Research Article

Effect of Sirtuin-1 and Wnt/β-Catenin Signaling Pathway in Rat Model of Spinal Cord Injury

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Sirtuin-1 (SIRT1) has anti-inflammatory and antioxidant effects and has been reported to be involved in spinal cord injury (SCI). Wnt/β-catenin signal has been shown to play a critical role in the pathogenesis of chronic diseases, and it participated in the recovery of nerve function after SCI. However, the specific link between them in SCI is unclear. In addition, targeting posttraumatic astrocyte apoptosis is crucial for improving neural degeneration and locomotor function. Therefore, in this article, we studied the relationship of β-catenin and SIRT1 using in the SCI rat model and primary astrocyte treated with hydrogen peroxide (H2O2) or lithium chloride (LiCl). Results showed that after SCI, SCI area and motor function recover over time, and β-catenin is gradually increased to the seventh day and then in turn decreases until 4 weeks, positively correlated with cell apoptosis. The expression of SIRT1 and downstream FOXO4 gradually increased, and β-catenin is negatively correlated with SIRT1 expression. Moreover, treatment with H2O2 in primary cultured astrocyte significantly increased β-catenin and Caspase-3 expression, while decreased SIRT1 and Forkhead box O- (FOXO-) 4. The immunofluorescence results are consistent with this. Administration of LiCl further aggravates the above results. These findings suggest that SIRT1 is negatively correlated with β-catenin in SCI, which promotes the apoptosis of motor neuron cells, which may be related to the participation of FOXO4.

1. Introduction

Spinal cord injury (SCI) is a serious neurological disorder characterized by complete or partial impairment of motor function leading to loss of motor function, which has caused chronic impairment and disability [1, 2]. SCI contains primary and secondary injuries. We cannot treat primary injury, so we mainly focus on secondary injury. The pathological process of secondary damage includes the following: a series of changes in mitochondrial dysfunction, infection, inflammation, oxidative stress injury, and apoptosis [3, 4]. Therefore, the current main therapeutic approach is to reduce or slow the pathological injury above secondary injury. For secondary SCI, neuronal apoptosis may provide a potential therapeutic target [5]. Apoptosis is a type of genetically controlled cell death [6]. It involves a series of gene activation, expression, and regulation of the role, while β-catenin [7] and Sirtuin-1 (SIRT1) [8] are involved in regulation. However, there is not yet a report about the relationship between SIRT1 and β-catenin in SCI.

SIRT1 is a protein deacetylase of class III that relies on nicotinamide adenine dinucleotide (NAD) [9]. SIRT1 protein has been proven to have anti-inflammatory and antioxidant effects, and it has a wide range of biological functions in growth regulation, stress response, oncogenesis, apoptosis, and prolongation of lifespan [10, 11]. And it can alleviate cell damage. Besides, SIRT1 has been reported to significantly inhibit inflammatory response-induced damage to neurons [12]. Studies have shown that SIRT1 plays a key role in regulating neuroinflammation following central nervous system injury and may be a new therapeutic target for post-SCI [13]. Yu et al. [12] show that SIRT1 inhibits apoptosis in vivo and in vitro models of SCI through miR-494. There are also studies show that resveratrol can protect...
SCI by activating SIRT1/AMPK signaling pathway-mediated autophagy and inhibiting apoptosis [14].

The Wnt/β-catenin signaling pathway is implicated in the growth of the nervous system, affecting cell patterning and proliferation, neuronal connectivity and survival, cell adhesion, cell motility and polarity, and axon guidance [15, 16]. Recent studies have found a molecular mechanism: β-catenin is negatively correlated with SIRT1 during neuronal apoptosis. And they found that upregulation of SIRT1 can inhibit hydrogen peroxide- (H₂O₂-) induced osteoblast apoptosis through the Forkhead box O (FOXO) 1/β-catenin pathway [17]. SIRT1 can be accessed via Wnt/β-catenin signal regulates apoptosis and extracellular matrix degradation of chondrocytes in osteoarthritis treated with resveratrol [18]. This research was to delve into the correlation between β-catenin and SIRT1 as well as its role and underlying mechanism in regulating apoptosis after SCI, which may help to develop effective treatment plan for SCI.

