A PPAR-Gamma Agonist Rosiglitazone Suppresses Fibrotic Response in Human Pterygium Fibroblasts by Modulating the p38 MAPK Pathway

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Purpose. Fibroblast activation may play an important role in pterygium progression. Synthetic peroxisome proliferator-activated receptor γ (PPAR-γ) ligands have been shown to be effective antifibrotic agents against transforming growth factor β1 (TGF-β1) induced fibrosis in several tissues. We aimed to investigate the antifibrotic effects of the PPAR-γ ligand rosiglitazone in pterygium fibroblasts and the underlying mechanisms.

Methods. Profibrotic activation was induced by TGF-β1 in primary cultured human pterygium fibroblasts and the effect of rosiglitazone treatment on α-smooth muscle actin (α-SMA), and extra cellular matrix proteins synthesis was detected by western blotting, real-time PCR, immunostaining, and flow cytometry. Pharmaceutical inhibition of PPAR-γ receptor was used to determine the dependency or otherwise of rosiglitazone’s action on PPAR-γ signaling. Major signaling pathways downstream of TGF-β1 were investigated by western blotting to assess their possible association with rosiglitazone’s effect. Cell viability and apoptosis were investigated to assess drug-induced cytotoxicity, and the effect of rosiglitazone treatment on cell migration was further determined.

Results. α-SMA and fibronectin synthesis induced by TGF-β1 were suppressed by rosiglitazone treatment in a dose-dependent manner. Rosiglitazone also inhibited intrinsic TGF-β1 expression. Smad2/3, ERK1/2, and P38 pathways were activated in response to TGF-β1. Rosiglitazone suppressed TGF-β1-induced P38 MAPK activation, while ERK1/2 and Smad2/3 signaling remained unaffected. The observed antifibrotic effect of rosiglitazone was not affected by the PPAR-γ antagonist GW9662, indicating it is not PPAR-γ dependent. Rosiglitazone also inhibited the proliferation and migration of pterygium fibroblasts.

Conclusions. Rosiglitazone suppresses TGF-β1-induced myofibroblast activation and extra cellular matrix synthesis in pterygium fibroblasts at least partly through the modulation of the p38 MAPK pathway.

Keywords: pterygium, cornea, fibrosis, PPAR-γ, rosiglitazone

Pterygium is an ocular disorder that clinically manifests as a benign growth of conjunctival epithelial tissues over the cornea. The characteristic origin and location of pterygium is the nasal limbus, from where it grows toward the cornea. Pterygium is characterized by extensive cell proliferation, connective tissue remodeling, angiogenesis and chronic inflammation resulting in redness and irritation in the area.1,2 The condition can significantly affect vision in advanced cases owing to a distortion of the shape in the cornea3 and can progress to ocular surface squamous neoplasia.4 Surgical excision is the standard treatment for pterygium. Wound healing after pterygium excision often leads to fibrosis characterized by recurrent pterygium. The bare sclera technique for instance is associated with high postsurgical recurrence rates reported to range from 38% to 88%.5-8 Because fibrovascular growth in recurrent pterygia is more extensive than that in primary disease,9 recurrence is of major concern as it worsens prognosis. To reduce postsurgical recurrences, mitomycin C application10 has been widely attempted, but numerous adverse effects such as scleral stromalysis11 and corneal melting12 have been reported. There is therefore the need for a safer and more targeted curative and preventive therapy.

Studies have shown the overexpression of transforming growth factor β1 (TGF-β1), a profibrotic agent, in pterygium tissues compared to normal conjunctiva cultures and in recurrent pterygium fibroblast cultures compared to primary pterygium fibroblast cultures.13,14 TGF-β1 signaling stimulates fibroblast migration, proliferation, and myofibroblast differentiation.15 The presence of myofibroblasts in pterygium is suggestive of the involvement of TGF-β1 in the pathogenesis and progression of the disorder.16 Other research findings also support the presence of myofibroblasts in the fibrovascular tissue of primary and recurrent pterygia.17,18

Previous research works have shown several synthetic ligands of peroxisome proliferator-activated receptor γ (PPAR-γ) that are effective in preventing fibrosis in various organs including the skin, kidney, and cornea.19-21 Notably, corneal fibrosis following
laser refractive ablation was suppressed by the topical application of a PPAR-γ ligand. The topical application of these synthetic PPAR-γ ligands and the possibility of preventing corneal haze are advantageous to their potential use in ophthalmology. We thus aimed to determine the antifibrotic properties of the PPAR-γ ligand rosiglitazone in cultured primary human pterygium fibroblasts and the possible signaling pathways mediating any observed effect.

