were passaged at the ratio of 1:2, cells after the third generation were used for all experiments. The information of which cells were used in each experiment were summarized and tied into the demographics data in Supplementary Table S1.

**Cultivation of Human Foreskin Fibroblasts (HFFs) and HeLa Cells**

HFFs (provided by Stem Cell Bank, Chinese Academy of Science) were used as positive control (standard fibroblasts), and HeLa cells were used as negative control (standard epithelial cells). HFFs were cultivated in DMEM/F-12 containing 20% FBS, 100 U/mL penicillin, and 100 g/mL streptomycin. HeLa cells were cultured in high glucose DMEM (4.5 g/L; Gibco Life Technologies) with 10% FBS, 100 U/mL penicillin, and 100 g/mL streptomycin. All of the cells were maintained in a humidified atmosphere of 5% CO2 at 37°C.

**Drug Treatments of Cells**

As mentioned above, the cells were incubated with DMEM/F-12 supplemented with 10% FBS before treatments, then the medium was changed to DMEM/F-12 containing 1% FBS to conduct the experiments. But in the wound healing assays, serum-free DMEM/F-12 was used. Bromfenac was used at nontoxic concentrations. To assess the antifibrotic effects of bromfenac on TGF-β1-induced cells, the cultured HPFs and HConFs were divided into the following groups and treated for 48 hours: the control group; the TGF-β1 group, in which the cells were exposed to 20 ng/mL TGF-β1 (200-1; PeproTech, Rocky Hill, NJ, USA); the bromfenac group, in which the cells were treated with 90 μg/mL bromfenac (bromfenac active components were from Senju Pharmaceutical, Osaka, Japan); the TGF-β1 + bromfenac group, in which the cells were treated with 20 ng/mL TGF-β1 plus 90 μg/mL bromfenac. For the dose-
effect relationship study, HPFs and HConFs were treated with three concentrations of bromfenac (30, 60, and 90 μg/mL) for 48 hours. For the inhibitors studies, 200 nM wortmannin (Selleck, Houston, TX, USA) (PI3K inhibitor) and 20 μM U0126 (Selleck, CST, Waltham, MA, USA) (MEK inhibitor) were added 2 hours before the TGF-β1 treatment, and the HPFs and HConFs were collected 48 hours after TGF-β1 treatment to evaluate the changes. TGF-β1 was dissolved in DMEM/F-12; bromfenac was dissolved in PBS, then filtered to eliminate the microorganisms; wortmannin and U0126 were dissolved in dimethyl sulfoxide (DMSO). The same volume of solvents DMEM/F-12, PBS, or DMSO was used as controls. The medium was changed every 24 hours to replenish the chemicals.

Western Blot Assay

After treatments, cell total proteins were extracted from the cells with the cell total protein extraction kit (Sangon, Shanghai, China), which contains lysis buffer (NaCl, SD8138 [1M Tris-buffer solution pH 7.4], 0.5M EDTA, 2Na, NDSB-201, and NP-40), phenylmethanesulfonyl fluoride, protease inhibitor, and phosphatase inhibitor. The total protein concentrations were determined by the bicinchoninic acid Protein Assay Kit (B9643–1L; Sigma-Aldrich, St. Louis, MO, USA). The same amounts of proteins were subjected to SDS-PAGE with certain concentrations, then transferred to a polyvinylidene fluoride (PVDF) membrane (Millipore, Bedford, MA, USA), and blocked for 2 hours with 5% BSA in Tris-buffered saline (TBS-T, 10 mM Tris-HCl [pH 7.5], 100 mM NaCl, and 0.1% Tween 20). The primary antibodies were diluted in TBS-T containing 5% BSA. The information of the antibodies used in our study is presented in Supplementary Table S2. The PVDF membranes were then incubated in the primary antibody diluents overnight at 4°C. After that, the PVDF membranes were washed and incubated with horseradish peroxidase-conjugated secondary antibodies for 1 hour at room temperature. Immunodetection was performed with the enhanced chemiluminescence Western Blot Detection System (Millipore). The immunoblots were visualized by the ChemiDoc XRS System (Bio-Rad, Hercules, CA, USA). The intensities of the bands were quantified by densitometry; the software Image J (http://image.nih.gov/ij/; provided in the public domain by the National Institutes of Health, Bethesda, MD, USA) was used. GAPDH served as an internal control.

