Transforming Growth Factor Beta 1 Regulates Fibroblast Growth Factor 7 Expression in Schwann Cells

Doan Nguyen, PhD, Olawale A. R. Sulaiman, MD, PhD, FRCS

Background: Our previous work demonstrated that application of transforming growth factor beta 1 (TGF-β1) and forskolin to the repair site after chronic denervation and axotomy has a mitogenic effect, reactivates Schwann cells (SCs), and supports axonal regeneration. We found decreased expression of fibroblast growth factor 7 (FGF-7), a factor involved in synaptic organization and maintenance. Using an in vitro system, we examined the molecular mechanism of TGF-β1 and forskolin on the regulation of FGF-7 expression in SCs.

Methods: SCs were prepared from the sciatic nerve and stimulated with forskolin (0.5 μM), TGF-β1 (1 ng/mL), or TGF-β1 + forskolin for 6 or 24 hours. SCs were also pretreated with LY2109761 (0.5 μM), a TGF-β receptor inhibitor, prior to stimulation with TGF-β1 + forskolin for 6 hours. Real-time TaqMan quantitative polymerase chain reaction analyses for FGF-7, myelin basic protein, and peripheral myelin protein 22 expression were performed. Cycle threshold (Ct) data were normalized to a reference gene, and fold changes relative to untreated SCs were determined using the 2^−ΔΔCt method. Statistical analysis was done using t test (P<0.05).

Results: TGF-β1 alone or in combination with forskolin for 24 hours resulted in a 3.3- and 2.8-fold decrease in FGF-7 expression in SCs, respectively. No change in FGF-7 expression was found with forskolin alone. TGF-β1 + forskolin treatment for 6 hours resulted in a 4.0-fold decrease in FGF-7 expression, while the addition of LY2109761 resulted in a 2.7-fold decrease in FGF-7 expression.

Conclusion: We showed that SC expression of FGF-7 is regulated by TGF-β1. The positive effect of TGF-β1 and forskolin on SC reactivation and axonal regeneration may involve modulation of FGF-7 expression and activity in SCs.

Keywords: Fibroblast growth factor 7, nerve injury, nerve regeneration, Schwann cells, transforming growth factor beta 1

INTRODUCTION

Although peripheral nerves retain the capacity to regenerate, injuries that involve large nerve trunks such as the sciatic plexus and brachial plexus often result in poor and suboptimal functional recovery.1,2 The reason for the lack of functional recovery resides in the lack of sufficient mature axons reaching their targets, a failure attributable to loss of the growth-supportive environment provided by Schwann cells (SCs) in the distal stump of injured nerves.3-5 SCs play a major role during the degenerative process termed Wallerian degeneration and are intimately involved in subsequent regenerative processes. Loss of axonal contact triggers SC transition from a myelinating cell type to a nonmyelinating proliferative cell type. Expression of myelin-associated genes or proteins is downregulated. SCs proliferate in the distal nerve stump to establish cellular channels (bands of Büngner) through which regenerating axons traverse in the distal nerve stump to get to the denervated muscle targets.6 Expression of regeneration-associated genes such as neurotrophins and their receptors7; adhesion molecules8; and cytokines such as transforming growth factor beta 1 (TGF-β1)9 is upregulated in SCs to support the growth of axons through the distal nerve stump. TGF-β1 mediates the recruitment of macrophages into the denervated distal nerve stump, as well as protein mediators that suppress fibroblast proliferation.10,11 However, axonal growth is a slow and turbulent process because axons take many months or even years to reach their targets. The prolonged process leads to a condition of chronic denervation that alters the growth-supportive environment in the distal nerve stump.

Axon outgrowth can be stimulated both in vivo and in vitro by increasing intracellular cyclic adenosine monophosphate (cAMP) levels using forskolin or by blocking phosphodiesterases with rolipram.12-15 In vitro experiments established that 1 ng/mL TGF-β1 and 0.5 μM forskolin...
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were optimal for maximal in vitro stimulation of SC proliferation. Ex vivo experiments demonstrated the benefit of combined TGF-β1 and forskolin on the regenerative effect in chronically denervated nerves as old as 6 months. The mechanism underlying the regenerative effect of TGF-β1 and forskolin on chronically denervated SCs is unknown.

