Inhibition of TGFBIp Expression by Lithium: Implications for TGFBI-Linked Corneal Dystrophy Therapy

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PURPOSE. The purpose of this study was to investigate the effects and molecular mechanisms of lithium on inhibition of TGFBIp expression as a potential therapy for TGFBI-linked corneal dystrophy.

METHODS. Primary culture corneal fibroblasts were isolated from the corneas of healthy subjects and patients with granular corneal dystrophy type 2 (GCD2) with a homozygous mutation in TGFBI. Levels of TGFBIp and its mRNA in corneal fibroblasts treated with various lithium (LiCl) concentrations were analyzed by Western blot, RT-PCR, and quantitative real-time PCR.

RESULTS. LiCl treatment reduced the expression levels of normal and mutant TGFBIp in a dose- and time-dependent manner. Furthermore, TGF-β1-induced TGFBIp expression decreased by 35% and 67% after treatment with 5 mM and 10 mM LiCl, respectively. LiCl decreased the level of pSmad3 (S423/425) in a dose- and time-dependent manner. Furthermore, LiCl increased the level of pGSK-3β (S21/9) in a dose-dependent manner. Also observed was the interaction between GSK-3β and Smad3, which was enhanced by lithium. In addition, Western blot analysis showed that the ratio of LC3-II/LC3-I in corneal fibroblasts increased after LiCl treatment. Cell viability at different doses was greater than 98%, indicating that LiCl did not induce significant corneal fibroblast death. Finally, the observed attenuating effects of LiCl on TGFBIp expression were not the results of cell death.

CONCLUSIONS. The accumulation of mutant TGFBIp ultimately leads to the histopathologic and clinical manifestations associated with TGFBI-linked corneal dystrophy. These data strongly suggest that lithium may be used for the prevention or treatment of this disease. (Invest Ophthalmol Vis Sci. 2011;52:3293–3300) DOI:10.1167/iovs.10-6405

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Materials and Methods

Isolation and Culture of Primary Corneal Fibroblasts

Normal and homozygote primary corneal fibroblasts were prepared using previously described methods. Donor confidentiality was maintained in accordance with the Declaration of Helsinki, and the conduct of the study was approved by the Severance Hospital IRB Committee (CR04124), Yonsei University. GCD2 was diagnosed by DNA sequence analysis for gene mutations. After removal of the corneal button (mut-TGFBIp) production through the regulation of TGF-β signaling with GSK-3 inhibitors may be a potential therapy for TFGBlip-mediated corneal dystrophy. Therefore, we investigated the effects of lithium treatment on TFGBlip expression. We found that these treatments reduced TFGBlip expression dramatically in corneal fibroblasts. To our knowledge, this is the first demonstration of decreased TFGBlip expression mediated through the inhibition of TGF-β signaling by lithium. These results suggest that the use of clinical trials of lithium to treat TFGBlip-mediated corneal dystrophy is a rational strategy.

Real-Time PCR

Primers for the human TGFBlip and β-actin specific genes were designed according to the published sequences available in GenBank using Primer3, as shown in Table 1. For the analysis of each sample, gene expression levels were calibrated using β-actin expression levels as an internal control. Two micrograms of total DNA was reverse-transcribed to cDNA using reverse transcriptase (Superscript One-Step RT-PCR System; Invitrogen Life Technologies) and primers (Table 1). Amplification products were visualized by electrophoresis in 1.2% agarose gels containing ethidium bromide.

Preparation of Cell Lysates, Immunoprecipitation, and Immunoblot Analysis

Cell lysates from corneal fibroblasts were prepared in radioimmunoprecipitation assay buffer (RIPA buffer; 150 mM NaCl, 1% NP-40, 0.5% deoxycholate, 0.1% SDS, 50 mM Tris-HCl pH 7.4) containing protease inhibitors (Complete Mini Protease Inhibitor Cocktail Tablet, catalog no. 1836170; Roche, Indianapolis, IN) and phosphatase inhibitors (PhosSTOP, catalog no. 04090845001; Roche). Crude cell lysates were centrifuged at 10,000g for 10 minutes at 4°C to remove nuclear fragments and tissue debris. A portion of the supernatant was used to determine the total protein concentration using a bicinchoninic acid kit (Pierce).

