Treadmill exercise alleviates depressive symptoms in rotenone-induced Parkinson disease rats

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Parkinson disease (PD) is characterized by selective loss of the dopaminergic neurons. The symptoms of depression following PD are closely associated with reduced activity of the serotonergic system in the dorsal raphe. We explored the anti-depressive effect of exercise and its possible mechanism using the rotenone-induced PD rats. PD rats were induced by subcutaneously injection with rotenone for 14 days. The rats in the exercise groups were made to run on a treadmill for 30 min once a day during 14 consecutive days. Forced swimming test, immunohistochemistry for serotonin (5-hydroxytryptamine, 5-HT), tryptophan hydroxylase (TPH), and western blot for serotonin _{1A} (5-HT_{1A}) receptor were conducted. Injection of rotenone induced PD rats. PD rats showed depressive state and treadmill exercise ameliorated this depressive state.

5-HT, TPH, and 5-HT_{1A} receptor expressions in the dorsal raphe were suppressed by rotenone injection and treadmill exercise increased the expressions of 5-HT, TPH, and 5-HT_{1A} receptor in the rotenone-injected rats. The present results show that treadmill exercise ameliorated depressive symptoms in the rotenone-induced PD rats. The antidepressive effect of treadmill exercise might be ascribed to the enhancement of serotonergic function through upregulation of 5-HT_{1A} expression in the dorsal raphe.

Keywords: Parkinson disease, Treadmill exercise, Rotenone, Serotonin, Serotonin _{1A} receptor

INTRODUCTION

Parkinson disease (PD) is characterized by selective loss of the dopaminergic neurons in the substantia nigra pars compacta and striatum. Intracytoplasmic inclusions, called Lewy bodies, in the remaining dopaminergic neurons is the hallmark of pathophysiology of PD, and resting tremor, rigidity, bradykinesia, and gait disturbance are the characteristics of PD patients (Blesa et al., 2012; Varçin et al., 2012). In addition to motor dysfunction, depression is the most frequent psychiatric complication in PD patients, and depression is one of the most important factors reducing quality of life of PD patients (Aarsland et al., 2011).

Depression is associated with functional impairment of serotonergic neurotransmitter system, such as serotonin (5-hydroxy-

tryptamine, 5-HT), tryptophan hydroxylase (TPH), and 5-HT transporters, which are involved in the regulation of mood (Christiansen et al., 2007). Of these, 5-HT is an important neurotransmitter in the many neuropsychiatric disorders, such as schizophrenia, depression, anxiety neuromodulator in the mammalian brain (Graeff, 1997; Lucki, 1998). As such, The *TPH* gene, coding for the rate-limiting enzyme in the biosynthesis of 5-HT, might be involved in pathogenesis events involving dysfunction of the 5-HT system. Thus, TPH is one of the major candidate genes for psychiatric and behavioral disorders, in particular, depression (Roy et al., 1999). Tryptophan depletion also increases symptom severity in depression patients through down-regulation of 5-HT biosynthesis (Neumeister et al., 2004; Ruhé et al., 2007). The symptoms of depression following PD are closely associated with re-

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duced activity of the serotonergic system (Kish et al., 2008). Reduction of serotonin and its metabolites was reported in brains of PD patients (Kish et al., 2008). Serotonergic neurons are closely correlated with depression (Brook and Pavese, 2011).

5-HT receptors are divided into 14 subtypes (Barnes and Sharp, 1999). Of receptor subtypes, serotonin $1A$ (5-HT $_{1A}$) receptor, a G protein-coupled receptor, plays a major role in the process of psychiatric disorders (Savitz et al., 2009). Positive correlation was appeared between depression and binding to the serotonin transporter and 5-HT $_{1A}$ receptor (Boileau et al., 2008; Brooks and Pavese, 2011). 5-HT $_{1A}$ receptor dysfunction causes depression (Leitch et al., 2003), and 5-HT $_{1A}$ receptor is implicated in the progression of psychiatric disorders (Savitz et al., 2009).

Exercise is known to exert antidepressive effect on depression patients (Strawbridge et al., 2002) as well as on animal model of depression (Shin et al., 2017b). The purpose of this study is to explore the antidepressive effect of treadmill exercise and its possible mechanism using the rotenone-induced PD rats.