Our previous work using a chronic denervation and axotomy rat model demonstrated that TGF-β1 alone and with forskolin reversed the deleterious effect of chronic SC denervation and reactivated SCs to support axonal regeneration. Previous gene expression profiling identified fibroblast growth factor 7 (FGF-7), also known as keratinocyte growth factor, in the distal nerve stump. FGF-7 has been identified as a secreted factor involved in cell proliferation and tissue repair, as well as in synaptogenesis. The significance of FGF-7 and whether FGF-7 is expressed in SCs is not known. We found that TGF-β1 and forskolin treatment at the site of injury and repair resulted in a 3-fold decrease in FGF-7 expression at the site of repair. This result suggests that the positive effect of TGF-β1 on SC reactivation and axonal regeneration may involve downregulation of FGF-7 expression at the site of repair and distal to the site of repair.

In this study, we used quantitative polymerase chain reaction (qPCR) to confirm the microarray results for FGF-7 expression at the site of repair and in the distal nerve stump. Using an in vitro system, we examined the molecular mechanism of TGF-β1 and forskolin regulation of FGF-7 expression in primary SCs.

METHODS

Chronic Injury and Delayed Repair Model

We used our rat model of tibial (TIB) nerve injury and delayed repair as previously described. All procedures were carried out on Sprague-Dawley (SD) rats and were approved and monitored by the Ochsner Institutional Animal Care and Use Committee. The first surgery involved incision of the rat hindlimb to expose the sciatic nerve, its TIB nerve, and common peroneal branches. The TIB nerve was transected, and its proximal stump was reflected backward and sutured into innervated muscle to prevent regeneration. The second surgery was performed after 8 weeks. The injured TIB nerve was reexposed, and end-to-end repair was done. After microsurgical repair, a 10 mm square piece of Surgifoam (Ethicon) presoaked with saline, forskolin (0.5 μM), TGF-β1 (1 ng/mL), or TGF-β1 + forskolin was applied by wrapping the Surgifoam around the TIB nerve repair site. The animals were allowed to recover, and regeneration of TIB nerves was allowed for 6 weeks. Nerve sections were optimal for maximal in vitro stimulation of SC proliferation. Ex vivo experiments demonstrated the benefit of combined TGF-β1 and forskolin on the regenerative effect in chronically denervated nerves as old as 6 months. The mechanism underlying the regenerative effect of TGF-β1 and forskolin on chronically denervated SCs is unknown.

In this study, we used quantitative polymerase chain reaction (qPCR) to confirm the microarray results for FGF-7 expression at the site of repair and in the distal nerve stump. Using an in vitro system, we examined the molecular mechanism of TGF-β1 and forskolin regulation of FGF-7 expression in primary SCs.