Quantitative Real-Time PCR

HPFs and HConFs having 80% confluency in a 6 cm-dish were treated and harvested, and the total RNA was extracted (Trizol Reagent; Invitrogen, CA, USA). RNA concentrations (200–500 ng/μl) and OD260/OD280 ratios (1.9–2.0) were determined using a Nanodrop 1000 UV spectrophotometer (Thermo Scientific, Waltham, MA, USA); approximately 1000 ng RNA was used per reverse transcription reaction (PrimeScriptRT Reagent Kit; Takara, Shiga, Japan). The SYBR Premix Ex Taq II Kit (TakaRa) was used for quantitative real-time PCR on a 7500 Fast Real-Time PCR System (ABI, Loma Linda, CA, USA), with a 2-step method. The primers are shown in Supplementary Table S3. The expression levels of the target genes were normalized to that of GAPDH and were calculated using the 2−ΔΔCT method. The data are presented as fold-changes.

Immunofluorescence (IF) Staining Assay

HPFs and HConFs were seeded on the cleaned coverslips for 24 hours. Afterward, the medium was changed to DMEM/F-12 with 1% FBS and the indicated drugs, respectively. After being treated for 48 hours, the cells were rinsed with PBS three times and then fixed in 4% formaldehyde solution for 20 minutes. After being rinsed with PBS three times again, the cells were permeabilized with 0.5% Triton X-100 (Sigma-Aldrich, St. Louis, MO, USA) for 15 minutes, and blocked with 10% normal goat serum for 1 hour at room temperature. Subsequently, the cells were incubated with primary antibodies against α-SMA, COL3, pan-keratin, vimentin, or corresponding isotype control overnight at 4°C. Then, after being rinsed with PBS, Alexa Fluor 555 Donkey Anti-Rabbit immunoglobulin (IgG) or Alexa Fluor 555 Donkey Anti-Mouse IgG solution was used to incubate the cells for 2 hours at room temperature. The nuclei were stained with DAPI (4',6'-diamidino-2-phenylindole) contained in the mounting medium. Eventually, the coverslips were mounted on the slides with a drop of mounting medium with DAPI (H-1200; Vector, Burlingame, CA, USA). Fluorescent images were then observed using a Leica inverted fluorescence microscope (DMI8/DFC 550; Leica, Wetzlar, Germany).

Wound Healing Assay

HPFs and HConFs were seeded in 12-well plates. After being cultured for 24 hours, the cell density of each well reached 90% confluency, and then the cells were scratched with a sterile 100-μL pipette tip. The scratched wells were then washed with PBS three times, and the cells were treated with the indicated drugs in serum-free medium. The wounds were photographed at 0 and 24 hours. The cellular migration area in each image was measured using Image J software. The data were quantified based on the area of the wounds at 0 hour, which was considered as 100%.

CCK-8 Cell Proliferation Assay

Cell proliferation ability was measured by Cell Counting Kit 8 (CCK-8; Dojindo, Japan). About 5 × 10^3 cells/well were seeded in a 96-well plate and cultivated for 24 hours, then treated with indicated drugs in medium free of FBS for 48 hours, respectively. Then, CCK-8 reagent (10 μl) was added to each well and incubated for 4h. Finally, the absorbance value of each well at a wavelength of 450 nm (OD450) was determined using a microplate reader.

Statistical Analysis

At least three independent experiments were performed at least in triplicate for each assay. The cells of an independent experiment were from the same batch of the same person. Different independent experiments used different batches of cells. The results of multiple experiments are presented as the mean ± SD. After confirming that all variables were normally distributed, the Kolmogorov-Smirnov test and visualized by Q-Q plot, statistical differences were determined by 1-way ANOVA followed by Bonferroni’s multiple comparison test for comparison among three or more groups. A P value < 0.05 was defined as statistically significant. All data were analyzed using SPSS (version 18.0; IBM Corp, Chicago, IL, USA) statistical software. The statistics are presented using GraphPad Prism 5.

RESULTS

Cultivation and Identification of Primary HPFs and HConFs

Primary HPFs (n = 21) and HConFs (n = 10) were isolated from human eyes. These cells were expanded in culture plates and their identities were confirmed using Western blotting and indirect IF microscopy for mesenchymal proteins FN, COL3,
vimentin, and absence of the expression of epithelial keratins using the pan-keratin antibody (total keratin 4, 5, 6, 8, 10, 13, and 18). Previous studies have reported the positive expression of pan-keratin in primary human pterygium epithelial cells. HFFs and HeLa cells were used as positive and negative controls, respectively. As a result, HFFs, HPFs, and HConFs clearly expressed FN and COL3, whereas HeLa cells had negative expression of FN and COL3. HFFs, HPFs, and HConFs expressed higher protein levels of vimentin than HeLa cells. HeLa cells expressed high protein level of pan-keratin, whereas HFFs, HPFs, and HConFs showed negative protein expression of pan-keratin (Supplementary Fig. S1B).