MATERIALS AND METHODS

Animals and treatments

Male Sprague-Dawley rats, weighing 250 ± 10 g (9 weeks of age), were used in this experiment. They were kept in a room with controlled temperature ($20^\circ\text{C} \pm 2^\circ\text{C}$) under a 12-hr light/12-hr dark cycle. The animals were allowed free access to food and water. All animal experimental procedures conformed to the regulations stipulated by the National Institutes of Health and the guidelines of the Korean Academy of Medical Science. The animals were randomly divided into four groups ($n = 10$ in each group): the control group, the control and treadmill exercise group, the rotenone-injected group, and the rotenone-injected and treadmill exercise group.

PD animal model

PD rats were induced, according to the previously described method (Shin et al., 2017a). Rotenone was purchased from Sigma Chemical Co. (St. Louis, MO, USA). The rats in the rotenone-injected groups were injected subcutaneously with rotenone (3.0 mg/kg) for 14 days, dissolved into dimethyl sulfoxide. The rats in the control group were injected subcutaneously with dimethyl sulfoxide daily for the same duration.

Exercise protocol

The animals in the exercise groups were forced to run on a mo-

torized treadmill for 30 min once a day during 14 consecutive days. The exercise load consisted of running at a speed of 2 m/min for the first 5 min, 3 m/min for the next 5 min, and then 4 m/min for the last 20 min.

Forced swimming test

In order to evaluate depression severity, we performed a modified forced swimming test (FST), according to the previously described method (Shin et al., 2017b). After cessation of exercise, the rats conducted a pre-test for 15 min to eliminate the acute stress by water and to adapt the animals to the water. Twenty-four hr after the pre-test, the rats were tested for 6 min. The animals were placed individually into the glass cylinder 20 cm in diameter and 90 cm in height, filled with water to a height of 75 cm. The temperature of the water was adjusted to $25^\circ\text{C} \pm 1^\circ\text{C}$. During the test session, the climbing time and immobility time were analyzed using a Smart version 2.5 video tracking system (Panlab, Barcelona, Spain). Immobility was defined to occur when no additional activity was observed other than the actions needed to keep the rat's head above the water. Climbing was defined to occur when the rat was in an active vertical motion with its forelegs above the water level.

Tissue preparation

After FST, the rats were deeply anesthetized with Zoletil 50 anesthesia (1 mg/kg, intraperitoneally; Vibac Laboratories, Carros, France). After complete anesthesia, the rats were transcardially perfused with 0.05-M phosphate-buffered saline (PBS), followed by 4% paraformaldehyde in 0.5-M sodium phosphate buffer at pH 7.4. The brain was removed, postfixed in the same fixative overnight, and transferred to a 30% sucrose solution for cryoprotection. Serial 40- μm -thick coronal sections were cut with a freezing microtome (Leica, Nussloch, Germany).

Immunohistochemistry for 5-HT and TPH

Immunohistochemistry for 5-HT and TPH was performed, according to the previously described method (Shin et al., 2017b). An average of eight sections was selected in each brain region spanning from Bregma -7.30 to -8.00 mm. The sections were incubated in PBS for 10 min and they were next washed three times with PBS. The sections were then incubated in 1% H_2O_2 for 30 min, and then they were incubated overnight with rabbit anti-5-HT antibody (Oncogene Research Product, Cambridge, UK) at a dilution of 1:500 or with mouse anti-TPH antibody (Oncogene Research Product) at a dilution of 1:500. The sections were incu-

bated for 1 hr with biotinylated anti-rabbit secondary antibody or with anti-mouse secondary antibody (Vector Laboratories, Burlingame, CA, USA), and they were subsequently incubated with avidin–biotin–peroxidase complex (Vector Laboratories) for 1 hr at room temperature. Immunoreactivity was visualized by incubating the sections in a solution consisting of 0.05% 3,3'-diaminobenzidine and 0.01% H₂O₂ in 50-mM Tris-buffer (pH, 7.6) for approximately 3 min. The sections were finally mounted on gelatin-coated glass slides. The slides were air-dried overnight at room temperature, and the coverslips were mounted using Permount (Thermo Fisher Scientific Inc., Waltham, MA, USA).

