EVALUATION OF SIRTUIN 3 BIOMARKER BEFORE AND AFTER EXERCISE REGIMEN IN CHRONIC UNPREDICTABLE MILD STRESS-INDUCED DEPRESSED RATS

SOWNDARYA R, DOSS VA*

Department of Biochemistry, PSG College of Arts and Science, Coimbatore, Tamil Nadu, India. Email: victordoss64@gmail.com

ABSTRACT

Objective: Depression is one of the most common psychiatric illnesses that cause disability and public health problems. Physical exercise has been shown to reduce the stress impairment and ameliorate depressive symptoms. The objectives of the present study were to confirm, whether swimming exercise alleviates the depression-like behavior induced by chronic unpredictable mild stress (CUMS) through the alteration in the expression of sirtuin 3 (sirt3) in brain.

Methods: This study was undertaken to evaluate the possible antidepressant activity of exercise. Female Wistar rats weighing about 120–150 g were divided into four groups of six animals in each: Control, CUMS-induced, venlafaxine-treated, and exercise-treated groups. The rat model of depression was established using CUMS protocol. Behavioral assessment was performed by sucrose consumption test and forced swim test. Semi-quantitative reverse transcription-polymerase chain reaction and Western blot analysis were performed to study the gene expression and the protein activity of SIRT3.

Results: Our results showed that the gene expression and protein activity of sirt3 were found to be significantly increased in the exercise-treated group when compared to the depressed group.

Conclusion: The interesting fact from this study is that swimming exercise increases sirt3 expression and possesses a beneficial effect in curing depression.

Keywords: Depression, Venlafaxine, Exercise, Sucrose consumption.

INTRODUCTION

Depression is a most prevalent multifaceted mental disorder characterized by apathy, loss of interest, decreased energy, retardation of thinking, disturbed sleep, and suicidal ideation [1,2]. It is a highly prevalent disabling condition associated with significant morbidity and mortality [3]. It is a common incidence worldwide which affects the quality of life of many people and has become a major cause of suicidal death [4]. Several studies have also shown the involvement of oxidative stress in depressive state [5,6]. Antidepressants are commonly prescribed for depression and other affective disorders. However, these synthetic antidepressants even taken in appropriate doses cause side effects [7].

Sirtuin 3 (SIRT3) is a protein that is encoded by the sirt3 gene. It exhibits NAD+ dependent deacetylase activity. SIRT3 expression is highest in metabolically active tissues including the brain, heart, liver, brown adipose tissue, and skeletal muscles. SIRT3 responds to both exercise and nutritional signals in skeletal muscles to correlate downstream molecular responses [8,9]. Exercise causes the cellular adenosine monophosphate (AMP):adenosine triphosphate ratio to increase. Increased levels of AMP trigger activation of AMP-activated protein kinase initiating a signaling cascade promoting SIRT3 expression. Exercise increases SIRT3 expression as well as associated cyclic AMP response element binding phosphorylation and peroxisome proliferator-activated receptor-γ coactivator-1α (PGC 1α) upregulation [10].

PGC 1α is also induced by exercise and is known to be a key regulator of mitochondrial biogenesis. PGC 1α may play a role in controlling SIRT3 gene expression [10-12]. Brain-derived neurotrophic factor (BDNF) is a protein that belongs to the neurotrophin (nerve growth factor-related proteins) superfamily. The physiological role of BDNF involves the development of the adult CNS to promote, modify, differentiate, and support the survival of neurons. BDNF influences development, survival, maintenance, and plasticity of neurons within the immature and adult nervous system, thus influencing neuronal excitability [13]. Chronic unpredictable mild stress (CUMS) is a well-validated animal model that mimics several human symptoms of depression [14]. This has been used widely for studying clinical depression and effect of antidepressants in diverse drugs [15].