2. Materials and Methods

2.1. Animals. A total of 90 mature male Sprague–Dawley (SD) rats (180–220 g) were acquired from the Experimental Animal Centre of Liaoning Medical University. All rats were kept in a room controlled temperature with a cycle of 12 h light/dark cycle exposure. All experiment processes abode by the Care and Use of Laboratory Animals released by the US National Institutes of Health.

Rats were separated to 5 groups randomly (N = 16 per group): sham group, 3 days after SCI group (3dSCI), 7 days after SCI group (7dSCI), 14 days after SCI group (14dSCI), 28 days after SCI group (28dSCI), sham group for laminectomy, and the other group using a modified rat model hammer method (diameter 2 mm, weight 10 g; Allen’s method).

2.2. Construction of the SCI Model. The SCI model was built followed by Allen’s method [1]. The rats were first anesthetized by pentobarbital sodium injection of 3% (0.2 ml/100 g, i.p.). The fur from the back of T8-T11 SD rats was shaved and disinfected. The skin was then cut, the muscles and skin were separated to 5 groups randomly (N = 16 per group): sham group, 3 days after SCI group (3dSCI), 7 days after SCI group (7dSCI), 14 days after SCI group (14dSCI), 28 days after SCI group (28dSCI), sham group for laminectomy, and the other group using a modified rat model hammer method (diameter 2 mm, weight 10 g; Allen’s method).

2.3. Primary Astrocyte Culture. Primary astrocytes were prepared from spinal cords of newborn SD rat (postnatal day 3). Spinal cords of neonatal rat were ejected from the vertebral column, treated with 0.25% trypsin (T4549, Gibco, USA) for 15 min, followed by mechanical trituration in DMEM (Sigma, USA)+10% heat-inactivated fetal bovine serum (FBS, Sigma). After centrifugation at 800 rpm for 5 min, the cells were suspended in DMEM, containing FBS and 1% penicillin/streptomycin [19]. The cultures were maintained in an atmosphere containing 5% CO₂ at 37°C.

2.4. Evaluation of the Functional Recovery of Motor Neuron. We used locomotor rating scales of Basso-Beattie-Bresnahan (BBB) [20] to evaluate the recovery in behavior on 0, 1, 3, 7, 14, and 28 days after strike. Scores of BBB ranged from 0 (complete paralysis) to 21 points (normal locomotion). Average BBB score was used to evaluate functional recovery after SCI.

2.5. Western Blot. RIPA lysis buffer (Beyotime, China) was used for dissolving the prepared tissue, followed by the quantification of the final protein concentration (2 g/L) by BCA kit (Enogene, China). After being separated by SDS-PAGE, the proteins (30 μg) were transferred onto polyvinylidene fluoride (PVDF) membrane. At room temperature, the membrane was blocked with 5% skimmed milk in the Tris-buffered saline with TBST for 2 h, and then, it was incubated at 4°C with the primary antibody overnight. The following primary antibody was contained in the hybridization solution: anti-β-catenin (1:1000; 8480, Cell Signaling Technology, USA), anti-FOXO4 (1:1000; 9472, Cell Signaling Technology), anti-Caspase-3 (NB100-56708; 1:1000, Novus, USA), and anti-SIRT1 (1:1000; 8469 Cell Signaling Technology). TBST was used to rinse the membrane three times, which was then incubated with the secondary antibody (1:10,000; Earthox, USA) for 2 h at room temperature. ChemiDoc-ITMTS2 Imager (UVP, LLC, Upland, CA, USA) was employed to develop immunoreactive band, and the ImageJ software (National Institute of Health, Bethesda, MD, USA) was applied for analyzing. The above steps were repeated three times.

2.6. Hematoxylin and Eosin Staining. Following staining with hematoxylin for 10 s, we quickly washed portrait sections of 5 μm with deionized water and then placed the slides inside the HCI/100% alcohol (1:50) for 5 seconds for differentiation, which had been stained by eosin after washing with deionized water for half of an hour. Again, the slices were rinsed and dehydrated in gradient alcohol. In the final, the slices were crystal clear, fixed with xylene to the upper neutral balsam.