METHODS

Ethical Statement

Pterygium tissues were obtained from patients who underwent pterygium excision surgery. This study protocol was approved by the Institutional Review Board of our institute (YWMR-15-0-053), and all subjects were treated in accordance with the Declaration of Helsinki. Written informed consent to participate in this research was obtained from each patient.

Isolation and Culture of Primary Human Pterygium Fibroblasts

The pterygium tissue sample was placed in a culture plate and allowed to attach. Dulbecco’s modified Eagle’s medium F12 (DMEM/F12; Welgene, Gyeongsan, Korea) with 10% fetal bovine serum (FBS) and 1% antibiotic/antimycotic (Gibco Life Technologies, CA) in dimethyl sulfoxide (DMSO) was added and the tissue incubated at 37°C in a humidified atmosphere of 5% CO2 during which cells migrated from the explant. Migrated cells were passaged with 0.25% trypsin (Gibco Life Technologies) and subcultured in DMEM (Welgene) with 5% FBS. Passage 3-5 cells were used for all experiments.

Western Blotting

The pterygium fibroblasts were seeded into culture plates in DMEM with 5% FBS. After attachment, serum-free media was incubated at 37°C for 24 hours. The cells were passaged with 0.25% trypsin and subcultured in DMEM (Welgene) with 5% FBS. Passage 3-5 cells were used for all experiments.

Flow Cytometry

The pterygium fibroblasts were cultured on glass chamber slides and pretreated with rosiglitazone (75 μM) in serum-free DMEM, followed by TGF-β1 (2.5 ng/mL). After 48 hours of incubation, the cells were fixed with 4% paraformaldehyde (PFA) and blocked with 10% fetal bovine serum. The slides were incubated with primary antibodies against α-SMA and fibronectin overnight at 4°C and then with Alexa Fluor labelled secondary antibodies (Thermo Fisher Scientific, Rockford, IL, USA) and incubated at room temperature in the dark for 2 hours. The nuclei were counterstained with 4’,6-diamidino-2-phenylindole (DAPI), and immunofluorescence was visualized and digitally captured using a confocal microscope (Leica Microsystems, GmbH, Wetzlar, Germany). Internal control was performed using PBS and secondary antibodies without the primary antibodies. The proportion of α-SMA positive cells was quantified using indirect flow cytometry. Brieﬂy, cell suspensions (approximately 1 × 10^6 cells/100 μL) prepared in stain buffer (ice cold PBS, 2% FBS, 5% sodium azide) were ﬁxed for 10 minutes with 1% PFA. After washing, the cells were suspended in 500 μL permeability buffer (stain buffer + 0.2% saponin), incubated in the dark for 5 minutes, and washed again. The cell pellets obtained after centrifugation at 1500 rpm for 5 minutes were resuspended in permeability buffer and incubated with primary antibodies against α-SMA for 30 minutes on ice in the dark. After washing, the cells were then incubated with an Alexa Fluor 488-conjugated secondary antibody for another 30 minutes, washed, and analyzed by flow cytometry.

Cell Viability Test

The pterygium fibroblasts were seeded into a 96-well plate at a density of 5000 per well and incubated for 24 hours. The cells were then treated with rosiglitazone (25, 50, 75, and 100 μM).
Control cells were vehicle treated. After 48 hours of incubation, the cell counting kit 8 (CCK-8) assay was performed according to kit manufacturer’s protocol (Dojindo Laboratories, Kumamoto, Japan). After a 2-hour incubation, the optical absorbance was read at 450 nm on a microplate reader. The experiment was done in triplicate for each concentration of rosiglitazone.

