Treadmill exercise improves behavioral outcomes and spatial learning memory through up-regulation of reelin signaling pathway in autistic rats

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Autism is a complex neurodevelopmental disability with impairments of social interaction and communication, and repetitive behavior. Reelin is an extracellular glycoprotein that is essential for neuronal migration and brain development. Neuroprotective effects of exercise on various brain insults are well documented, however, the effects of exercise on autism in relation with reelin expression are not clarified. In the present study, we investigated the effects of treadmill exercise on the functional recovery and on the expressions of reelin and its downstream molecules, phosphatidylinositol-3-kinase (PI3K), phosphorylated Akt (p-Akt), phosphorylated extracellular signal-regulated protein kinase 1 and 2 (p-ERK1/2), using autistic rats. For the induction of autism-like animal model, 400 mg/kg valproic acid was subcutaneously injected into the rats on the postnatal day 14. The rat in the treadmill exercise groups were forced to run on a treadmill for 30 min once a day, five times a week for 4 weeks, starting postnatal day 28. To investigate autism-like behaviors and memory deficit, open field, social interaction, and radial 8-arm maze were performed. Immunohistochemistry and western blotting were conducted. In the present results, treadmill exercise alleviated aggressive tendency and improved correct decision in the spatial learning memory in the autistic rats. Treadmill exercise increased neurogenesis and the expressions of reelin and its downstream molecules, PI3K, p-Akt, and p-ERK1/2, in the hippocampus of the autistic rats. The present study showed that treadmill exercise ameliorated aggressive behavior and improved spatial learning memory through activation of reelin signaling pathway in the valproic acid-induced autistic rats.

Keywords: Autism, Valproic acid, Treadmill exercise, Reelin, Hippocampus, Neurogenesis

INTRODUCTION

Autism is a complex neurodevelopmental disability with a variety of symptoms including impairments of social interaction and communication, and repetitive behavior (Fatemi et al., 2005; Vernaza-Martin et al., 2005). Autism is generally diagnosed between the ages of six months and three years old (Rogers, 2009), and it has been hypothesized that autism’s symptoms may be resulted from the maturation-related histological alterations in many areas of brain, and these changes are closely associated with dysregulation of biochemical molecules expressed in the autistic brains (Penn, 2006). Since autism was first recognized as a disorder in 1943, speculation about its etiology has ranged from biological to psychological and back again. However unknown causes account for 80-90% of cases.

Apoptosis is a form of cell death that constitutes part of a common mechanism in cell replacement, tissue remodeling, and removal of damaged cells (Thompson, 1995). Apoptosis has two main pathways (the extrinsic and intrinsic pathways). A class of cysteine proteases, such as caspase-8, caspase-9, and caspase-3, is commonly involved in these pathways (Nuñez et al., 1998). Of these, caspase-3 is the most widely studied member of the caspase
family and it is one of key executors of apoptosis (Cohen, 1997). Excessive neuronal apoptosis contributes to dysfunction of the central nervous system (Baek et al., 2012; Kuhn et al., 2005).

Reelin is a large secreted extracellular matrix glycoprotein that regulates migration and positioning of neurons as well as synaptic plasticity in the developmental brain (Fatemi, 2005; Fatemi et al., 2001). Its activity is regulated by a signaling transduction pathway binding to the two members of very low density lipoprotein receptor (VLDL) and the apolipoprotein E receptor s (ApoER2). VLDL and ApoER2 phosphorylate intracellular adaptor protein Disabled-1 (Dab-1), which facilitates Src-tyrosine kinase family. These changes activate specific molecules such as phosphatidylinositol-3-kinase (PI3K) and protein kinase B (Akt), and extracellular signal-regulated protein kinase 1 and 2 (ERK1/2), then lead to inhibition of neuronal apoptosis in the brain (Beffert et al., 2002, 2005). Fatemi et al. (2001) reported the deficits in the brain levels of reelin mRNA and protein in the subjects with schizophrenia, major depression, and autism.