Schwann Cell Culture

Rat SCs were isolated from the contralateral sciatic nerve of adult female SD rats, modified from an established protocol without the use of 2-deoxyuridine to inhibit fibroblast proliferation and a flame-narrowed borosilicate Pasteur pipette to triturate the tissues after enzymatic digestion. The epineurium sheath of the sciatic nerve was removed using fine forceps. The stripped nerve was cut into 1 mm segments, and 10-15 pieces were placed on the central portion of a 35 mm tissue dish containing 750 μL of DMEM/10% fetal bovine serum (FBS) (Atlanta Biologicals, Inc.) plus penicillin-streptomycin (PS). After 1 week, fibroblast outgrowth reached a near confluent monolayer around the nerve explants, and the nerve explants were removed and transferred to a new 35 mm tissue dish. By the fourth explants transfer, cellular outgrowth consisted principally of SCs, and explants were enzymatically dissociated overnight using Dispase II (0.25%, Sigma Aldrich) and collagenase (0.05%, Worthington). The dissociated SCs were plated onto a poly-L-lysine-coated 60 mm tissue dish and maintained in complete DMEM/10% FBS, PS, pituitary extract (20 μg/mL Thermo Fisher Scientific), heregulin (10 ng/mL, BioLegend), forskolin (2 μM, Sigma Aldrich), and ascorbic acid (50 μg/mL, Sigma Aldrich). SCs were subcultured and used between passages 3-6. SCs (5 x 10⁴ cells) were maintained on poly-L-lysine-coated tissue clusters for 6 days, starved for 17 hours, and stimulated with forskolin (0.5 μM), TGF-β1 (1 ng/mL, R&D Systems), or TGF-β1 + forskolin for 24 hours. SCs were also treated with LY2109761 (0.5 μM), a TGF-β receptor type I/II (TβRI/II) dual inhibitor (Selleckchem), for 15 minutes prior to stimulation with TGF-β1 and forskolin for 6 hours or 24 hours.

Real-Time Polymerase Chain Reaction

Total RNA was extracted from SCs using the RNeasy Micro Kit per manufacturer’s instruction (Qiagen) and quantified using a Nanodrop spectrophotometer (Thermo Fisher Scientific). Total RNA (100 ng) was used for complementary DNA (cDNA) synthesis using the iScript cDNA Synthesis Kit (Bio-Rad), and 10 ng cDNA was used for real-time TaqMan qPCR (Thermo Fisher Scientific). Reactions were run in triplicate using the ABI 7500 thermal cycler (Applied Biosystems, Inc.). The cycle threshold (Ct) was determined using the default setting. The Ct of target genes was normalized to the ribosomal RPLS reference gene. Fold changes relative to untreated controls were analyzed using the 2−ΔΔCt method. A ≥2-fold change was used to represent differential gene expression. Statistical significance was analyzed using the t-test (P < 0.05) and expressed as mean ± standard deviation.

Immunohistochemistry

SCs (5 x 10⁴ cells) were plated and maintained at 37°C for 24 hours. SCs were fixed in 4% paraformaldehyde at 4°C for 30 minutes and washed in phosphate buffered saline (PBS, Corning) 3 times for 5 minutes at room temperature (RT). SCs were blocked in 5% normal goat serum PBS for 1 hour at RT, incubated with anti–FGF-7 (Santa Cruz Biotechnology) or anti-S100B (Protein Tech) primary antibody at 4°C overnight and washed in PBS 3 times for 5 minutes at RT. The horseradish peroxidase (HRP) secondary antibody (Enzo Life Sciences) was added and incubated for 30 minutes, washed 3 times in PBS, and incubated for 30 minutes with avidin–biotin complex reagents (Vector Labs). SCs were washed in PBS and stained with 3, 3’-diaminobenzidine (DAB, Vector Labs), hematoxylin and eosin counterstained, and photographed on a Nikon E600 photomicroscope. The parallel control (without the primary antibody) was also analyzed.
The expression of FGF-7 at the site of repair and a 24.2-fold β vs forskolin. Compared to forskolin treatment alone, TGF-
expression was less than the 2-fold cutoff. Expression of TGF-
downward trend after forskolin-only treatment, but the 
downward trend with TGF-
expression was less, with a decrease of 2.7-fold relative to the control (Figure 3). FGF-7 expression levels after 6 hours 
with the dual inhibitor and TGF-β1 + forskolin were similar to 
FGF-7 expression at 24 hours.

LY2109761 had no effect on MBP expression after TGF-β1 + forskolin stimulation, whereas TGF-β1 + forskolin treatment 
decreased PMP-22 expression similar to the effect observed at 24 hours, and PMP-22 expression showed an 
upward trend in the presence of LY2109761.

