Treadmill exercise activates ATF3 and ERK1/2 downstream molecules to facilitate axonal regrowth after sciatic nerve injury

Ji-Eun Kim, Yeong-Hyun Cho, Tae-Beom Seo*

Department of Kinesiology, College of Natural Science, Jeju National University, Jeju, Korea

The purpose of this study was to investigate the effect of treadmill exercise on activating transcription factors such as activating transcription factor 3 (ATF3) and extracellular signal-regulated kinase (ERK1/2) signaling pathway to facilitate axonal regrowth after sciatic nerve injury (SNI). The experimental rats divided into the normal control (n = 10), sedentary groups for 7 (n = 10) and 14 days (n = 10) post crush, exercise group for 7 (n = 10) and 14 days (n = 10) post crush (dpc). The rats in exercise groups run on treadmill device at a speed of 8 m/min for 20 min once a day according to exercise duration. In order to evaluate specific regeneration markers and axonal elongation in injured sciatic nerve, we applied immunofluorescence staining and western blot techniques. Treadmill exercise further increased growth-associated protein (GAP-43) expression and axonal regrowth at 7 and 14 dpc than those in sedentary group. Among mitogen-activated protein kinase downstream molecules, phospho-ERK1/2 (p-ERK1/2) was enhanced by treadmill exercise at only 7 dpc and decreased to basal level 14 days later. But c-Jun N-terminal kinase, c-Jun, and phospho-cyclic adenosine monophosphate response element-binding protein showed a tendency to increase continuously until 14 dpc by exercise. ATF3 expression in exercise group was upregulated at both 7 and 14 dpc compared to the sedentary group. These results indicate that treadmill exercise had beneficial effect on expression of regeneration-related proteins after SNI, suggesting that exercise might be one of various therapeutic strategies for sciatic nerve regeneration.

Keywords: Sciatic nerve, Regeneration, Axonal regrowth, Treadmill exercise, Activating transcription factor 3, Mitogen-activated protein kinase

INTRODUCTION

Peripheral nervous system (PNS) axons can voluntary regenerate the injured axons, but central nervous system (CNS) axons cannot. This reason is because Schwann cells in PNS afford essential support by regenerative response of cellular and molecular changes in contrast to the oligodendrocytes and astrocytes in CNS (Gordon, 2016). But some previous studies suggested that spontaneous axonal regrowth after sciatic nerve injury (SNI) not only takes a long time to achieve a satisfied functional recovery but also does not lead to a complete recovery (Park and Hoke, 2014). In general, the distal stump of the injured sciatic nerves undergoes Wallerian degeneration (Arthur-Farraj et al., 2012; Grinsell and Keating, 2014), which Schwann cells and myelin break down and macrophages infiltrate around the injury area.

The regeneration process of the injured sciatic nerve has been well documented in human and animal experiments (Ko et al., 2018; Wanner et al., 2017). Specifically, Schwann cell proliferation is generated at the early stage of the regeneration and proliferating cells migrate toward the distal region to the injury site by releasing neurotrophic factors. Then migrating Schwann cells differentiate for re-myelination. Finally, regeneration is finished by nerve innervation on target muscle (Glenn and Talbot, 2013), representing an important role of Schwann cells in sciatic nerve regeneration.

Schwann cell proliferation after SNI is regulated by genes asso-
related with the cell cycle and cell division, and these cell biological changes facilitate axonal regeneration of the injured peripheral nerves (Jiang et al., 2017). In previous studies on cell proliferation and nerve regeneration, cell division cycle-2 (Cdk2 or Cdk1) promoting G2-M phase transition in the cell cycle has been demonstrated to enhance axonal elongation of the injured sciatic nerve by activating Schwann cell proliferation and neurotrophic factors including brain-derived neurotrophic factor (Han et al., 2007) as well as ERK1/2 are reported to be activated by SNI to enhance Schwann cell proliferation (Seo et al., 2009). In addition, ATF3 or CREB for activating regeneration associated genes (RAGs) are upregulated in cell body of injured peripheral nerve. It has been known that these RAGs are increased by inhibition of free oxygen radicals (Kalender et al., 2009), cell transplantation therapy (Chen et al., 2007) and physical activity (Gómez-Pinilla et al., 2002) after SNI.

Various physical activities including treadmill, swimming, resistance, and wheel running exercise have frequently recommended as a therapeutic modality for improving histological changes and functional recovery after SNI (Armada-da-Silva et al., 2013). According to related studies, treadmill training stimulates proliferation of Schwann cells that support axonal regeneration at early stage of sciatic nerve regeneration (Seo et al., 2009), and physical activity improves the capacity of axonal growth and re-innervation with target muscles (English et al., 2009; Udina et al., 2011a).