Apoptosis Test

The pterygium fibroblasts were pretreated with rosiglitazone (25 and 75 μM) in serum-free DMEM after which TGF-β1 (2.5 ng/mL) was added. Cells with only TGF-β1 treatment were used as control. After a 24-hour incubation, the cells were washed with cold PBS and resuspended in 1X binding buffer at a concentration of 1 x 10^6 cells/mL. One hundred microliters of the solution with approximately 1 x 10^5 cells was transferred to a 5-mL culture tube and 5 μL each of FITC annexin V and propidium iodide (PI) (BD Pharmingen, San Jose, CA, USA) added. After 15 minutes of incubation at room temperature in the dark, 400 μL 1X binding buffer was added to each tube and analyzed by flow cytometry within 1 hour.

Cell Migration Test

The pterygium fibroblasts were seeded into culture plates and incubated for 24 hours to allow the cells to attach to the plate surface. Further, scratch wounds were induced with a pipette tip (1000 μL), and the suspended cells were washed off. The cells were then pretreated with rosiglitazone (75 μM), followed by treatment with TGF-β1 (2.5 ng/mL) in serum-free media. The wounds were monitored using phase-contrast microscopy and photographed at 12, 24, and 48 hours. The ImageJ software was used to determine the wound area. Vehicle-treated cells were used as controls.

Statistical Analysis

The data were analyzed using Prism 5.0 software. Statistical comparisons were done using Student’s t-test or 1-way analysis of variance (ANOVA), as appropriate. All results are expressed as mean ± SD. Statistical significance was defined as P < 0.05 in all cases.

RESULTS

Rosiglitazone Decreased TGF-β1-Induced Synthesis of ECM Proteins and Myofibroblast Differentiation in Pterygium Fibroblasts Independent of PPAR-γ

Basal levels of α-SMA, a key marker for myofibroblast differentiation, and fibronectin, an ECM protein, in the pterygium fibroblasts were detectable (Fig. 1). There was a marked increase of approximately threefold in the expression α-SMA (P < 0.0001, F[5, 12] = 31.75), (C) fibronectin (P < 0.0001, F[5, 12] = 145.1), and (D) vimentin (P = 0.1989, F[5, 12] = 1.745). #Represents P < 0.05 compared to vehicle-treated control cells; *represents P < 0.05 compared to only TGF-β1-treated cells.

The observed antifibrotic effect of rosiglitazone was seen to be independent of PPAR-γ receptor as treatment with rosiglitazone together with GW9662 did not affect its antifibrotic effect. These western blot results were confirmed by the relative mRNA expression of these proteins, as well as collagen 1 and serum response factor (SRF), which is a transcription factor for actin cytoskeletal reorganization and myofibroblast transformation.24 The mRNA expression levels of α-SMA, collagen 1, fibronectin, and SRF also tended to show a dose-dependent antifibrotic effect of rosiglitazone in pterygium fibroblasts, whereas no significant change was observed in
the expression of vimentin (Fig. 2). Immunostaining results showed decreased expression of α-SMA and fibronectin in rosiglitazone-treated cells (Fig. 3A). Indirect flow cytometry results revealed that rosiglitazone treatment decreased the proportion of α-SMA-positive cells, which further confirmed its antifibrotic effect in pterygium fibroblasts (Fig. 3B).

**Rosiglitazone Decreased Intrinsic TGF-β1 Production in Pterygium Fibroblasts Independent of PPAR-γ**

Our results showed a dose-dependent decrease in the intrinsic expression of TGF-β1 when the cells were treated with rosiglitazone, indicating a modulatory effect of rosiglitazone on TGF-β1 protein synthesis. In addition, this effect was independent of PPAR-γ receptor, as cotreatment with rosiglitazone and GW9662 did not reverse the decreased synthesis of TGF-β1. The basal levels of αSMA and fibronectin proteins also tended to decrease in a dose-dependent manner subsequent to the decreased TGF-β1 expression (Fig. 4).

**Rosiglitazone Inhibited TGF-β1-Induced Activation of p38 MAPK Signaling Pathway in Pterygium Fibroblasts**

To determine the possible mechanisms underlying the antifibrotic effect of rosiglitazone, the major signal transduction pathways downstream of TGF-β1 were examined. As shown, TGF-β1 increased the activation of p38 MAPK, ERK1/2, and Smad2/3 from the basal level indicating increased signaling. However, the p38 MAPK signaling was decreased by rosiglitazone treatment as seen by the decreased phosphorylated p38 MAPK protein levels. This effect of rosiglitazone was not observed with ERK1/2 and Smad2/3 signaling as the phosphorylated protein levels remain unaltered (Fig. 5).