Total cell lysate (5 mg protein) was subjected to immunoprecipitation with anti-GSK-3 and anti-Smad3 antibodies, as indicated, overnight at 4°C with gentle agitation, followed by incubation (Protein A/G
Plus-Agarose (Invitrogen) for 2 to 4 hours at 4°C. After binding, the beads were washed extensively three times with RIPA buffer and then mixed with 2× SDS sample buffer and boiled for 5 minutes.

For immunoblot analysis, total cellular protein was subjected to electrophoresis on 10% Tris-glycine SDS polyacrylamide gels. Proteins were transferred onto polyvinylidene difluoride membranes (Millipore Corp., Bedford, MA), blocked in 5% dry milk in Tris-buffered saline containing Tween-20 (TBS-T) (0.02 M Tris/0.15 M NaCl, pH 7.5, 0.1% Tween 20) at room temperature for 1 hour, washed three times with TBS-T, and incubated with antibodies against TGFBIp (0.2 μg/mL; catalog no. AF2955; R&D Systems), β-actin (1:5000 dilution; catalog no. A-5441; Sigma-Aldrich), and GSK-3α/β, pGSK-3α/β, Smad3, and pSmad3 (1:1000 dilution; Cell Signaling Technology, Beverly, MA) overnight at 4°C. After three washes with TBS-T, blots were incubated with HRP-conjugated secondary antibody (anti-mouse IgG [1:5000 dilution; catalog no. NA931V] or anti-rabbit IgG [1:5000 dilution; catalog no. NA934V]; Amersham Pharmacia Biotechnology, Piscataway, NJ) at room temperature for 1 hour. Western blot analysis was visualized using enhanced chemiluminescence (Pierce). Immunoreactive protein bands were scanned at two intensities, and the optical densities of the bands were quantified computationally with the use of ImageJ software (developed by Wayne Rasband, National Institutes of Health, Bethesda, MD; available at http://rsb.info.nih.gov/ij/index.html). The background was corrected through subtraction, and protein levels were normalized relative to the intensity of the corresponding β-actin protein bands.

Cell Viability

Corneal fibroblasts were plated in 96-well plates at a density of 10,000 cells/well and incubated overnight. Cell proliferation was determined using the CellTiter 96 AQuosce One Solution Reagent Cell Proliferation Assay Kit (Promega, Madison, WI) and tetrazolium (3-[4,5-di-methylthiazol-2-yl]-5-[3-carboxymethoxyphenyl]-2-[4-sulphophenyl]-2H-tetrazolium, inner salt; MTS) (catalog no. G5380; Promega). Briefly, the culture medium was removed, and 20 μL MTS solution was added to each well containing 100 μL culture medium. Cultures were incubated at 37°C for 2 to 4 hours under 95% humidity and 5% CO2. Optical density was measured at 450 nm using a plate reader.

Statistical Analysis

The results were evaluated statistically for significance (P < 0.05) using one-way analysis of variance followed by the Newman-Keuls multiple comparison test. Data are expressed as mean ± SD. All data were processed using scientific graphing analysis software (Prism, version 4.0; GraphPad Software Inc., San Diego, CA).

RESULTS

TGFBIp Expression Is Downregulated by Lithium in Corneal Fibroblasts

Recently, several studies have revealed that inhibitors of GSK-3 can downregulate TGFBIp expression by blocking TGF-β signaling.30 –32 To test this hypothesis, we analyzed levels of TGFBIp in primary corneal fibroblasts treated with lithium, a known GSK-3 inhibitor. Western blot analysis showed that lithium reduced levels of TGFBIp in a dose- (Fig. 1A) and a time-dependent manner (Fig. 1B). We next examined the effect of lithium on TGFBIp expression using quantitative reverse transcription-PCR (RT-PCR). As shown in Fig. 1C, lithium reduced TGFBIp mRNA levels in a dose-dependent manner. The results from RT-PCR were confirmed by Western blot analysis (Fig. 1D). The Western blot data were quantified using NIH Image J and normalized to β-actin levels. The background was corrected through subtraction and normalized to the control. All data were expressed as mean ± SD of at least three independent experiments. Statistical analysis was performed using one-way analysis of variance followed by the Newman-Keuls multiple comparison test. *P < 0.05; **P < 0.01 with LiCl versus without LiCl.
effects of lithium on TGFBI mRNA expression. Quantitative RT-PCR showed that treatment with LiCl for 12 hours resulted in decreased TGFBI mRNA expression in a dose-dependent manner (Fig. 1C). Quantitative analysis revealed that treatments with 10 mM, 30 mM, and 50 mM LiCl significantly decreased TGFBI mRNA expression by approximately 81.43% ± 8.73%, 74.10% ± 4.41%, and 70.60% ± 2.69%, respectively, relative to β-actin, which was used as a control (P < 0.05) (Fig. 1D). Taken together, these results suggest that TGFBIp regulation occurs at the level of transcription.