DISCUSSION
In previous work, we demonstrated the positive effect of TGF-β1 and forskolin treatment on SC reactivation and 
axonal regeneration in the chronically denervated nerve stump.17,19,20 The decreased expression of FGF-7 two 
months after TGF-β1 and forskolin treatment of a chronically 
denervated nerve stump suggests negative regulation of FGF-7 by TGF-β1, and forskolin may be needed to pro-
mote or maintain axonal regeneration. To our knowledge, ours is the first report of FGF-7 expression in the TIB nerve and SCs.

In addition, we showed that TGF-β1 and forskolin decreased MBP and PMP-22 expression, suggesting that 
TGF-β1 and forskolin treatment maintain SCs in a non-
myelinating proliferative state in vivo. TGF-β1 + forskolin 
decreased MBP expression at 24 hours but did not decrease

Figure 1. Immunocytochemical detection of (A) S100B and 
(B) fibroblast growth factor 7 (FGF-7) in primary Schwann 
cells (SCs) isolated from rat sciatic nerves. The positive staining 
indicates that these SCs are S100B-positive and express 
FGF-7 (×10).

RESULTS
TGF-β1 Decreases FGF-7 Expression in the 
Chronically Denervated Nerve Stump
Gene expression profiling of distal nerve stumps of chron-
ically denervated SCs (8 weeks) that were repaired/treated 
with TGF-β1 + forskolin or forskolin alone (6 weeks) 
showed that only TGF-β1 + forskolin treatment resulted in 
a decreased expression of FGF-7. As stated in Methods, 
we used real-time TaqMan qPCR analysis to confirm the 
microarray results. We compared FGF-7 expression at the 
site of repair and distal nerve stump for TGF-β1 + forskolin 
vs forskolin. Compared to forskolin treatment alone, TGF-
β1 + forskolin treatment resulted in a 34.6-fold decrease in 
the expression of FGF-7 at the site of repair and a 24.2-fold decrease in the distal nerve stump.

TGF-β1 Decreases FGF-7 Expression in Schwann 
Cells
Primary culture of rat SCs prepared from a normal sci-
atic nerve displayed a typical bipolar morphology and was immunopositive for S100B, a biomarker of SCs (Figure 1A). 
FGF-7 immunoreactivity was also positive in primary SCs (Figure 1B). To test the effect of TGF-β1 on FGF-7 expres-
sion, SCs were cultured in the presence of TGF-β1 (1 ng/mL), 
forskolin (0.5 μM), TGF-β1 + forskolin, or media alone for 
24 hours. As stated in Methods, the expression of FGF-7 in 
primary SCs was determined by real-time TaqMan qPCR, 
and fold change was analyzed relative to the untreated con-
trol. Expression of FGF-7 significantly decreased 3.3-fold 
with TGF-β1 treatment and 2.8-fold with TGF-β1 + forskolin 
(P<0.05) after 24 hours (Figure 2). Forskolin treatment alone 
also decreased FGF-7 expression, but the decrease did not reach the 2-fold cutoff.

Expression of myelin basic protein (MBP) and peripheral myelin protein 22 (PMP-22), markers for SC differen-
tiation, were also analyzed by real-time TaqMan qPCR. The expres-
sion of MBP decreased 3.3-fold with TGF-β1 treatment and 
5.8-fold with TGF-β1 + forskolin. MBP expression showed a 
downward trend after forskolin-only treatment, but the 
expression was less than the 2-fold cutoff. Expression of 
PMP-22 was not different from the control after forskolin or 
TGF-β1 treatment. However, PMP-22 expression showed a 
downward trend with TGF-β1 + forskolin treatment relative to 
the control.