The effect of exercise on depression has long been of interest. Many studies have demonstrated the antidepressant effect of exercise intervention. They support the belief that exercise has been proven effective in improving depression. The latest neuroscience studies have reported that physical activity and exercise can change the levels of brain monoamines and neurotrophic factors, increase synaptic plasticity and neurogenesis, and alter intracellular signaling proteins and neuronal activity [16,17]. The beneficial effects of exercise on brain function have been demonstrated in animal models and in a growing number of clinical studies on humans [18]. A study has reported that routine physical exercise prevents depression relapse so much better than antidepressant medication [19].

It is a well-known that exercise can enhance mental ability, cognition, and mental balance. It is proven that exercise induces PGC-1α. Another well-known fact is that PGC-1α induces SIRT3 level in muscle. However, the same in brains is not studied much. Furthermore, it is proven recently that SIRT3 induces the secretion of BDNF, the all-important and newly discovered brain neurotrophin. This study aims to learn the gene expression of sirt3, pgc-1α, and bdnf and the protein expression of SIRT3 before and after exercise regimen.
METHODS

Animals
Young female Wistar rats were purchased from KMCH, Tamil Nadu, India. Animals were maintained at constant temperature (37°C), humidity controlled room, and 12 h light/dark cycle with access to food and water. The ethical clearance for handling of experimental animals was obtained from the Institutional Animal Ethics Committee constituted for the purpose and care of laboratory animals as per guidance of the Committee for the Purpose of Control and Supervision of Experiment on Animals (CPCSEA), Ministry of Social Justice and Empowerment, Government of India (CPCSEA/No: 158/1999/CPCSEA).

CUMS procedure
The CUMS protocol was performed according to previous study [20]. Rats were randomly divided into four groups (n=6 per group): Control group, depression-induced group, venlafaxine-treated group, and exercise-treated group. The control rats were housed together without any disturbance, while the model rats were exposed to following CUMS procedure: Cage tilting (tilt at 45°C angle) for 24 h, crowded housing (5 rats per cage), noises for 1 h (alternative periods of noise 10 min and silence 10 min), 24 h food, 24 h water deprivation, soiled bedding (wetting cage with water), removal of bedding materials for 12 h, and reversed light-dark cycle, respectively. One stressor was applied per day and the whole stress process was done in random order for 4 weeks. After 4 weeks, the depression behavior was confirmed by forced swim test (FST) and sucrose consumption test (SCT).

Behavior test
FST
The FST was carried out according to Porsolt et al., 1977 [21]. Here, the experimental rats were made to swim individually in a cylindrical container of 35 cm in height and 12 cm, such that the rat could not touch the bottom of the cylinder with its limb or tail or climb over the edge of the chamber.

Trials sessions were conducted before the actual 6 min test started. The initial 2 min in the total 6 min was not considered, because during the time period, which the animals try to escape out. A rat was considered immobile when it remained floating in water keeping all its limbs motionless, making only little movements to keep its head above water. The total duration of immobility during final 4 min was recorded. The rats were then allowed to dry in a pre-washed enclosure (~52°C) before being returned to their cages. The decrease in the duration of immobility is a measure of antidepressant activity [22].

SCT
The test was performed as described previously with few modifications [23]. After 24 h period of food and water deprivation, each rat was subjected to be in individual cage and was given two bottles containing 1% sucrose solution and water. The preference for sucrose was calculated as a percentage of consumed sucrose solution of the total amount of liquid consumed.

Exercise protocol
Rats were trained in a moderate swimming program according to previous procedure [24]. Daily swimming exercise was performed in a large plastic barrel (45 cm in diameter and 60 cm in height) filled with fresh water (32±2°C) to the depth of 50 cm. Exercise was performed at the same time daily (between 10.00 am to 12.00 pm). After swimming, rats were towel dried and kept warm by electric drier (heater).

The swimming program included two phases: Adaptation and training. During the 1st week (adaptation), the training was begun with 15 min on the 1st day, and it was increased gradually till 60 min on the last day [25].

Sample collection
Brain tissues were harvested to examine the SIRT3 gene expression using semi-quantitative and Western blot analysis.