Exercise enhances learning ability and memory functions, protects from neurodegeneration, delays age-related cognitive decline, and alleviates symptoms of the developmental and neuropsychiatric disorders (Corman et al., 2007; Kim et al., 2010, 2011; Seo et al., 2013). Neuroprotective effects of exercise on various brain insults are well documented (Kim et al., 2011; Lang et al., 2010; Petrus et al., 2008), however, the effects of exercise on autism in relation with reelin expression are not clarified. In the present study, we investigated the effects of treadmill exercise on the functional recovery and on the expressions of reelin and its downstream molecules, PI3K, phosphorylated Akt (p-Akt), phosphorylated ERK1/2 (p-ERK1/2), using valproic acid (VPA)-induced autistic rats.

MATERIALS AND METHODS

Experimental animals and treatments

Male Sprague-Dawley rats, weighing 26±8 g (2 weeks of age) were used for the experiment. The experimental procedures were performed in accordance with the animal care guidelines of the National Institutes of Health (NIH) and the Korean Academy of Medical Sciences. The rats were housed under the controlled temperature (20±2°C), humidity (60%), and the lighting (08:00-20:00 h) conditions. Food and water were made available ad libitum. The animals were randomly assigned to each of four groups (n=15 in each group): the control group, the treadmill exercise group, the VPA-treated group, and the VPA-treated and treadmill exercise group.

For the induction of autism-like animal model, 400 mg/kg valproic acid (Sigma Chemical Company, St. Lois, MO, USA) dissolved in saline at a concentration of 0.1 mL/kg was subcutaneously injected into the rat pups on the postnatal day 14, according to the previous described method (Wagner et al., 2006). Day of birth was recorded as day 0, and all rats were labeled for individual identification. The rats in the control groups received subcutaneous injection of saline in the same volume and on the same schedule.

After starting of treadmill exercise, 5-bromo-2’-deoxyuridine (BrdU) (Sigma Chemical Company) administrated to all animals via intraperitoneal injection (50 mg/kg in saline), once daily for 3 times during a week. The animals were sacrificed immediately after finishing behavior tests.

Treadmill exercise protocol

The rat in the treadmill exercise groups were forced to run on a treadmill for 30 min once a day, five times a week for 4 weeks, starting postnatal day 28. Exercise load for the exercise groups consisted of running at a speed of 2 meters/min for the first 5 min, at a speed of 5 m/min for the next 5 min, and then at a speed of 8 m/min for the last 20 min, with the 0’ inclination.

Open field test

For open field test, the animals were randomly assigned to an order of testing and placed in a white square open field arena (100 cm×100 cm) made of wood. As previously reported (Kim et al., 2011), it was enclosed with 40-cm high walls and placed under strong illumination (200 lux). The arena was divided into 25 squares (each square is 20×20 cm), defined as 9 central and 16 peripheral squares. The animal was placed in the center of the arena and left free to explore the environment for 1 min. After that time, the numbers of squares that the rat crossed were recorded for 5 min.

Social interaction test

The social interaction test was conducted in a 120×120×60 cm black wooden arena illuminated by a 40 W red lamp. Age- and treatment-matched pairs of rats with approximately equal body weight, but from different home cages, were placed together in the arena for 20 min. Social behaviors include: (1) following/chasing: one rat following the other within a distance of two body-lengths, (2) anogenital interactions, (3) adjacent interactions: including play-fighting, climbing over/under, and adjacent lying, (4) head-to-head interactions, and (5) total social interaction: time
spent in all of the above categories. The apparatus was cleaned with 50% ethanol between test sessions.

**Radial 8-arm maze test**

Spatial learning memory was tested using a radial 8-arm maze apparatus, as the previously described method (Kim et al., 2010). The radial 8-arm maze apparatus consisted of a central octagonal plate (30 cm in diameter) and radiating eight arms (50 cm in length and 10 cm in width). The apparatus was placed 1 m above the floor. A small receptacle filled with water (3 cm in diameter and 1 cm in depth) was located at the end of the arms. The rats were trained three times before the spatial learning test. In the training sessions, the rats deprived of water for 24 h were allowed to explore the water and to drink for 5 min. The time spent in seeking water at the end of the arms was counted. The test was terminated when a rat found water in all eight arms or over 6 min elapsed. The number of correct choice before the first error was counted. Re-entering to the previously visited arms was counted as the error.