Figure 2. Fold changes (FC) in expression of fibroblast 
growth factor 7 (FGF-7), myelin basic protein (MBP), and pe-
ripheral myelin protein 22 (PMP-22) relative to untreated 
controls (Con) after 24-hour treatment with forskolin (Fsk), 
transforming growth factor beta 1 (TGFβ), or transforming 
growth factor beta 1 + forskolin (T/F). Expression of FGF-
7 and MBP significantly decreased with TGFβ and T/F treat-
ments (n=3, P<0.05).
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MBP expression at 6 hours. The difference may be that MBP, which is constitutively expressed by SCs in vitro, may require extended treatment longer than 24 hours to be affected by TGF-β1 + forskolin treatment. The negative effect on FGF-7 expression by TGF-β1 appeared to be partially mediated by TβRII activation of the TGF-β1–mediated signaling pathway, as FGF-7 expression was not completely suppressed. TGF-β1 and forskolin, which activate different signaling pathways, may act in an antagonistic way in regulating FGF-7 expression as shown with TβRII inhibition in SCs, and suppression of FGF-7 may be an important aspect of chronic neural repair and regeneration.

The effects of TGF-β on SCs are complex, and outcomes are dependent on the metabolic state as well as the presence or absence of other mitogenic factors. TGF-β acts as a mitogen in SC culture, in the normal milieu of the nerve and activated macrophages, and in neurons. Furthermore, SCs also synthesize and secrete TGF-β. The effects of TGF-β on SCs in the presence of neurons are more of a modulatory role and are in part responsible for the switch from a quiescent myelinating phenotype to a more active proliferative phenotype in the absence of neurons as happens during Wallerian degeneration.

The positive effect of TGF-β1 and forskolin on SC reactivation and axonal regeneration may involve the modulation of FGF-7 expression. FGF-7 is a member of the FGF intercellular signaling molecule involved in a wide range of biologic processes, including cell proliferation, migration, tissue repair, and response to injury. FGF-7 was first identified as a protein secreted from mesenchymal cells to repair epithelial tissues. FGF-7, FGF-10, and FGF-22 are closely related members of the FGF family. These factors have been identified as target-derived factors and as synaptic organizers involved in the formation and maintenance of nerve terminals at synaptic sites. During neuromuscular junction development, TGF-β1 mediates synaptic formation similar to other astrocyte-derived synaptogenic factors and leads to doubling of acetylcholine receptor clusters at the nerve-muscle contact. In vitro studies using SC-conditioned media found the positive effect on synaptogenesis at the neuromuscular junction and showed that TGF-β1 acts as SC-derived synaptogenic factors.

In a study published in 2012, FGF-7 was shown to function as an inhibitory regulator of synaptogenesis, as FGF-7 knockout mice are prone to epileptogenesis. Glial cells, including SCs, not only serve as support cells but are also active participants in modulating presynaptic nerve terminals and postsynaptic specialization. FGF-7 has also been identified in small-diameter neurons of the dorsal root ganglion and localized in large dense-core vesicles in nociceptive neurons. FGF-7 is postulated to exert an excitatory postsynaptic response as FGF-7–deficient mice have reduced acute and inflammatory nociceptive responses.

The molecular mechanism underlying TGF-β1 modulation of FGF-7 expression is not known. Our results are consistent with prior in vitro work in which the combination of TGF-β1 + forskolin added to cultures of chronically denervated isolated SCs for 48 hours was sufficient to increase axon survival when used in vivo as a nerve bridge. Our results are also consistent with the hypothesis that while forskolin is sufficient to activate cAMP functions in SCs and hence mimic the action of various growth factors such as glial cell-derived neurotrophic factor, the effect of TGF-β1 and its modulatory action on other growth factors are important for the full effect to be seen.

As such, understanding the role of FGF-7 in axonal repair and regeneration could provide the basis for development of alternative approach to modulate SC response and synaptic organization.

CONCLUSION

We showed that SC expression of FGF-7 is regulated by TGF-β1. The positive effect of TGF-β1 and forskolin on SC reactivation and axonal regeneration may involve modulation of FGF-7 expression and activity in SCs.

ACKNOWLEDGMENTS

The authors have no financial or proprietary interest in the subject matter of this article. A portion of this work was presented at the Society for Neuroscience Annual Meeting (October 2015, Chicago, IL) and at the Louisiana Neurosurgical Society (January 2016, New Orleans, LA).

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