In addition, we analyzed the expression of mut-TGFBIp in GCD2 homozygous (HO) corneal fibroblasts after lithium treatment to test whether the inhibitor also blocked the expression of GCD2-linked mutant TGFBIp. Treatment of wild-type (WT) corneal fibroblasts with 10, 30, and 50 mM LiCl for 12 hours reduced TGFBIp levels by 58.11% ± 9.06%, 32.46% ± 5.79%, and 14.80% ± 3.94%, respectively, compared with samples treated only with TGF-B1.

### Lithium Inhibits TGF-β Signaling by Inhibition of Smad3 Phosphorylation

At least two mechanisms are conceivable for how LiCl decreases TGFBIp protein levels. LiCl enhances TGFBIp degradation or decreases TGFBIp biosynthesis. After identification of Smad3/GSK-3 complexes, Guo et al.37 suggested that GSK-3 can activate Smad3. Further, lithium is well established as a GSK-3 inhibitor. Therefore, we examined the effects of LiCl on TGFBIp expression by investigating the TGF-β signaling pathway. Treatment of corneal fibroblasts with 10, 30, and 50 mM LiCl reduced Smad3 phosphorylation (5423/425) by 46.09% ± 5.53%, 31.29% ± 6.70%, and 20.37% ± 6.29%, respectively (Figs. 3A, 3B). In addition, LiCl interacts with GSK-3α/β and negatively regulates its activity (Fig. 3A, 3C, 3D). Treatment of corneal fibroblasts with 10, 30, and 50 mM LiCl for 12 hours increased GSK-3α phosphorylation significantly by 317.3% ± 38.00%, 401.0% ± 43.86%, and 512.0% ± 23.90%, respectively (P < 0.01; Fig. 3A, 3C). GSK-3β phosphorylation also increased significantly (Figs. 3A, 3D). Taken together, these results suggest that the observed lithium-induced decrease in TGFBIp levels was caused by the inhibition of TGF-β signaling.

### Interaction of Smad3 with GSK-3β

To examine whether Smad3 interacts with GSK-3β in corneal fibroblasts, we used healthy corneal fibroblasts. Endogenous GSK-3β was detected by the anti-GSK-3β antibody. When the lysates were immunoprecipitated with the anti-GSK-3β antibody, Smad3 was detected in the GSK-3 immune complex (Fig. 4A, lane 3). To examine the effect of LiCl on the interaction between GSK-3β and Smad3, we used lysates of corneal fibroblasts treated with 0, 30, and 50 mM LiCl concentration. When the lysates were immunoprecipitated with the anti-Smad3 antibody, detection of GSK-3β increased in the GSK-3

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**Figure 2.** TGF-β1-induced TGFBIp expression is inhibited by LiCl. (A) A healthy corneal fibroblast cell line was treated with the indicated amounts of TGF-β1 for 12 hours. Cell lysates and media were analyzed for TGFBIp by Western blot analysis. (B) A healthy corneal fibroblast cell line was treated with 5 ng/mL TGF-β1 for the indicated times. TGFBIp levels were analyzed by Western blot. (C) Healthy corneal fibroblasts pretreated with 5 ng/mL TGF-β1 for 2 hours were cultured with 10 to 50 mM LiCl for 12 hours. TGFBIp levels were analyzed by Western blot. (D) Quantification of data presented in (C). The levels of TGFBIp were quantified using NIH Image J software and normalized to β-actin. Values represent the mean ± SD of three independent experiments. Statistical analysis was performed using one-way analysis of variance followed by the Newman-Keuls multiple comparison test. *P < 0.05; **P < 0.01 with LiCl versus without LiCl.
immune complex obtained by LiCl treatment compared with the GSK-3 immune complex from no LiCl treatments (Fig. 4B, lane 3). In our hands, interactions between Smad3 and GSK-3 were not different during treatment with 30 and 50 mM LiCl. Together, these results suggest that lithium may regulate the intracellular TGF-β signaling by the interactions of GSK-3 and Smad3 proteins.