Cell phenotypes were confirmed by light microscopy. HFFs, HPFs, and HConFs presented elongated and fusiform morphology, whereas primary HConFs and HeLa cells had typical multilateral epithelial cell morphology (Supplementary Fig. S1B).

**The Effects of Bromfenac on TGF-β1–Induced ECM Synthesis and Myofibroblast Activation in HConFs and HPFs**

HPFs from three patients and HConFs from three donors were treated with TGF-β1 (20 ng/mL) and bromfenac (90 μg/mL) for 48 hours. The effects of bromfenac and TGF-β1 on the synthesis of FN, COL3, and α-SMA were tested by Western blotting. In HPFs, after the treatment of TGF-β1, the protein levels of FN (2.62 ± 0.34 folds, P < 0.001), COL3 (1.98 ± 0.27 folds, P = 0.002), and α-SMA (2.32 ± 0.34 folds, P < 0.001) increased significantly, as compared with the control group; TGF-β1 showed similar profibrotic effects on HConFs (Fig. 1A). Besides, the protein levels of FN, COL3, and α-SMA in HConFs and HPFs increased with time after exposure to TGF-β1 (Supplementary Fig. S2A).

Meanwhile, in HPFs, TGF-β1 + bromfenac group expressed lower protein levels of FN (0.59 ± 0.07 folds, P = 0.008), COL3 (0.48 ± 0.08 folds, P = 0.001), and α-SMA (0.61 ± 0.03 folds, P = 0.008), in comparison with the TGF-β1 group; similar results were found in HConFs (Fig. 1A). The results of IF assays reflected that the protein distributions of COL3 and α-SMA in the TGF-β1 + bromfenac group were more compact and wider than those in the TGF-β1 + bromfenac group and the control group, in HConFs (Supplementary Fig. S3A) and HPFs (Supplementary Fig. S3B). FN and COL3 are important compositions of ECM, and α-SMA is a reliable marker for myofibroblast activation, which enhances fibroblast contractile activity significantly. Collectively, these results indicated that bromfenac inhibits TGF-β1–induced ECM synthesis and myofibroblast activation in HConFs and HPFs.

The effects of using bromfenac alone were validated in HPFs and HConFs. After the treatment of bromfenac for 48 hours, the protein (Supplementary Fig. S4A) and mRNA (Supplementary Fig. S5) expression levels of FN, COL3, α-SMA, and survivin decreased in a dose-dependent manner. Proliferative marker survivin may play a role in the development of pterygium. The protein and mRNA levels of matrix metalloproteinase 3 were not changed by bromfenac in our study.

**Effects of Bromfenac on the Signaling Pathways in TGF-β1–Treated HConFs and HPFs**

HPFs from three patients and HConFs from three donors were treated with TGF-β1 (20 ng/mL) and bromfenac (90 μg/mL) for 48 hours; the effects of bromfenac and TGF-β1 on the signaling pathway were tested by Western blotting. The expression levels of p-AKT, p-p38 MAPK, and p-S9 in HConFs and HPFs were below the linear range for ECL detection in Western blotting, but became detectable after the treatment of TGF-β1. Besides, TGF-β1 rapidly (approximately 30 minutes) activated Smad2/3, p38 MAPK, AKT, and ERK1/2 by phosphorylating them (Supplementary Fig. S2B), as reported before.21 Phosphorylation at serine residue 9 of GSK-β3 represses its activity.41 Our data also showed that TGF-β1 inactivates GSK-β3 by phosphorylating GSK-3β-S9.

In HPFs, in comparison with the TGF-β1 group, the TGF-β1 + bromfenac group expressed lower levels of p-AKT (0.66 ± 0.08 folds, P = 0.032), p-ERK1/2 (0.69 ± 0.11 folds, P = 0.003), and p-GSK-3β-S9 (0.65 ± 0.10 folds, P = 0.002); similar results were seen in HConFs (Fig. 1B). The expression of p-Smad2/3, p-p38 MAPK, and the total protein levels of AKT, ERK1/2, GSK-3β, Smad3, and p38 MAPK were not changed by bromfenac in HConFs and HPFs. In addition, at 30 minutes after the treatments, TGF-β1–induced expression of p-AKT, p-ERK1/2, and p-GSK-3β-S9 in HConFs and HPFs was reduced by bromfenac in a dose-dependent manner; however, the increase of p-Smad2/3 and Smad3 induced by TGF-β1 was not changed by bromfenac (Supplementary Fig. S6).

Collectively, these results suggested that bromfenac inhibits TGF-β1–induced ECM synthesis and myofibroblast activation in HConFs and HPFs, and might through dephosphorylating AKT, ERK1/2, and GSK-3β-S9.