Isolation of total RNA from brain tissue using TRI reagent
Total RNA was isolated using TRI reagent (sigma) according to manufacturer's protocol. The collected brain tissue (100 mg) was homogenized in 1 ml of TRI reagent aseptically. The mixture was centrifuged at 12,000 g for 10 min at 4°C, the supernatant was transferred to a fresh tube. Then, added 0.2 ml of chloroform and centrifuged at 12,000 g for 10 min at 4°C. Following centrifugation, the mixture separated into three phases, from which the upper aqueous phase was aspirated carefully into the fresh tube without disturbing the interphase containing DNA. The RNA pellet was precipitated by adding 0.5 ml of ice-cold isopropanol, mixed, and incubated for 10 min on ice. After incubation, the tubes were centrifuged at 12,000 g for 10 min at 4°C and the supernatant was discarded completely. To the RNA pellet, 1 ml of ice-cold 75% ethanol was added and centrifuged at 7500 g for 10 min at 4°C to wash off the remaining TRI reagent. The supernatant was decanted; the pellet was dried, dissolved in 25 µl of RNase-free water, and stored at −20°C. RNA concentration was determined using a UV spectrophotometer (Shimadzu) at 260 nm. Quality of RNA was confirmed by calculating 260/280 ratio for detection of protein contamination and by running samples on agarose gel to confirm that the samples are DNA free.

Reverse transcription (RT)
cDNA synthesis was performed using iScript cDNA synthesis kit (BioRad) according to manufacturer's protocol. Briefly, 20 µl reactions were prepared by combining 4 µl cDNA, 4 µl 30X reaction mix (containing random and oligo-dT primers), 2 µl RNA, 1 µl reverse transcriptase, and 13 µl nuclease-free water. The reaction was performed by incubating for 5 min at 25°C, followed by 30 min at 42°C and 5 min at 85°C.

Semi-quantitative RT polymerase chain reaction (RTPCR)
Semi-quantitative analysis of SIRT3 was performed according to previous procedure with few modifications [26,27]. Briefly, simultaneous amplification of both sirt3 and glyceraldehyde-3-phosphate dehydrogenase (gapdh) (housekeeping gene) was performed. Primers (Table 1) for the selected genes were designed using Oligo perfect primer designer (Invitrogen). Gapdh was used as control gene. Primers were purchased from Merck millipore.

The PCR reaction mixture comprising 2 µl of cDNA, 5 µl of 10X assay buffer, 1 µl of 30 mM 4NTP mix, 2 µl of each forward and reverse oligo nucleotide primers, 1 µl of Taq DNA polymerase and 17 µl of nuclease free water to a final volume of 30 µl. PCR was carried out under the following conditions: Initial denaturation at 94°C for 2 min, denaturation at 94°C for 45 s, and extension at 72°C for 1 min. After 35 cycles, the aliquots of PCR products were analyzed by gel electrophoresis with 2% agarose gel stained in ethidium bromide, photographed under UV light, and the bands were analyzed by ImageJ software.

Table 1: Primers for RT-PCR

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer</th>
<th>Product length</th>
</tr>
</thead>
<tbody>
<tr>
<td>gapdh</td>
<td>Fwd: ACCACAGTGAGGAGAGCTG</td>
<td>267</td>
</tr>
<tr>
<td></td>
<td>Rve: GTGAGACTGAGCCAGATGC</td>
<td></td>
</tr>
<tr>
<td>sirt3</td>
<td>Fwd: TATCCGACGTTCAGAGTCCAG</td>
<td>227</td>
</tr>
<tr>
<td></td>
<td>Rve: ACCCATGACTTCACACACAC</td>
<td></td>
</tr>
<tr>
<td>pgl1 alpha</td>
<td>Fwd: TGCACATAGTGGTGGTGGCTT</td>
<td>258</td>
</tr>
<tr>
<td></td>
<td>Rve: CCTTCGAAATGTGTGGGCA</td>
<td></td>
</tr>
<tr>
<td>bdnf</td>
<td>Fwd: GGGCGAGAATAAAAAGACTGC</td>
<td>238</td>
</tr>
<tr>
<td></td>
<td>Rve: GCGAGAACGCTCCTGTTT</td>
<td></td>
</tr>
</tbody>
</table>