**Lithium Induces Autophagy**

Reports have shown that lithium induces autophagy by way of the PI3K signaling pathway.38 Previously, we demonstrated that activation of autophagy enhances the cytosolic clearance of accumulated mut-TGFBIp in primary cultured GCD2 corneal fibroblasts (Choi et al., manuscript submitted). Thus, we investigated whether lithium activates autophagy in corneal fibroblasts. During autophagy, a cytosolic form of the microtubule-associated protein light chain 3-I (LC3-I) is conjugated to phosphatidylethanolamine to form LC3 phosphatidylethanolamine conjugate (LC3-II), which is recruited to autophagosomal membranes. Both the ratio of LC3-II to LC3-I and the amount of LC3-II can be used to monitor autophagosome formation.39,40 Therefore, it was used to monitor autophagy in this study. After treatment with 10, 30, and 50 mM LiCl for 12 hours in corneal fibroblasts, the ratio of LC3-II/LC3-I increased by 1.09 ± 0.1-, 1.60 ± 0.19-, and 2.56 ± 0.24-fold, respectively (P < 0.01), over the control (Figs. 5A, 5B). These results suggest that lithium not only inhibits TGFBIp expression, it also enhances the cytosolic clearance of accumulated mut-TGFBIp in corneal fibroblasts.

**Lithium Is Not Toxic to Primary Cultured Corneal Fibroblasts**

To assess the cytotoxicity of LiCl, primary corneal fibroblasts were incubated with LiCl, and cell viability was measured by the MTT assay. As shown in Figure 6, cell viability was greater...
than 98% after treatment with four different doses of LiCl, indicating that LiCl did not result in significant cell death with the concentrations tested. Taken together, the observed attenuating effects of LiCl on TGFBI expression were not due to corneal fibroblast cell death.

**DISCUSSION**

There is no prevention or cure for TGFBI-linked corneal dystrophy. Thus, there is great hope for the development of novel therapies. To our knowledge, this is the first study describing the impact of lithium on specimens from TGFBI-linked corneal dystrophy patients. Here, we show an effective decrease in TGFBI expression with lithium without cytotoxic effects. Furthermore, we confirmed lithium’s role in the stimulation of autophagy in corneal fibroblasts. These findings strongly suggest that lithium may be used for the prevention or treatment for TGFBI-linked corneal dystrophies.

TGFBI is a secreted protein induced by TGF-β in human adenocarcinoma cells and other human cell types. This protein exhibits tumor suppressive function. We have also shown that TGFBI expression can be induced by TGF-β in a dose-dependent manner and that lithium inhibits this induction in corneal fibroblasts. Thus, these results suggest that lithium may also have a potential role in inhibiting the TGF-β signaling pathway.

Lithium was discovered in 1996 by Klein and Melton as a reversible GSK-3 inhibitor. Studies have established that lithium reduces GSK-3 activity in two ways. As a direct inhibitor, lithium competes with magnesium ions for GSK-3 binding. Alternatively, as an indirect inhibitor, lithium increases the inhibitory N-terminal serine phosphorylation of GSK-3, which inactivates enzyme function. More recently, Millet et al. have suggested that GSK-3 activity negatively regulates the TGF-β signaling pathway. These results suggest that lithium can affect genes regulated by TGF-β signaling. Based on these results, we hypothesized that a GSK-3 inhibitor can be used as a potential therapeutic agent for TGFBI-linked corneal dystrophies. Therefore, we examined the role of lithium on the expression of TGFBI in WT and GCD2 corneal fibroblasts. Here, we have shown that lithium may prevent or treat TGFBI-linked corneal dystrophies by inhibiting TGF-β signaling.

The TGF-β signaling pathway has been implicated as a regulator in numerous cellular and physiological processes, including extracellular matrix homeostasis. This pathway is initiated by the binding of TGF-β to type I (TβRI) and type II TGF-β receptors (TβRII), both of which are serine/threonine kinases. The activated receptor complex phosphorylates the downstream transcription factors Smad2 and Smad3, leading to their association with Smad4. The Smad complex then becomes concentrated in the nucleus and regulates the expression of target genes. These steps are tightly controlled to ensure that Smad2/3 transmits signals from the plasma membrane to the nucleus. Therefore, the decrease in TGFBI expression induced by lithium treatment is consistent with the observed reduction in Smad3 phosphorylation in the presence of the inhibitor (Fig. 3A).