HPFs from three patients and HConFs from three donors were taken to validate the effects of using bromfenac alone. After the treatment of bromfenac for 48 hours, the phosphorylated protein levels of AKT, ERK1/2, and GSK-3β-S9 declined with dosage (Supplementary Fig. S4B).

**The PI3K/AKT and MEK/ERK Signaling Pathways Mediate TGF-β1–Induced ECM Synthesis and Myofibroblast Activation in HConFs and HPFs**

HPFs from three patients and HConFs from three donors were treated with TGF-β1 (20 ng/mL) and the indicated inhibitors for 48 hours; the effects of the inhibitors and TGF-β1 were tested by Western blotting. AKT signaling was blocked with the PI3K/AKT inhibitor wortmannin; ERK signaling was blocked with the MEK/ERK inhibitor U0126. In HPFs, in comparison with the TGF-β1 group, TGF-β1 + wortmannin group had lower expression level of p-AKT (0.41 ± 0.11 folds, P = 0.002), and the TGF-β1 + U0126 group had lower expression level of p-ERK (0.44 ± 0.05 folds, P < 0.001), respectively. Similar results were found in HConFs (Fig. 2A).

In HPFs, as compared with the TGF-β1 group, the TGF-β1 + wortmannin group had lower expression levels of FN (0.69 ± 0.05 folds, P = 0.001), COL3 (0.64 ± 0.06 folds, P = 0.007), and α-SMA (0.64 ± 0.09 folds, P = 0.002); wortmannin had similar effects on HConFs (Fig. 2B). In addition, in HPFs, TGF-β1 + U0126 group had lower expression levels of FN (0.60 ± 0.09 folds, P = 0.002), COL3 (0.53 ± 0.08 folds, P = 0.001), and α-SMA (0.48 ± 0.05 folds, P < 0.001), in comparison with the TGF-β1 group; U0126 had similar effects on HConFs (Fig. 2B).

Therefore, blocking either PI3K/AKT or MEK/ERK signaling pathway could reduce the TGF-β1–induced ECM synthesis and myofibroblast activation in HConFs and HPFs. From the above, a conclusion can be drawn that bromfenac inactivates the AKT and ERK pathways, both of which could modulate the expression of FN, COL3, and α-SMA.

**Effects of Bromfenac on the TGF-β1–Induced Cellular Migration and Proliferation on HConFs and HPFs**

The cellular migration ability was detected using wound healing assays. TGF-β1 increased the migration ability...
FIGURE 1. Effects of bromfenac on the TGF-β1-induced HConFs and HPFs. HPFs from three patients and HConFs from three donors were treated with vehicle, TGF-β1 (20 ng/mL) only, TGF-β1 (20 ng/mL) plus bromfenac (90 μg/mL), or bromfenac (90 μg/mL) only for 48 hours. Representative Western blot images of (A) FN, COL3, and α-SMA protein expression and (B) p-AKT, total AKT, p-ERK1/2, total ERK1/2, p-GSK3β-S9, total GSK3β, p-smad2/3, total Smad3, p-p38 MAPK, and total p38 MAPK expression. Quantitative analysis of immunoblots presents the mean values ± SD of three independent experiments. Total protein levels were normalized to GAPDH for loading control, whereas the phosphorylated proteins were normalized to their total proteins. One-way ANOVA was used to test the variations among groups, and followed by Bonferroni’s multiple comparison test, which was used for comparison between two groups. *P < 0.05, **P < 0.01, ***P < 0.001 versus vehicle; #P < 0.05, ##P < 0.01, ###P < 0.001 versus with TGF-β1.
The recurrence of pterygium is a process of pathological fibrosis, excessive wound healing accompanies the production of large amounts of ECM after pterygium excision surgeries. In our study, bromfenac inhibited the production of \( \alpha \)-SMA, FN, and COL3, and suppressed cellular migration and proliferation in TGF-\( \beta \)-induced HPFs and HConFs. Therefore, bromfenac has nice antifibrotic effects on HPFs and HConFs.

CCK-8 assays were performed to detect the cellular proliferation ability. TGF-\( \beta \) increased the proliferation ability (presents as a higher value of OD450) of HPFs (1.28 ± 0.13 folds, \( P = 0.034 \)), as compared with the control group. Similarly, TGF-\( \beta \) increased the proliferation ability of HConFs induced by TGF-\( \beta \). In HPFs, TGF-\( \beta \) + bromfenac group had lower proliferation ability (0.64 ± 0.03 folds, \( P = 0.002 \)), as compared with the TGF-\( \beta \) group. Similar effects of bromfenac were seen in HConFs (Fig. 3B).