Western blot analysis

Western analysis was performed, according to the previously described method (Shin et al., 2017b). Dorsal raphe tissues were dissected. Sample tissues were stored at -70°C until analysis. Sample tissues were lysed in ice-cold lysate 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 ethyleneglycol-bis-(β -aminoethyl ether)-N,N'-tetraacetic acid, 1-mM phenylmethylsulfonyl fluoride, 2- μ g/mL leupeptin, 1- μ g/mL pepstatin, 1-mM sodium orthovanadate, and 100-mM sodium fluoride, after which the mixture was incubated for 30 min at 4°C. The tissues debris was then removed by microcentrifugation followed by quick freezing of the supernatant. The protein concentration was measured using a Bio-Rad colorimetric protein assay kit (Bio-Rad, Hercules, CA, USA). Protein of 30 μ g was separated on sodium dodecyl sulfate-polyacrylamide gels and transferred onto a nitrocellulose membrane (Whatman, Clifton, NJ, USA). Rabbit anti-5-HT_{1A} antibody (1:1,000; Abcam, Cambridge, UK) were used as a primary antibody. Horseradish peroxidase-conjugated anti-rabbit antibody (1:3,000; Vector Laboratories) was used to probe for 5-HT_{1A}. Band detection was performed using the enhanced chemiluminescence detection system (Santa Cruz Biotechnology, Santa Cruz, CA, USA). The bands were quantified using an Image-Pro Plus computer-assisted image analysis system (Media Cybernetics Inc., Bethesda, MD, USA).

Statistical analysis

All data were analyzed using IBM SPSS Statistics ver. 23.0 (IBM Co., Armonk, NY, USA). The data were expressed as the mean $\pm$ standard error of the mean. For the comparison among the groups, One-way analysis of variance and Duncan *post hoc* test were performed with $P < 0.05$ as an indication of statistical significance.

Fig. 1. Effect of treadmill exercise on immobility and climbing time in the forced swimming test. (A) Immobility time. (B) climbing time. CON, control group; CON-EX, control and treadmill exercise group; ROTE, rotenone-injected group; ROTE-EX, rotenone injected and treadmill exercise group. Data are presented as the means $\pm$ standard error of the mean. * $P < 0.05$ compared to the control group. # $P < 0.05$ compared to the rotenone-injected group.

RESULTS

Effect of treadmill exercise on immobility and climbing time in the forced swim test

The immobility time and climbing time in the FST are presented in Fig. 1. The immobility time was increased by injection of rotenone ($P < 0.05$) and treadmill exercise decreased immobility time in the rotenone-injected rats ($P < 0.05$). The climbing time was decreased by injection of rotenone ($P < 0.05$) and treadmill exercise increased climbing time in the rotenone-injected rats ($P < 0.05$).

Effect of treadmill exercise on 5-HT expression in the dorsal raphe

Photomicrographs of 5-HT-positive cells in the dorsal raphe are presented in Fig. 2. The number of TPH-positive cells in the dorsal raphe was 287.10 ± 18.37 /section in the control group,

Fig. 2. Photomicrographs showing 5-hydroxytryptamine (5-HT) expressions in the dorsal raphe. The scale bar is 100 μ m. CON, control group; CON-EX, control and treadmill exercise group; ROTEX, rotenone-injected group; ROTEX, rotenone injected and treadmill exercise group.

285.30 $\pm$ 16.89/section in the control and treadmill exercise group, 217.40 $\pm$ 18.55/section in the rotenone-injected group, 283.10 $\pm$ 14.80/section in the rotenone-injected and treadmill exercise group. 5-HT expression in the dorsal raphe was decreased by injection of rotenone ($P < 0.05$) and treadmill exercise increased 5-HT expression in the rotenone-injected rats ($P < 0.05$).

Effect of treadmill exercise on TPH expression in the dorsal raphe

Photomicrographs of TPH-positive cells in the dorsal raphe are presented in Fig. 3. The number of TPH-positive cells in the dorsal raphe nuclei was 474.90 $\pm$ 17.13/section in the control group, 496.00 $\pm$ 13.88/section in the control and treadmill exercise group, 361.00 $\pm$ 20.44/section in the rotenone-injected group, 420.70 $\pm$ 9.78/section in the rotenone-injected and treadmill exercise group. TPH expression in the dorsal raphe was decreased by injection of rotenone ($P < 0.05$) and treadmill exercise increased TPH expression in the rotenone-injected rats ($P < 0.05$).