2.7. Nissl Staining. We immersed the coronal slices of 5 μm into the solution of chloroform and ethanol (1:1) and hydrated the slices through decreasing the concentration of alcohol (100% alcohol, 95% alcohol, and water) in order to acknowledge the numbers of neurons and notice the changes of morphology in different groups. Cresyl violet solution of 0.1% was used to rinse immerse the slices for 40 minutes at 40°C, and then, the slices were rinsed with deionized water and applied with 95% alcohol for differentiation, 100% alcohol for dehydration, and xylene for 5 minutes for being transparent. In the final, cover glass was covered on the slices using DPX.

2.8. Immunofluorescence (IF). At the room temperature, primary neurons pretreated were fixed by PFA of 4% for 30
minutes. The transverse slices of 5 μm were dried in air for 2 h at the room temperature, followed by the incubation with normal goat serum of 5% for 2 h at the room temperature, and the cells were also treated. The primary antibodies were as follows: anti-β-catenin (1:200; 8480, Cell Signaling Technology, USA) and anti-SIRT1 (1:200; 8469, Cell Signaling Technology), and the secondary antibodies were as follows: FITC goat anti-mouse IgG (1:300; Bioss, China) and Cy3 goat anti-rabbit IgG (1:300; Bioss, China). In the final, DAPI solution (1:800) was used to restain nucleus, and all the images above were captured by a fluorescence microscope (Leica, Germany).

2.9. Statistical Analysis. Data were all analyzed by the SPSS 22.0 software. T test was used for comparison between the two groups, and one-way ANOVA was used for comparison between multiple groups. Correlations between the expression of β-catenin and SIRT1 or Caspase-3 were performed using the Spearman correlation analysis. The data were expressed as mean ± standard deviation (SD), and $P < 0.05$ is considered as a significant difference.

3. Results

3.1. Motor Functional Recovery after SCI in Rats. H&E and Nissl staining were accomplished at 3, 7, 14, 21, and 28 days after SCI to assess the neurological self-recovery. The results showed that the spinal cord of rats was structurally intact with normal cell morphology in the sham group. At 3 days after SCI, the central gray matter and

![Image](image-url)
dorsal white matter were obviously damaged, necrosis and large-area cavities appeared, some nerve cells had nuclear pyknosis, and a large number of inflammatory cells infiltrated. But the damaged area recovered over time (Figure 1(a)).

The number of motor neurons in the anterior horn of the spinal cord was determined by Nissl staining after SCI. These results showed that neuron degeneration occurred 3 days after SCI, and some neuron necrosis appeared after 7 days and then recovered at 28 days (Figure 1(b)). It indicated that SCI in rats might be a self-healing process that reduced motor neuron loss and SCI.

After SCI, we measured BBB scores at 0, 1, 3, 7, 14, 21, and 28 days to evaluate functional recovery. The results in Figure 1(c) showed that after SCI, the locomotor scores of BBB of all rats fell straightly from $21 \pm 0$ to $0 \pm 0$. A few days later, gradually, the rats gradually recovered from SCI and the BBB scores were gradually restored.
3.2. The Relationship of SIRT1 and β-Catenin in SCI Rats. In order to study the relationship between β-catenin and SIRT1 at different time points after SCI and its relationship with astrocyte apoptosis, we used Western blot and immunofluorescence to measure the expression changes of β-catenin, SIRT1, FOXO4, and Caspase-3 protein. The results of Figures 2(a) and 2(b) showed that β-catenin protein increased first and then decreased, while the protein SIRT1 showed a gradual upward trend (Figures 2(a) and 2(b)). And Western blot showed the same trend of results.

Figure 3: Correlation analysis of β-catenin and SIRT1 expression in spinal cord injury rats. (a) Correlation analysis of the relationship between SIRT1 and β-catenin. (b) Correlation analysis of the relationship between Caspase-3 and β-catenin. N = 16 per group.