### DISCUSSION

The recurrence of pterygium is a process of pathological fibrosis, excessive wound healing accompanies the production of large amounts of ECM after pterygium excision surgeries. In our study, bromfenac inhibited the production of \( \alpha \)-SMA, FN, and COL3, and suppressed cellular migration and proliferation in TGF-\( \beta \)-induced HPFs and HConFs. Therefore, bromfenac has nice antifibrotic effects on HPFs and HConFs.

The present study also demonstrated that blocking the PI3K/AKT or MEK/ERK pathway suppressed the synthesis of FN, COL3, and \( \alpha \)-SMA induced by TGF-\( \beta \). This indicated that the two pathways are essential to pterygium pathogenesis, and that inhibitions of them give promise of other treatment routines for pterygium. TGF-\( \beta \) has been identified as an inducer of myofibroblast activation in pterygia\(^{17} \); however, knowledge...
about the molecular mechanisms underlying the effects of TGF-β on pterygium is still incomplete. The p38 MAPK pathway has been reported to participate in the TGF-β-induced ECM protein synthesis and myofibroblast activation of HPFs. Previous studies have shown that the canonical Smad2/3 signaling and several other noncanonical pathways participate in the TGF-β1-induced fibrosis in many other diseases, such as breast cancer. TGF-β shows profibrotic effects through activation of the AKT/GSK-3β pathway in liver cancer cells. In TGF-β-induced lung fibroblasts and cutaneous microvascular endothelial cells, ERK/GSK-3β pathway participates in the fibrogenesis.

Our study indicated that bromfenac inhibits the phosphorylated protein levels of AKT, ERK1/2, and GSK-3β-S9 in HPFs and HConFs; meanwhile, blocking the PI3K/AKT or MEK/ERK pathway has similar effects with bromfenac, which leads to a decrease of ECM synthesis and myofibroblast activation. Generally, NSAIDs act by inhibiting COX-2 to block the
synthesis of prostaglandin E2 (PGE2). PGE2 plays a key role in the process of fibrosis and cell proliferation. It has been reported that PGE2 promotes cell proliferation and ERK phosphorylation in esophageal squamous cell carcinoma.36,37 Downregulation of ERK by RNA interference substantially weakens PGE2-induced cell proliferation.46 PGE2 induces the activation of the PI3K/AKT pathway, thus promoting cell migration and the invasion of colorectal carcinoma cells.48 Fjui et al.49 showed that both PKA and PI3K could function downstream of PGE2 to phosphorylate GSK-3β in the transformed HEK293 cell line. Activation of the E-series prostaglandin 2 (EP2) receptor by PGE2 leads to the binding of an α subunit of Gs protein to axin, which promotes the release of GSK-3β from the β-catenin degradation complex; meanwhile, the free βγ subunits of Gs protein stimulate the PI3K/AKT signaling cascade, which results in the phosphorylation and inactivation of GSK-3β.50 Deletion or inactivation of GSK-3β leads to the accumulation and nuclear translocation of β-catenin.51 In the nucleus, β-catenin binds to TCF/LEF family proteins to form a transcriptional activation complex and then activates the β-catenin dependent fibrosis processes,52,53 which include promoting collagen gel contraction, α-SMA expression, and cell migration in human dermal fibroblasts.54 In addition, there is a crosstalk network among ERK, PI3K/AKT, and Wnt/β-catenin signaling pathways.55

The present study demonstrated that the antiﬁbrotic effects of bromfenac are achieved at least in part by regulating the AKT and ERK pathways. Further research is needed to investigate the relation between GSK-3β and pterygium pathogenesis. Using speciﬁc activator of PI3K/AKT or MEK/ERK, which has the potential profibrotic effects, would be more satisfactory to further prove the mechanisms of bromfenac. Meanwhile, the mechanisms underlying the antiﬁbrotic effects of bromfenac are also needed to be fully understood. It is also necessary to investigate the effects of bromfenac on pterygium epithelial cells. Comparing pterygium with normal conjunctiva by histological experiments is needed to study the pathogenesis of pterygium. Because our study is purely in vitro, further preclinical validation is necessary.

In summary, bromfenac inhibits TGF-β1-induced synthesis of the ECM proteins, myofibroblast activation, and cellular migration and proliferation in HPFs and HConFs, at least in part through inactivating AKT and ERK1/2. Besides, bromfenac has already been used clinically for its analgesic and antiinflammatory effects. These ﬁndings suggest that bromfenac may be a promising adjuvant drug for pterygium surgery to prevent recurrence.

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