**TGF-β1 Induces Synthesis of ECM Proteins and Differentiation of Myofibroblasts via p38 MAPK Signaling in Pterygium Fibroblasts**

To test whether p38 was involved in the TGF-β1-dependent phenotypic transformation of myofibroblasts in pterygium fibroblasts, we first investigated the time course of p38 MAPK activation. As shown in Figure 6A, the TGF-β1-induced activation of p38 MAPK signaling peaked at 45 minutes and lasted for at least 24 hours. This indicated that rosiglitazone downregulated the p38 MAPK pathway. Maintenance of this downregulation by rosiglitazone on p38 activation for at least 24 hours (Fig. 6B) confirmed the role of p38 signaling in the antifibrotic effect of rosiglitazone. Further, the kinase inhibitor, SB203580, blocked the p38 signaling pathway and led to a dose-dependent decrease in the expression of αSMA and fibronectin proteins (Figs. 6C, 6D, 6E). This indicated that TGF-β1-induced myofibroblast transformation and ECM synthesis occurs at least partly through p38 MAPK signaling in pterygium fibroblasts.

**FIGURE 2.** Real-time PCR analysis. Quantitative analysis of mRNA expression levels of (A) α-SMA ($P < 0.0001$, $F[5, 12] = 18.78$), (B) collagen 1 ($P = 0.0396$, $F[5, 12] = 3.471$), (C) fibronectin ($P = 0.0011$, $F[5, 12] = 9.447$), (D) SRF ($P = 0.0398$, $F[5, 12] = 9.447$), and (E) vimentin ($P = 0.2412$, $F[5, 12] = 1.593$). *Represents $P < 0.05$ compared to vehicle-treated control cells; *represents $P < 0.05$ compared to only TGF-β1-treated cells.
Rosiglitazone Reduced Pterygium Fibroblasts Proliferation and Cell Migration, But Did Not Induce Cell Death

Results of fluorescence-activated cell sorting (FACS) analysis of cells treated with rosiglitazone compared with vehicle-treated controls and cells treated with only TGF-β1 showed approximately the same percentage of distribution of live cells (80%), cells at early apoptosis (6%), cells at late apoptosis (11%), and necrotic cells (3%), indicating that rosiglitazone treatment did not induce apoptosis. However, CCK-8 assay results showed that rosiglitazone treatment significantly caused a reduction in cell number dose dependently (Fig. 7) indicative of an inhibitory effect of rosiglitazone on pterygium fibroblast proliferation. Rosiglitazone also suppressed the TGF-β1-induced cell migration in pterygium fibroblasts (Fig. 8).

Rosiglitazone Reduced the TGF-β1-induced Increase in MMP-9 in Pterygium Fibroblasts

We investigated the effect of rosiglitazone on the expression of MMP-1, -3, and -9, which are known to be highly expressed in pterygium fibroblasts. Results showed that TGF-β1 increased the expression of MMP-3 and -9, but not MMP-1. In addition, rosiglitazone significantly suppressed the TGF-β1-induced expression of MMP-9. However, it had no effect on the expression of MMP-1 and -3 (Supplementary Fig. S1).

DISCUSSION

Recent studies identifying PPAR-γ as an important cellular antifibrotic mechanism and its importance in the negative regulation of connective tissue biosynthesis during both physiologic and pathological matrix remodeling, and the subsequent consistent findings in different tissue types with different PPAR-γ ligands have increased research focus in this area in the quest to finding potential therapeutic agents for the myriad chronic debilitating fibrotic disorders. In this regard, PPAR-γ ligands have also been reported to be effective corneal antifibrotics in vitro and in vivo a PPAR-γ-independent manner. However, their effects on other ocular fibrotic disorders, such as pterygium, which is known to overexpress TGF-β1, is not known. Our results showed that the PPAR-γ ligand rosiglitazone is an effective antifibrotic agent in pterygium fibroblasts as well in vitro.