However, although controlling Smad activity is crucial for proper TGF-β signaling and its related factors, the specific mechanism(s) involved in the lithium-induced decrease in Smad3 phosphorylation remains unknown. More recently, Guo et al. have shown that nonactivated Smad3, but not Smad2, undergoes proteasome-dependent degradation caused by the concerted action of the scaffolding protein Axin and its associated kinase, GSK-3β. Smad3 interacts physically with Axin and GSK-3β only in the absence of TGF-β. Reduction in the expression or activity of Axin/GSK-3β leads to increased Smad3 stability and transcriptional activity without affecting TGF-β receptors or Smad2. In contrast, overexpression of these proteins promotes Smad3 basal degradation and desensitizes cells to TGF-β. However, in this study, the mechanism of TGFBI reduction in response to lithium may be mediated by the inhibition of TGF-β signaling through decreased Smad3 phosphorylation without a reduction in total Smad3 protein. In addition, Millet et al. demonstrated that TGF-β induces phosphorylation at three sites within the Smad3 linker region in addition to two C-terminal residues. GSK-3 is responsible for phosphorylating one of these sites, namely Ser204.
we did not assay Smad3 (Ser204) phosphorylation, our results showed decreased Smad3 phosphorylation at S423/425 and increased interaction between Smad3 and GSK-3β in lithium-treated cells. Thus, our findings reveal a novel aspect of Smad3 signaling that controls the final amplitude of TGFBI expression and the cellular responses to TGF-β.

The proteasome and autophagy/lysosomal pathways are the major routes for the clearance of cytotoxic protein. Although the narrow proteasome barrel precludes entry of oligomers/aggregates of aggregate-prone intracellular proteins, such substrates can be degraded by autophagy. We have previously shown that TGFBIp is degraded by the lysosome/autophagy degradation pathway, and its degradation is delayed in GCD2 corneal fibroblasts. Further, the autophagy inducer rapamycin reduced the levels of cytotoxic mutant TGFBIp and attenuated its toxicity in GCD2 homozygous corneal fibroblasts. Furthermore, these data demonstrate that the aggregate-prone form of mutant TGFBIp, unlike the WT protein, is strongly dependent on autophagy for clearance. In addition, we found that mutant TGFBIp accumulates in lysosomes or autolysosomes in GCD2 corneal fibroblasts because of an impairment in autophagy (Choi et al., manuscript submitted). Therefore, these findings suggest that an autophagy inducer can also be used as a therapeutic drug for TGFBI-linked corneal dystrophies in the future.

The autophagy pathway is strongly regulated by levels of IP3, which acts as an endogenous autophagy inhibitor. In contrast, lithium promotes autophagy by blocking IP3 activity. In this study, we found that lithium induces autophagy in corneal fibroblasts. Although we did not determine the levels of decreased TGFBIp after the activation of autophagy with lithium, these findings suggest the therapeutic benefits of lithium in TGFBI-linked corneal dystrophies. In fact, given that lithium did not induce significant death in corneal fibroblasts even at very high concentrations (100 mM), it is highly likely that lithium can be used as a therapeutic agent for treating GCD2.

Studies have also demonstrated that the cytoprotective effects of lithium against oxidative stressors are dependent on the blockage of cytochrome c release and caspase-3 activation. Previously, we showed that oxidative stress is involved in the pathogenesis of GCD2. One possible relationship between GSK-3β activity and oxidative stress has been well described. Oxidative stress may promote degeneration and ultimate cell death through DNA fragmentation, lipid peroxidation, and induction of mitochondrial proapoptotic pathways involving caspases and GSK-3β. In addition, efforts to understand the molecular mechanisms underlying the resistance of cells to hydrogen peroxide-induced oxidative stress have shown that reduced GSK-3β activity may be essential to cell survival. In addition, the treatment of cells with lithium as a GSK-3β inhibitor resulted in increased resistance to hydrogen peroxide-induced oxidative stress. Interestingly, evidence exists showing that GSK-3β inhibition specifically protects cells from intrinsic oxidative stress. Thus, GSK-3β plays an important role in P3K- and Akt-mediated cell survival pathways. Lithium is known to have antiapoptotic effects by reducing the expression of multiple apoptotic proteins, such as p53, Bax, and caspases, and by increasing Bcl-2 levels. Therefore, we also suggest that lithium treatment of TGFBI-linked corneal dystrophies may have multifunctional benefits. Ultimately, lithium treatment may be effective in regulating TGF-β signaling, autophagy, and oxidative stress in numerous diseases.

References