**Tissue preparation**

The experimental animals were fully anesthetized using Zoletil 50® (10 mg/kg, i.p.; Vibac Laboratories, Carros, France), transcardially perfused with 50 mM phosphate-buffered saline (PBS), and fixed with a freshly prepared solution consisting of 4% paraformaldehyde in 100 mM phosphate buffer (PB, pH 7.4). Brains were dissected, post-fixed in the same fixative overnight, and transferred to 30% sucrose for cryoprotection. Sagittal sections of 40 μm thickness in each section of cerebellum were made with a freezing microtome (Licea, Nussloch, Germany). The sections were finally mounted onto gelatin-coated slides. The slides were air-dried at 65°C for 2 h, denaturated in 2N HCl at 37°C for 30 min, and they were rinsed twice in 100 mM sodium borate (pH 8.5). Afterwards, the sections were incubated overnight at 4°C with BrdU-specific mouse monoclonal antibody (1:600; Roche, Mannheim, Germany). The sections were then washed three times with PBS and incubated for 1 h with the biotinylated mouse secondary antibody (1:200; Vector Laboratories, Burlingame, CA, USA). Then, the sections were incubated for another 1 h with avidin-peroxidase complex (1:100; Vector Laboratories). For visualization, the sections were incubated for 5 min in 50 mM Tris-HCl (pH 7.6) containing 0.02% 3,3’-diaminobenzidine (DAB), 40 mg/ml nickel chloride, and 0.03% hydrogen peroxide. The slides were air-dried overnight at room temperature, and the coverslips were mounted using Permount® (Fisher Scientific, New Jersey, NJ, USA).

**Caspase-3 immunohistochemistry**

Caspase-3 immunohistochemistry was performed, according to a previously described method (Kim et al., 2010). In brief, the sections were incubated overnight with mouse anti-caspase-3 antibody (1:500; Santa Cruz Biotechnology, Santa Cruz, CA, USA) and then they were incubated for another 1 h with the biotinylated mouse secondary antibody. The bound secondary antibody was then amplified using a Vector Elite ABC kit® (1:100; Vector Laboratories). The antibody-biotin-avidin-peroxidase complex was visualized using 0.03% DAB. The sections were finally mounted onto gelatin-coated slides. The slides were air-dried overnight at room temperature, and the coverslips were mounted using Permount® (Fisher Scientific).

**Western blot analysis**

Western blotting was conducted a previously described method (Kim et al., 2010). Tissue samples harvested from the hippocampus were lysed in the protein lysis buffer containing 50 mM HEPEs (pH 7.5), 150 mM NaCl, 10% glycerol, 1% Triton X-100, 1.5 mM magnesium chloride hexahydrate, 1 mM phenylmethylsulfonyl fluoride (PMSF), 2 mg/mL leupeptin, 1 mg/mL pepstatin, 1 mM sodium ortho vanadate, and 100 mM sodium fluoride. Protein concentration was measured using a Bio-Rad colorimetric protein assay kit (Hercules, CA, USA). Protein of 40 μg was separated on SDS-polyacrylamide gels and transferred onto a nitrocellulose membrane (Schleicher & Schuell GmbH, Dassel, Germany). Mouse anti-reelin antibody (1:1,000; Millipore Corporation, Billerica, MA, USA), mouse anti-PI3K antibody (1:1,000; Santa Cruz Biotechnology), rabbit anti-p-Akt antibody (1:2,000; Cell Signaling Technology, Beverly, MA, USA), rabbit anti-total Akt antibody (1:1,000; Cell Signaling Technology), mouse anti-p-ERK1/2 antibody (1:1,000; Santa Cruz Biotechnology), and rabbit anti-total ERK1/2 antibody (1:2,000; Cell Signaling Technology) used as primary antibodies. Horseradish peroxidase-conjugated anti-mouse antibody (1:3,000; Santa Cruz Biotechnology)
for actin, and horseradish peroxidase-conjugated anti-rabbit antibody (1:4,000; Vector Laboratories) or horseradish peroxidase-conjugated anti-mouse antibody (1:3000; Vector Laboratories) used as secondary antibodies. Band detection was performed using as enhanced chemiluminescence (ECL) detection system (Amersham Pharmacia Biotech GmbH, Freiburg, Germany).

**Statistical analysis**

The numbers of caspase-3-positive and BrdU-positive cells in the hippocampal dentate gyrus were counted using confocal laser-scanning microscopy with LSM 510 META (Carl Zeiss, Oberkochen, Germany). The numbers of caspase-3-positive and BrdU-positive cells in the hippocampal dentate gyrus were expressed as the number of cells per mm² of the cross sectional area of the granular layer. The final amount of western blot product for protein expression was calculated densitometrically using Imaging-Pro® Plus (Media Cybernetics Inc., Silver Spring, MD, USA).