Figure 4: The effect of Wnt/β-catenin pathway activation on SIRT1 protein and cell apoptosis. (a) Western blot was utilized to detect β-catenin, SIRT1, Foxo4, and Caspase-3 protein expression in the different cell groups. (b) The β-catenin and SIRT1 expression in different groups of cells detected by immunofluorescence. Magnification: 200x. *P < 0.05; **P < 0.01; ***P < 0.001.
(Figures 2(c)). And FOXO4 protein was also gradually increased after SCI. However, the apoptotic marker Caspase-3 first increased and then decreased, and the peak is also on the 7 days after SCI (Figure 2(c)). Furthermore, the correlation analysis revealed that β-catenin expression was negatively correlated with SIRT1 expression (Figure 3(a)) but positively correlated with Caspase-3 expression (Figure 3(b)).

3.3. Activation of β-Catenin Decreased SIRT1 Expression and Increased Apoptosis Expression. Furthermore, we built an injury and apoptosis model by exposing glial cells to H2O2, which was used to crystallize the mechanism of β-catenin with SIRT1 involvement in neuroprotection after injury. At the same time, we used Wnt/β-catenin pathway activator LiCl (20 mM) to clarify the relationship between the Wnt/β-catenin pathway and SIRT1 and FOXO4. The results showed that apoptosis induced by H2O2 increased proapoptotic protein Caspase-3 expression, and LiCl significantly increased the level of β-catenin protein expression. In contrast, SIRT1 protein and its downstream FOXO4 protein significantly downregulated after the addition of LiCl along with H2O2 induction (Figure 4(a)). The immunofluorescence results were consistent with Western blot (Figure 4(b)).

4. Discussion

This research primarily showed that the protein β-catenin and protein SIRT1 were negatively correlated in the SCI rat model of secondary injury. LiCl, the Wnt pathway activator, increased the expression of β-catenin and inhibited the expression of SIRT1 and FOXO4.

Sirtuins are class III of histone deacetylases [21]. Among the seven sirtuins, SIRT1 regulates a variety of physiological processes, including apoptosis, DNA repair, inflammatory response, metabolism, cancer, and stress [22, 23]. In recent years, numerous studies have shown that SIRT1 has a protective role in SCI [14, 24]. Chen et al. [13] showed that agonists of SIRT1 reduced neuroinflammation and promoted neuronal survival after spinal cord injury. Some studies have shown [25] that SIRT1 is negatively correlated with β-catenin expression after SCI. Consistent with these studies, in the present study, we found that β-catenin was negatively correlated with SIRT1 expression in SCI rats and that β-catenin showed a positive correlation with the expression of the apoptotic protein Caspase-3.

The canonical Wnt pathway exerts a significant influence on cell development and differentiation [5]. Apoptosis was controlled by Wnt/β-catenin pathway, involving in human disease, tumor, and embryonic development [26]. However, the final result of the Wnt signal is determined by genes whose activities are controlled by TCF and β-catenin. When it is activated, the β-catenin accumulated in the cytoplasm will move into the nucleus to activate the TCF transcription factor participating in the DNA transcription [27, 28]. Therefore, β-catenin is the key regulator of the Wnt pathway. More recently, lithium has been demonstrated as a Wnt signaling pathway activator in animal and cell models of neurodegenerative diseases and tumors [29–31]. In our study, β-catenin was significantly upregulated compared with the decrease in SIRT1 expression downstream of the protein FOXO4 after lithium-treated glial cells. In line with this observation, there was a negative correlation between β-catenin and SIRT1 in the regulation of neuronal apoptosis in rats. Some reports have reported that activated SIRT1 reduces the β-catenin expression through phosphorylation of mesenchymal stem cells [32, 33]. It is consistent with the results of this article.

In this paper, we found SIRT1 and β-catenin showed an inverse relationship in apoptosis in SCI rats, and there was a regulatory relationship between them. These results suggested that we offered a new molecular mechanism for the potential clinical application of SIRT1 to treat SCI by activating or inhibiting β-catenin, although the complex interregulatory mechanism of SIRT1 and β-catenin is not yet clear to us. However, it lays the foundation for our next study.

**Data Availability**

The data used to support the findings of this study are available from the corresponding author upon request.

**Conflicts of Interest**

The authors declare that they have no conflicts of interest.

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