RT-PCR: Reverse transcription-polymerase chain reaction, bdnf: Brain-derived neurotrophic factor, gapdh: Glyceraldehyde-3-phosphate dehydrogenase, sirt3: Sirtuin 3
WESTERN BLOT ANALYSIS

Total protein from brain tissue was extracted in RIPA lysis buffer and quantified by bicinchoninic acid protein assay. Protein samples were separated on 12.5% sodium dodecyl sulfate polyacrylamide gel electrophoresis and transferred to nitrocellulose membrane by electroblotting (100 V current for 2 h). The membranes were blocked with 5% non-fat dry milk in TBS (10 mM Tris, 150 mM NaCl, pH 7.5) overnight at 4°C. All steps were followed by 3 times washing with TBST (0.05% Tween-20, 10 mM Tris, 150 mM NaCl, pH 7.5) buffer. Next day, the primary antibody (rabbit anti-SIRT3 polyclonal antibody - 1:1000 dilutions) was added to the membranes and kept for 1 h at 4°C. After being washed, the membranes were incubated with appropriate anti-rabbit secondary antibody (diluted 1:5000) for 1 h at room temperature. Bands were visualized using the ECL substrate. Band intensity was quantified using ImageJ software.

Statistical analysis

Data were expressed as mean ± standard deviation. The levels of significant between the groups were determined by performing one-way analysis of variance and Duncan’s multiple range test for multiple comparisons among different groups using Statistical Package for the Social Sciences (version 16.0). Differences with p<0.05 were considered to be statistically significant.

RESULTS AND DISCUSSION

In FST (Table 2), there was a significant (p<0.05) increase in the immobility time of Group II (depressed rat) compared to the control rats (Group I). After subjecting to swimming exercise, the stress-induced rats showed a significant (p<0.05) decrease in immobility time when compared to the Group II rats.

In SCT (Table 2), the percentage of sucrose consumption was found to be decreased significantly (p<0.05) in Group II rats when compared to control group (Group I) and after 4 weeks of exercise regimen, sucrose consumption was found to be increased significantly when compared to the Group II stressed rats, which shows that the exercise reduces the CUMS-induced depression-like behavior in rats.

From Figs. 1 and 2, we found that the gene expression levels of sirt3, proliferator-activated receptor-γ (pgc1α), and bdnf in depressed rats were much lower than the normal rats. After 4 weeks of exercise regimen, the gene expression levels of sirt3, pgc1α, and bdnf in treated groups (III and IV) were all improved compared to the depressed groups. The results showed that swimming exercise can increase the gene expression of sirt3, which may be due to the activity of sirt3 by protecting the brain cells against oxidative stress.

![Fig. 1: Effect of exercise on mRNA expression of sirtuin 3 (sirt3) in brain tissue using semi-quantitative reverse transcription-polymerase chain reaction (RT-PCR). (a) Semi-quantitative RTPCR - gene expression of sirt3: Groups - normal control, chronic unpredictable mild stress-induced depression, venlafaxine-treated rats, and exercise-treated rats. Glyceraldehyde-3-phosphate dehydrogenase, peroxisome proliferator-activated receptor-γ (pgc1α) coactivator-1α, sirt3 - sirtuin 3, and brain-derived neurotrophic factor (bdnf). Depressed rats showed decreased pgc1α, sirt3, and bdnf expression than the normal rats. Drug- and exercise-treated rats showed increased expression of pgc1α, sirt3, and bdnf compared to untreated rats. (b) Band intensities were quantified using peak area, *p<0.05 - significant](image1)

![Fig. 2: Western blot of sirtuin 3 (SIRT3). (a) The circles indicate the 28 kDa SIRT3 protein expression in the experimental groups. It is to be highlighted that the SIRT3 normal (A) expression is reduced during depression (B) and increases efficiently during venlafaxine- (C) and exercise-treated rats (D). (b) - Band intensities were quantified using peak area, *p<0.05 - significant](image2)
Furthermore, the increased expression of SIRT3 in exercised groups was confirmed by Western blotting analysis (Fig. 2). We found that the protein expression level of SIRT3 was significantly decreased in untreated groups. After the exercise treatment, the SIRT3 level was found to be increased significantly in treated groups (III and IV), which were essentially in agreement with that of the gene expression level of sirt3. Therefore, this assay confirmed that the exercise possessed neuroprotective effect by increasing the sirt3 level in brain during depressed conditions.