With these results mentioned by previous studies, physical activity is essential for axonal regeneration and biochemical alteration after SNI. However the rationale of treadmill exercise on activation of ATF3 and ERK1/2 signaling pathway at early stage of sciatic nerve regeneration is still not clear. Therefore, the purpose of this study was to investigate effect of treadmill training on axonal regrowth at 7 and 14 days after SNI through activation of ATF3 and ERK1/2 downstream molecules.

MATERIALS AND METHODS

Experimental animals

Male Sprague-Dawley rats (160 g, 6 weeks old) were used in this experiment. They were allocated into 5 groups: the normal control (n = 10), sedentary groups for 7 (n = 10) and 14 days (n = 10) postinjury, exercise group for 7 (n = 10) and 14 days (n = 10) post-injury. Animals were maintained at a constant room temperature of 22°C and 60% of humidity under 12/12-hr light-dark cycle. They were accepted to eat commercial rat chow (Samyang Co., Seoul, Korea) and water ad libitum. The experimental procedures must be approved by the ethics committee of Jeju National University and followed animal experiment guide in our institution before this study.

Sciatic nerve injury

For the surgical procedures, the rats were anesthetized using an animal inhalation narcosis control (Jeungdo Bio & Plant, Seoul, Korea). First of all, the rats were placed into the chamber with 2%–2.5% concentration of isoflurane for anesthesia and then 1.5%–1.8% concentration for maintenance during SNI. The left sciatic nerve was crushed with a pair of forceps for 1 min and 30 sec at intervals (Seo et al., 2006). After surgery, anesthetized animals were then placed on a heating pad maintained at 37°C, and then they were put in their cages for resting. All rats were sacrificed 7 and 14 days later.

Treadmill exercise

The experimental rats were adapted to treadmill exercise about 10 min for a week before starting the experiment. Sciatic nerve surgery was performed to exercise and sedentary groups but rats in control group were not crushed. The animals had rest for 2 days after sciatic crush injury. Treadmill exercise was conducted on third postoperative day. The animals in exercise group ran on the treadmill device (Jeungdo Bio & Plant) at 8 m/min for 20 min with no inclination according to exercise period.

Western blot analysis

The dissected sciatic nerves were rinsed with phosphate-buffered saline (PBS), and lysed in Triton lysis buffer. Being denatured proteins were separated on sodium dodecyl sulfate-polyacrylamide gel and then transferred onto polyvinylidene difluoride membrane on ice at 200 mA for 2 hr. The membranes were blocked in TBS including 5% skim milk, 0.1% Tween 20. Then, the membranes were incubated overnight with primary antibodies at 4°C. Protein (20 μg) was used for western blot analysis using anti-GAP-43 mouse monoclonal antibody (1:1,000, Santa Cruz Biotechnology, Santa Cruz, CA, USA), antiphosphorylated ERK1/2 rabbit polyclonal antibody (1:2,000, Cell Signaling Biotechnology, Danvers, MA, USA), antiphosphorylated CREB rabbit monoclonal antibody (1:1,200, Cell Signaling Biotechnology), anti-p38MAPK rabbit polyclonal antibody (1:700, Cell Signaling Biotechnology), anti-ATF3 rabbit polyclonal antibody (1:700, Santa Cruz Biotechnology), antiphosphorylated c-Jun rabbit polyclonal antibody (1:1,200, Cell Signaling Biotechnology), anti-c-Jun N-terminal kinase (JNK) rabbit polyclonal antibody (1:1,200, Cell Signaling Biotechnology).
Biotechnology), anti-β-actin mouse monoclonal antibody (1:2,000, Santa Cruz Biotechnology), and goat anti-mouse or goat anti-rabbit horseradish peroxidase conjugated secondary antibody (1:1,000, GeneTex Inc., Irvine, CA, USA) were used. The blotting proteins were detected by using Westar enhanced cheminescence substrates (Cyanagen, Bologna, Italy). Analysis of protein density was performed using Chemidoc (Bio-Rad, Hercules, CA, USA).

Immunofluorescence staining

Nerve segments were longitudinally sliced on a cryostat at -20°C. For immunofluorescence staining, sections were incubated in PBS with 4% paraformaldehyde and 4% sucrose for 40 min and rinsed 3 times. 0.5% NP-40 in PBS was treated for cell permeability and 2.5% horse serum and 2.5% bovine serum albumin were treated for inhibition of background for 4 hr at room temperature. The following primary antibodies were used for immunostaining: anti-GAP-43 (1:1,000, Santa Cruz Biotechnology) and anti-neurofilament-200 (NF-200) rabbit polyclonal antibody (1:700, Sigma-Aldrich, Darmstadt, Germany) antibodies. Then they were incubated with rhodamine-goat anti-rabbit secondary antibody (1:600, Molecular Probes, Eugene, OR, USA) or fluorescein-goat anti-mouse antibody (1:500, Molecular Probes) for 90 min at room temperature. The sections were viewed with a Nikon fluorescence microscope (Nikon model E-600, Nikon, Kawasaki, Japan), and the images were analyzed using Adobe Photoshop (Adobe CS6, San Jose, CA, USA).