All data were analyzed using the statistical software SPSS (version 18.0). The data are expressed as the mean±standard error of the mean (SEM). For the comparison among the groups, one-way analysis of variance (ANOVA) and Duncan’s post-hoc test were performed and differences were considered statistically significant at \( P < 0.05 \).

**RESULT**

**Expression of caspase-3 in the hippocampus one day after VPA treatment**

One day after VPA treatment, caspase-3 expression was detected using caspase-3 immunohistochemistry. The number of caspase-3-positive cells in the hippocampal dentate gyrus was 7.4 ± 1.3 in the control group and 48.7 ± 3.1 in the one day after VPA-treated group (\( P < 0.05 \)). Many caspased-3-labeled cells were observed in the VPA-treated rats compared to the control group. The present results indicate that VPA injection induced apoptotic cell death in the hippocampus.

**Effect of treadmill exercise on the activity in the open field test and on the behaviors in the social interaction test**

The effect of treadmill exercise on the activity in the open field test is presented in Fig. 1A. The activity level was 85.1 ± 7.4 in the control group, 97.5 ± 9.6 in the control and treadmill exercise group, 53.6 ± 10.4 in the VPA-treated group, and 67.3 ± 11.1 in the VPA-treated and treadmill exercise group. The present results indicated that the activity in the VPA-treated group was decreased compared to the control group (\( P < 0.05 \)). However, treadmill exercise did not significantly increase the activity in the VPA-in-
jected rats ($P > 0.05$).

The effect of treadmill exercise on the behaviors in the social interaction test is presented in Fig. 1B, C. The number of non-aggressive behavior was $20.0 \pm 2.7$ in the control group, $19.2 \pm 1.8$ in the control and treadmill exercise group, $10.4 \pm 1.9$ in the VPA-treated group, and $14.2 \pm 0.9$ in the VPA-treated and treadmill exercise group. The number of aggressive behavior was $3.8 \pm 0.7$ time in the control group, $5.8 \pm 1.1$ time in the control and treadmill exercise group, $16.8 \pm 1.6$ time in the VPA-treated group, and $10.2 \pm 2.9$ time in the VPA-treated and treadmill exercise group. The present results indicated that non-aggressive behavior was decreased and aggressive behavior was increased in the VPA-treated group compared to the control group ($P < 0.05$). Treadmill exercise did not enhance non-aggressive behavior ($P > 0.05$), however, treadmill exercise suppressed aggressive behavior in the VPA-injected rats ($P < 0.05$).

**Effect of treadmill exercise on the spatial learning memory in the radial 8-arm maze test**

The effect of treadmill exercise on spatial learning memory in the radial 8-arm maze test is presented in Fig. 2. The time taken to complete eight successful performances was $249.3 \pm 30.6$ sec in the control group, $194.9 \pm 15.4$ sec in the control and treadmill exercise group, $368.6 \pm 24.9$ sec in the VPA-treated group, $294.1 \pm 47.2$ sec in the VPA-treated and treadmill exercise group. The number of correct choices to complete eight successful performances was $5.2 \pm 0.5$ in the control group, $7.2 \pm 0.1$ in the control and treadmill exercise group, $3.0 \pm 0.4$ in the VPA-treated group, and $4.7 \pm 1.9$ in the VPA-treated and treadmill exercise group. The number of error to complete eight successful performances was $4.6 \pm 0.5$ in the control group, $2.7 \pm 0.7$ in the control and treadmill exercise group, $6.0 \pm 1.4$ in the VPA-treated group, and $5.1 \pm 0.8$ in the VPA-treated and treadmill exercise group. The present result indicated that time and error number were increased and correct number was decreased in the VPA-treated group compared to the control group ($P < 0.05$). Treadmill exercise did not decrease time and error number ($P > 0.05$), however, treadmill exercise increased correct number in the VPA-injected rats ($P < 0.05$).