Notably, our results also showed that rosiglitazone can reduce the basal levels of α-SMA and fibronectin, thereby making it a very suitable candidate for therapy. This assertion is further consolidated by the additive downregulation of the intrinsic expression of TGF-β1 in pterygium fibroblasts, which
FIGURE 4. Effect of rosiglitazone on intrinsic TGF-β1 synthesis in pterygium fibroblast. (A) Representative western blot images of α-SMA and fibronectin protein expression. Quantitative analysis of immunoblots with summary data from three independent experiments from (B) α-SMA ($P = 0.0004$, $F[4, 10] = 14.05$) and (C) fibronectin ($P = 0.0007$, $F[4, 10] = 12.18$). (D) Representative western blot images of TGF-β1 expression. (E) Quantitative analysis of immunoblots with summary data from three independent experiments ($P = 0.0068$, $F[4, 10] = 6.714$). *Represents $P < 0.05$ compared to control cells.

FIGURE 5. Effects of rosiglitazone on signaling pathways following exposure to TGF-β1 for 2 hours. (A) Representative western blot images. Quantitative analysis of immunoblots for (B) P-P38/T-P38 ($P = 0.0008$, $F[5, 12] = 9.378$), (C) P-ERK/T-ERK ($P = 0.1599$, $F[5, 12] = 1.946$), and (D) P-Smad/T-Smad ($P = 0.0638$, $F[5, 12] = 2.848$), with summary data from three independent experiments. *Represents $P < 0.05$ compared to only TGF-β1-treated cells.
Role of p38 signaling in the TGF-β1-mediated fibrosis and the effect of rosiglitazone. (A) Representative western blot images of the time course of p38 protein activation, (B) effect of rosiglitazone on p38 signaling following exposure to TGF-β1 for 24 hours, and (C) protein expression of α-SMA and fibronectin after blocking p38 signaling with SB203580. Quantitative analysis of the immunoblots for (D) α-SMA \(P < 0.0001, F[5, 12] = 36.09\) and (E) fibronectin \(P < 0.0001, F[5, 12] = 97.39\) with representative data from three independent experiments. #Represents \(P < 0.05\) compared to vehicle-treated control cells; *represents \(P < 0.05\) compared to only TGF-β1-treated cells.

The effect of rosiglitazone on cell viability. FACS results of (A) vehicle-treated control cells, (B) cells treated with only TGF-β1, (C) cells treated with TGF-β1 and 25 μM rosiglitazone, (D) cells treated with TGF-β1 and 75 μM rosiglitazone, and (E) their quantitative representation. (F) Quantitative representation of CCK-8 assay results after treatment with different concentrations of rosiglitazone compared with vehicle-treated control cells \(P < 0.0001, F[4, 10] = 56.74\). *Represents \(P < 0.05\) compared to vehicle-treated control.
suggests that rosiglitazone modulates TGF-β1 induced fibrogenesis. Given that pterygium fibroblasts exhibit aberrant ECM remodeling and are associated with MMPs, the effect of rosiglitazone on the production of MMPs provides another promising perspective. Investigation of the expression of MMP-1, -3, and -9 in pterygium fibroblasts showed that rosiglitazone significantly suppressed the TGF-β1-induced expression of MMP-9. However, it had no effect on MMP-1 and -3. Because MMP production is mainly associated with NF-kB pathway, further studies to understand the underlying mechanism are warranted. The noncytotoxic and antiproliferative effects of rosiglitazone treatment as shown by the cell viability assays are also an added advantage compared to the currently used therapeutic applications. Thus, rosiglitazone’s effect appears to be mediated by limiting the proliferation of pterygium fibroblasts and their transformation into myofibroblasts.