Statistical analysis

All the data is presented as a mean±standard error. Statistical analysis was performed using one-way analysis of variance followed by Duncan post hoc test. The significance level was set at P < 0.05. All graphs were shown by using Prism 6 (GraphPad, La Jolla, CA, USA).

RESULTS

Treadmill exercise increases GAP-43 and axonal elongation in the distal area of the injured sciatic nerve at 7 and 14 dpc

To investigate the effect of treadmill exercise on axonal re-growth, we analyzed time-dependent changes in GAP-43 known as a specific marker for peripheral nerve regeneration. GAP-43 was expressed at 7 and 14 after SNI, and treadmill exercise significantly increased GAP-43 induction at all time points compared to those in the sedentary group (Fig. 1A). At 7 dpc, the immunohistochemistry images in 5 mm distal to the damage site showed increased axonal elongation stained with NF-200. And colocalization of GAP-43 with NF-200 was higher in treadmill group than those seen in the sedentary group (Fig. 1B).

Treadmill exercise upregulates activation of ERK1/2 and CREB signaling pathway at early stage of sciatic nerve regeneration

To examine the possible link between mitogen-activated protein

![Fig. 1. Treadmill exercise increased GAP-43 expression levels and axonal elongation in the injured sciatic nerve. (A) GAP-43 was expressed in the injured sciatic nerve at 7 and 14 days post crush (dpc) and was further upregulated by treadmill exercise. In the quantitative graph, GAP-43 in exercise group (Ex) was significantly enhanced at each time point compared to sedentary group (Sed). (B) Immunofluorescence images of NF-200 and GAP-43 proteins in 5 mm distal to the injury site at 7 dpc. The GAP-43 signals generally overlapped with NF-200-labeled axons, and were higher in Ex than Sed. *P<0.05 vs. the control group. dpc, days post crush; Cont, control; Ex, treadmill exercise; Sed, sedentary.](https://www.e-jer.org/10.12965/jer.2040188.094)
kinase (MAPK) and phospho-CREB (p-CREB), we identified time-dependent alterations in phosphorylated ERK1/2 and CREB in the injured sciatic nerves. As shown in Fig. 2. Phospho-ERK1/2 expression levels were transiently increased at 7 dpc, and then decreased by a basal level at 14 dpc. At 7 dpc, treadmill exercise significantly increased p-ERK1/2 activity compared to those in the sedentary group. Phospho-CREB didn’t show significant differences between control and sedentary groups, but in treadmill exercise group, p-CREB expression levels were significantly increased compared to the sedentary group. Moreover, expression levels of p38MAPK, a negative regulator of Schwann cell differentiation, were significantly decreased in the treadmill exercise group when compared to the sedentary group.

Treadmill exercise increases phosphorylation of ATF3-mediated c-Jun at early stage of sciatic nerve regeneration

To examine proliferating Schwann cell and axonal elongation after SNI, we examined time-dependent changes in ATF3, JNK, and p-c-Jun levels in the damaged sciatic nerves. As shown in Fig. 3. ATF3 expression levels were further increased at only 14 dpc in the sedentary than the control group, but treadmill exercise significantly increased induction levels of ATF3 at 7 and 14 dpc compared to those in the sedentary group. At 14 dpc, JNK protein levels were significantly upregulated in the exercise group. In addition, phosphorylated c-Jun levels showed significant differences in the exercise group than those in the control and sedentary group at 7 and 14 dpc.

**DISCUSSION**

Recent studies have shown that performing regular exercise can improve somatosensory functions after peripheral nerve injury (Martínez de Albornoz et al., 2011; Teodori et al., 2011). But these studies have limitations in reporting the exact biochemical mechanisms of the stimulation caused by physical activity. Our data suggest that GAP-43 protein activated by treadmill exercise may contribute to promoting axonal elongation at 7 and 14 days after SNI. GAP-43 has been known as RAGs accumulated regeneration proteins that stimulate axonal sprouting from proximal to distal stump of the injured sciatic nerve (Moore and Goldberg, 2011). In particular, GAP-43 is upregulated in Schwann cells of the injured motor axons (Vaudano et al., 1995), and physical activity including treadmill exercise not only increases Schwann cell
proliferation at early stage of sciatic nerve regeneration but also upregulates GAP-43 expression levels in proliferated Schwann cells (Udina et al., 2011a; Wilhelm et al., 2012), emphasizing the importance of the timing and duration of exercise for improving sciatic nerve regeneration. The results of these previous studies support our findings that treadmill exercise facilitated regrowth of the injured axons through activation of GAP-43 protein.