DISCUSSION

In this present study, we validated, whether swimming exercise alleviates the depression-like behavior induced by chronic stress through the alteration in the expression of sirt3 in brain. Stress is known to exacerbate depression in susceptible individuals [28]. Here, we aimed to establish CUMS rat model with depression-like behavioral changes by performing stress procedures, as the CUMS-induced animal model mimicked the human depression [29]. After analyzing the depressed conditions, swimming exercise procedure was followed for 28 days. Semi-quantitative RT-PCR and Western blot analysis were performed to study the gene expression and the protein activity of SIRT3.

FST is a commonly used behavioral despair model in depression [30]. Sucrose preference is considered as one of the symptoms of depression, leading to anhedonia indicating loss of interest [31,32]. Both tests are quite sensitive and relatively specific to confirm the depressive symptoms [21]. Here, we found increase in immobility time in FST and a remarkable reduction in sucrose preference in Group II (depressed) rats reflecting the state of behavioral despair, which were claimed to reproduce a condition similar to human depression [33]. After the exercise treatment, there was a significant decrease in immobility time and significant increase in the sucrose consumption.

In gene expression study, the expression of sirt3 in brain was significantly decreased in CUMS-induced depressed rats, which may be caused due to the damage induced by CUMS in brain cells. After the treatment, the recent study shows that the enzyme SIRT3 protects brain cells against stresses and also found that physical exercises increase the expression of SIRT3, helping to protect the brain against degeneration [34]. Moreover, it was reported that sirt3 overexpression protects cultured motor neurons against the cell death promoting effect of 50D1 mutation that suggests the neuroprotection role of SIRT3 [35]. SIRT3 promotes activation of antioxidant systems, fatty acid oxidation, and neuroprotection. Thus, SIRT3 activity can reduce reactive oxygen species (ROS) levels by directly modulating key antioxidant enzymes, thereby acting as a shield against oxidative damage. Hence, it is possible that exercise activates PGC-1α and SIRT3. Increased SIRT3 lowers ROS-mediated DNA damage. Furthermore, overexpression of SIRT3 has been shown to significantly increase neuronal life span [36]. The antioxidant and metabolic effects mediated by SIRT3 suggest a potential protective role through improved mitochondrial function, which subsequently results in increased neuronal survival and reduced aging effects. The Western blot analysis of SIRT3 also showed the increased protein expression in treated groups when compared to the depressed groups. Thus, our data prove the antidepressant-like effect of exercise in CUMS-induced depressed rats.

CONCLUSION

It is found that the exercise could reverse the anhedonic behavior before and after exercise. CUMS-induced behavioral changes were ameliorated by chronic exercise and antidepressant medications. In this study, correlating sirt3 to depression through semi-quantitative RT-PCR and Western blotting revealed decreased expression in the depressed group when compared to treated (drug and exercise) groups that show increased expression than the depressed group and similar to the normal group. Thus, our data suggest that sirt3 expression can be associated to pathogenesis of depression and exercise can be used to treat depression.

ACKNOWLEDGMENT

The authors are much thankful to University Grant Commission (UGC), New Delhi, for providing financial support under Minor Research Project (F.No:4-4/2013-14(MRP-SEM/UGC-SERO)).

AUTHORS’ CONTRIBUTION

Sowndarya. R and Dr. Doss VA designed the experiment. Sowndarya. R performed the experiment and prepared manuscript. Final approval of the article was done by (Associate Prof) Dr. Doss VA.

CONFLICTS OF INTEREST

There are no conflicts of interest.

REFERENCES