Effect of treadmill exercise on 5-HT_{1A} receptor expression in the dorsal raphe

When the level of 5-HT_{1A} receptor (50 kDa) in the control group was set at 1.00, the level of 5-HT_{1A} receptor was 1.11 $\pm$ 0.08 in the control and treadmill exercise group, 0.30 $\pm$ 0.09 in the rotenone-injected group, 0.96 $\pm$ 0.20 in the rotenone-injected

Fig. 3. Photomicrographs showing tryptophan hydroxylase (TPH) expressions in the dorsal raphe. The scale bar is 100 μ m. CON, control group; CON-EX, control and treadmill exercise group; ROTEX, rotenone-injected group; ROTEX, rotenone injected and treadmill exercise group.

Fig. 4. Effect of treadmill exercise on 5-hydroxytryptamine_{1A} (5-HT_{1A}) receptor expression in the dorsal raphe. Upper panel: Representative 5-HT_{1A} expression. Lower panel: Relative 5-HT_{1A} expression in each group. CON, control group; CON-EX, control and treadmill exercise group; ROTEX, rotenone-injected group; ROTEX, rotenone injected and treadmill exercise group. Data are presented as the means $\pm$ standard error of the mean (SEM). * $P < 0.05$ compared to the control group. # $P < 0.05$ compared to the rotenone-injected group.

and treadmill exercise group (Fig. 4). The expression of 5-HT_{1A} were decreased by injection of rotenone ($P < 0.05$) and treadmill exercise increased 5-HT_{1A} expression in the rotenone-injected rats ($P < 0.05$).

DISCUSSION

PD is closely associated with depression (Berghauzen-Maciejewska et al., 2014). FST is most commonly used test for assessment of depression in animal models of depression (Oitzl et al., 2000; Shin et al., 2017b). PD rats showed increased immobility time in FST (Shin et al., 2017b). In the present study, immobility time was increased and climbing time was decreased by rotenone injection and treadmill exercise suppressed immobility time and enhanced climbing time in the rotenone-injected rats. These results demonstrate that rotenone injection induced depressive state and treadmill exercise ameliorated this depressive state.

In the brain, serotonin cell bodies are located in the raphe nuclei of the brainstem, where they projected into the caudal brainstem and to the spinal cord (Hornung, 2003). Altered serotonergic neurotransmission contributes to the motor and non-motor features commonly associated with PD (Loane et al., 2013). Changes in the 5-HT system are associated with psychiatric disorders, including major depression and bipolar disorder (Fava and Kendler, 2000). Depressed patients showed lowered 5-HT concentration in the cerebrospinal fluid (Hou et al., 2006). Loss of tryptophan contents also increased symptom severity of depression patients, inducing downregulation of 5-HT biosynthesis (Ruhé et al., 2007). Antidepressant agents increased TPH expression as well as 5-HT synthesis (Shishkina et al., 2007; Yang et al., 2010). Dorsal raphe nuclei are involved in the anxiolytic effects of physical activity (Greenwood et al., 2005). In the present study, 5-HT and TPH expressions in the dorsal raphe were suppressed by rotenone injection and treadmill exercise increased the expressions of 5-HT and TPH in the rotenone-injected rats. These results demonstrate that rotenone induced 5-HT depletion in the dorsal raphe and treadmill exercise restored the content of 5-HT in the rotenone-induced PD rats.

Greenwood et al. (2005) reported that wheel running increased 5-HT_{1A} mRNA expression in dorsal raphe. 5-HT_{1A} receptor is closely associated with depression and antidepressant effect (Cryan and Leonard, 2000). In the present study, 5-HT_{1A} receptor was decreased by rotenone injection and treadmill exercise increased 5-HT_{1A} receptor expression in the rotenone-induced PD rats. These results demonstrate that rotenone down-regulated 5-HT_{1A} receptor in the dorsal raphe and treadmill exercise up-regulated 5-HT_{1A} receptor in the rotenone-induced PD rats.

In the present study, treadmill exercise ameliorated depressive symptoms in the rotenone-induced PD rats. The antidepressant effect of treadmill exercise might be ascribed to the enhancement

of serotonergic function through upregulation of 5-HT_{1A} expression in the dorsal raphe.

CONFLICT OF INTEREST

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

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