It has been known that GAP-43 activation is closely related with MAPKs signaling pathway secreted in regenerating nervous system (Cui et al., 2017; Fan et al., 2017). MAPKs, a family of serine/threonine specific kinases, are a signaling pathway to play a role in various cellular responses ranging from proliferation and differentiation to apoptosis, which is like a double-edged sword. MAPKs consist of three subfamilies such as ERK1/2, JNK, and p38 kinase. In the nerve regeneration field, Zhang et al. (2018) reported that ERK1/2 could regulate phosphorylation of CREB for promoting neurite outgrowth and cell survival in the damaged nerves. In our study, p-ERK1/2 was enhanced by treadmill exercise at only 7 dpc and decreased to normal level 14 dpc. But its downstream molecules including JNK, c-Jun, and p-CREB showed a tendency to increase continuously until 14 dpc by exercise. These data implicate that ERK1/2-regulated Schwann cell proliferation might be associated with axonal regrowth at early stage of regeneration. Seo et al. (2009) demonstrated that treadmill exercise is one of the most therapeutic tools for enhancing cell proliferation and axonal elongation of the injured sciatic nerve via phosphorylation of ERK1/2 protein. In addition, many researchers suggested that the other main MAPK signaling pathways including JNK and c-Jun (Barr and Bogoyevitch, 2001; Davis, 1994) as well as phosphorylation of CREB (Sahatier and English, 2015; Udina et al., 2011b) are activated by peripheral nerve injury and regular exercise after SNI further facilitates these proteins in the damaged Schwann cells compare to those in the sedentary group. Base on the biochemical similarity of MAPK and CREB in nerve regeneration after exercise, we consider that ERK1/2 and CREB signaling pathways contribute to axonal elongation with Schwann cell proliferation.

In addition to activation of MAPK pathway, ATF3 also is a regulator for inducing GAP-43 expression in the injured peripheral nerves (Abe and Cavalli, 2008; Allodi et al., 2012). In general, ATF3 is examined in neurons and Schwann cells after crush or transection of sciatic nerve in rats (Saito and Dahlin, 2008), and it is deeply related with JNK and c-Jun activation. Blom et al. (2014) noted that JNK inhibition by SP600125 and (D)-JNK1 dramati-

![Fig. 3. Treadmill exercise-activated ATF3-mediated c-Jun phosphorylation at early stage of sciatic nerve regeneration. (A) ATF3 protein was increased in exercise group (Ex) compared to those in the sedentary group (Sed) at both 7 and 14 dpc. Treadmill exercise significantly upregulated JNK expression levels at 14 dpc. Also p-c-Jun was further increased in exercise group than those in the sedentary group (Sed). (B) The lower panel shows quantitation of western blot analysis. *P<0.05, **P<0.01, ***P<0.001 vs. the control group. dpc, days post crush; Cont, control; Ex, treadmill exercise; Sed, sedentary.](https://www.e-jer.org)
cally down-regulated ATF3 induction in the injured neurons and JNK-dependent activation of c-Jun is one of a variety of cellular reactions in response to nerve damage, which mean delay of nerve regeneration in injured nerve. Our study observed increased ATF3 expression in exercise group at both 7 and 14 dpc compared to the sedentary group, and ATF3 showed the same pattern of expression in JNK and p-c-Jun proteins in present study. According to the previous studies, it has been suggested that a number of different transcription factors including ATF3, c-Jun, CREB, and Oct-6 were founded in achieving anatomical and functional regeneration after SNI (Patodia and Raivich, 2012), and ATF3 might mediate GAP-43 expression in the damaged Schwann cells. Furthermore, exercise for 1 hr at 50% maximum oxygen uptake transiently increased ATF3 induction in skeletal muscle (Fernández-Verdejo et al., 2017) as well as in nerves (Nakagomi et al., 2003).

Given these findings reported in previous and present studies, physical activity such as treadmill exercise would be important in developing therapeutic strategies for facilitating regeneration-related molecules in the damaged sciatic nerve. In conclusion, the present finding on exercise-activated ATF3 and MAPK pathway during peripheral nerve regeneration should provide a more precise biochemical mechanism on axonal regeneration.

CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

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