**Effect of treadmill exercise on the hippocampal neurogenesis**

The effect of treadmill exercise on the neurogenesis in the hippocampal dentate gyrus is presented in Fig. 3. The number of BrdU-positive cells in the hippocampal dentate gyrus was $198.5 \pm 8.8$ in the control group, $214.1 \pm 6.9$ in the control and treadmill exercise group, $62.6 \pm 5.4$ in the VPA-treated group, and $102.3 \pm 4.6$ in the VPA-treated and treadmill exercise group. The
present result indicated that neurogenesis in the hippocampal dentate gyrus was lower in the VPA-treated group compared to the control group (P < 0.05). In contrast, treadmill exercise increased neurogenesis in the VPA-injected rats (P < 0.05).

**Effect of treadmill exercise on the co-localization of reelin in the hippocampus**

Reelin protein was expressed in the CA1, CA3, hilus, and dentate gyrus in the hippocampus. Co-localization of reelin with BrdU-positive cells was observed in the hippocampal dentate gyrus (Fig. 4). Reelin expression in the hippocampus was lower in the VPA-treated group compared to the control group. Treadmill exercise increased reelin expressions in the VPA-injected rats.

**Effect of treadmill exercise on the expressions of reelin and its down-stream molecules in the hippocampus**

BrdU-positive neurons were co-localized with reelin in the dentate gyrus of the hippocampus (Fig. 5, upper). The effect of treadmill exercise on the expressions of reelin and its down-stream molecules, PI3K, p-Akt, and p-ERK1/2, is presented in Fig. 5 (lower). Band intensity of each protein expressed in control group was set at 1.00.

When the level of reelin in the control group was set at 1.0 ± 0, the level of reelin was 1.1 ± 0.02 in the control and treadmill exercise group, 0.3 ± 0.04 in the VPA-treated group, and 0.5 ± 0.06 in the VPA-treated and treadmill exercise group. When the level of PI3K in the control group was set at 1.0 ± 0, the level of PI3K was 1.7 ± 0.15 in the control and treadmill exercise group, 0.4 ± 0.04 in the VPA-treated group, and 0.7 ± 0.06 in the VPA-treated and treadmill exercise group. When the level of p-Akt in the control group was set at 1.0 ± 0, the level of p-Akt was 1.3 ± 0.10 in the control and treadmill exercise group, 0.5 ± 0.04 in the VPA-treated group, and 0.9 ± 0.04 in the VPA-treated and treadmill exercise group.

When the level of phosphorylated (p-ERK1/2) in the control group was set at 1.0 ± 0, the level of p-ERK1/2 was 1.9 ± 0.25 in the control and treadmill exercise group, 0.2 ± 0.03 in the VPA-treated group, and 0.6 ± 0.06 in the VPA-treated and treadmill exercise group. The present result indicated that the expressions of reelin and its down-stream molecules, PI3K, p-Akt, and p-ERK1/2, were decreased in the VPA-treated group compared to the control group (P < 0.05). In contrast, treadmill exercise increased the expressions of reelin and its down-stream molecules in the VPA-injected rats (P < 0.05).

**DISCUSSION**

In the present study, we used 400 mg/kg of VPA to induce au-

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**Fig. 3.** Effect of treadmill exercise on the neurogenesis in the dentate gyrus of the hippocampus. (A) Photomicrographys of neuronal nuclei and 5-bromo-2’-deoxyuridine (BrdU)-positive cells. The scale bar represents 500 μm. (B) The number of BrdU-positive cells. (A) Control group, (B) control and exercise group, (C) valproic acid (VPA)-treated group, (D) VPA-treated and treadmill exercise group. The data are represented as the mean ± SEM. Symbols a, b, c represent statistical significance (P<0.05) in each group.
Autistic symptoms in the rats on the postnatal day 14, and we observed caspase-3 activation in the hippocampal dentate gyrus one day after VPA exposure. VPA is primarily used for the treatment of epilepsy, major depression, and bipolar disorder. However, exposure of VPA during pregnancy results in a fetal valproate syndrome similar to autism (Arnon et al., 2000; Mawer et al., 2002). Administration of VPA to the rats on the postnatal day 14 showed autism-like symptoms including impairment of social and cognition and restricted-repetitive behavior (Yochum et al., 2008). Many previous studies reported that less than 500 mg/kg VPA injection induced severe autism-like state without death of experimental animals (Markram et al., 2008; Yochum et al., 2010). In the present results, the dose of 400 mg/kg valproic acid was relevant to the inducing autism in the rats.