Our evaluation of the possible molecular mechanisms underlying the observed antifibrotic effects of rosiglitazone indicated that rosiglitazone inhibits TGF-β1-induced fibrosis in pterygium fibroblasts at least in part by modulating the noncanonical p38 MAPK cellular signaling pathway. As noticed, TGF-β1-induced phosphorylated Smad protein levels remained unaltered with rosiglitazone treatment, while phosphorylated P38 protein levels were downregulated with a consequent significant reduction in ECM synthesis and myofibroblast differentiation in a dose-dependent manner, albeit not to the baseline levels. Blocking of the p38 signaling in pterygium fibroblasts led a dose-dependent decrease in myofibroblast differentiation and ECM synthesis similar to the effect of rosiglitazone treatment. The MAPK signaling pathways, which include ERK1/2, jun kinase (JNK/SAPK), and p38 pathways, are known to play a key role in the transduction of extracellular signals to elicit cellular responses including proliferation, differentiation, development, inflammation, and apoptosis. These pathways relay, amplify, and integrate signals from different stimuli, such as growth factors by phosphorylating proteins, which then translocate to the nucleus to activate transcription factors to regulate the expression of target genes. Indeed, TGF-β1 stimulated p38 MAPK activation has been reported to be involved in fibrosis and scar tissue formation, and research findings in human cornea and tenon fibroblasts indicate that rosiglitazone inhibits TGF-β1-induced activation of these fibroblasts via the p38 MAPK signaling pathway. The involvement of this pathway in the TGF-β1 stimulated fibrotic process in pterygium fibroblasts as our results also indicate is consistent with these reports. However, both rosiglitazone and p38 inhibition failed to completely block myofibroblast differentiation and ECM synthesis. This indicated that the TGF-β1-induced fibrosis is only partially dependent on p38 MAPK signaling. Previous studies have shown that the canonical Smad2/3 signaling and several other noncanonical pathways are involved in TGF-β1-induced fibrosis, thus indicating a complex regulatory linkage between TGF-β1 signaling and the development of fibrosis.

The observed antifibrotic effects of rosiglitazone were also seen to be independent of PPAR-γ receptors as treatment with the PPAR-γ receptor antagonist GW9662 did not reverse the effects. This is consistent with results obtained in cornea fibroblasts treated with rosiglitazone and other PPAR-γ ligands. PPAR-γ agonists modulate multiple cellular functions both in a PPAR-γ-dependent and PPAR-γ-independent manner as indicated by several research findings with different PPAR-γ ligands from different tissues. The dependence or otherwise of rosiglitazone on PPAR-γ signaling with regard to its antifibrotic effects seems to be tissue specific as similar results demonstrated in autosomal dominant polycystic kidney disease.

**Figure 8.** Effect of rosiglitazone on pterygium fibroblasts. (A) Phase-contrast microscopic images of the scratch wound area of cells treated with vehicle, TGF-β1 (2.5 ng/mL) only, and TGF-β1 (2.5 ng/mL) along with rosiglitazone (75 μM), taken at 12, 24, and 48 hours after treatment. (B) Quantitative analysis of the scratch wound area with representative data from three different sites of the vehicle-treated control cells (P = 0.0292, F[2, 6] = 6.739), cells treated with only TGF-β1 (P < 0.0001, F[2, 6] = 432.2), and cells treated with TGF-β1 and rosiglitazone (P = 0.0101, F[2, 6] = 10.89). *Represents P < 0.05 compared to the scratch wound area at 12 hours after treatment.
disease (ADPKD) cyst-lining epithelial cells were observed to be PPARγ dependent. In human corneal fibroblasts, the antifibrotic effects of two other synthetic PPARγ ligands, CDDO and 15d-PGJ2, were seen to be largely independent of PPARγ, also indicative of the different mechanisms by which these ligands function. Further research is necessary to evaluate the antifibrotic effect of other PPARγ ligands to obtain the best alternative for therapeutic use. Although the PPARγ-independent nongenomic effect of rosiglitazone has been reported, the receptors possibly targeted and underlying mechanisms are not fully understood. We speculate that Ca2+ acts as a possible mediator for the cross-talk between rosiglitazone and TGF-β1 signaling. Previous studies have shown that TGF-β1 evokes Ca2+ transients in human pulmonary fibroblasts, which when disrupted interfere with ECM production, rosiglitazone is known to affect Ca2+ homeostasis. In corneal fibroblasts, activation of transient receptor potential cation channel subfamily V member 1 (TRPV1) with a subsequent increase in intracellular Ca2+ is associated with myofibroblast transformation, and it is also upregulated in the pterygium. Further research is necessary to fully understand the mechanism underlying the antifibrotic effect of rosiglitazone.

In summary, our study demonstrated that rosiglitazone is effective in reducing ECM production and myofibroblast differentiation independent of PPARγ. These findings may present the basis for its use as a promising candidate drug for adjuvant therapy after surgery to prevent recurrent pterygium.

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References