In the present results, treadmill exercise suppressed VPA-induced aggressive behavior, in contrast treadmill exercise exerted no significant effect on VPA-induced hyperactivity and non-aggressive behavior. The present results also showed that treadmill exercise alleviated VPA-induced decrement of correct number, in contrast, treadmill exercise exerted no significant effect on time and error number. Our preset study revealed that treadmill exercise alleviated aggressive tendency and improved correct decision in the spatial learning memory in the autistic rats.

It is well documented that physical exercise enhances cognitive function, learning ability, and memory function (Baek et al., 2012; Brisswalter et al., 2002; Kim et al., 2010, 2012). Exercise amelio-

Fig. 4. Photomicrographs of reelin expressions in the CA1, CA3, hilus, and dentate gyrus of the hippocampus. The scale bar represents 100 μm. (A) Control group, (B) control and exercise group, (C) valproic acid (VPA)-treated group, (D) VPA-treated and treadmill exercise group. Enlarged images are to magnify the asterisk area in photo A. The scale bar represents 100 μm.
rates memory deficits, improves social behavior problems, and relieves hyperactivity in autism (Allison et al., 1991; Bahrami et al., 2012; Petrus et al., 2008). Following bouts of physical activity, children with autism experienced the decrement in negative behaviors, such as stereotypy, and the increment in positive behaviors, such as time on task (Levinson and Reid, 1993; Lang et al., 2010). Petrus et al. (2008) suggested that exercise provides reduction of stereotypic behaviors in the children with autism spectrum disorder. Children with autism are more inactive in physical activity compared to their peers without autism (Macdonald et al., 2011).

In the present study, we carried out BrdU assay to investigate the level of neurogenesis in the hippocampus of autistic rats. The present results showed that neurogenesis in the hippocampal dentate gyrus was decreased in the autistic rats, in contrast, treadmill exercise increased neurogenesis in the autistic rats.

In the present results, reelin expression in the hippocampus was suppressed in the autistic rats, in contrast, treadmill exercise enhanced reelin expression in the autistic rats.

Several factors such as schizophrenia-1 (DISC-1), cyclin-dependent kinase 5 (cdk5), NeuroD, Prox1, and reelin are implicated in the controlling of neurogenesis in the adult dentate granule cells (Gao et al., 2009; Karalay et al., 2011). Of them, reelin is essential for neuronal survival and differentiation. Reelin is an extracel-
lular protein that is essential for neuronal migration and brain development (Fatemi et al., 2001; Fatemi, 2005). Dysregulation of reelin may be responsible for some of the brain structural and behavioral abnormalities observed in autism (Fatemi et al., 2001). In the adult brain, reelin is involved in the signaling pathway which underlies neurotransmission, memory formation, and synaptic plasticity (Fatemi, 2005). The activation of reelin signaling may inhibit excitotoxic neurotransmission and Tau phosphorylation, and may activate neurogenesis. This may lead to diminished brain injury and to increased brain injury repair (Delahaye et al., 2007).

In the present study, we confirmed the changes of down-stream molecules of the reelin signaling pathway in the autistic rats. Expressions of PI3K, p-Akt, and p-ERK1/2 in the hippocampus were decreased in the autistic rats, in contrast, treadmill exercise increased the expression of PI3K, p-Akt, and p-ERK1/2 in the autistic rats.

PI3K-Akt signaling pathway is under the control of reelin signaling (Jossin and Goffinet, 2007; Waite and Eickholt, 2010). The relation of PI3K-Akt pathway with autism spectrum disorders has been suggested. Kwon et al. (2006) reported that abnormal activation of the PI3K-Akt pathway might be the one of the underlying mechanisms of macrocephaly and behavioral abnormalities in human autism spectrum disorders. PI3K signaling pathway is involved in the development of the brain, and aberrant PI3K signaling is an important contributing factor to the pathophysiology of disorders such as schizophrenia and autism (Waite and Eickholt, 2010). Weaken PI3K-Akt-Bcl-xL results in an augmented cell death, in contrast, activation of PI3K-Akt signaling pathway increases cell survival (Jeon et al., 2011).

In the present study, treadmill exercise ameliorated aggressive behavior and improved spatial learning memory through activation of reeling signaling pathway in the VPA-induced autistic rats. The present study suggests the possibility that exercise can be applied for the symptom relief of autistic patients.

**CONFLICT OF INTEREST**

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

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