Research Article

Research on Polycaprolactone-Gelatin Composite Scaffolds Carrying Nerve Growth Factor for the Repair of Spinal Cord Injury

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Objective. This study was to investigate the mechanism of action of polycaprolactone/gelatin (PCL/GE) composite fiber scaffold with nerve growth factor (NGF) in the recovery of spinal cord injury (SCI).

Methods. Sixty female Sprague-Dawley (SD) rats were randomly assigned to the negative control group, the positive control group, the PCL/GE scaffold group, and the collagen-binding structural domain nerve growth factor (CBD-NGF)/PCL/GE scaffold group, with 15 rats in each group. Spinal cord transection was used to establish SCI models in rats. The negative control group received sham surgery, while the other three groups were given spinal cord transection at the tenth thoracic vertebra (T10) segment. The rats in the PCL/GE scaffold group were implanted with a 4 mm PCL/GE composite fiber scaffold, and those in the CBD-NGF/PCL/GE scaffold group were implanted with a CBD-NGF/PCL/GE composite fiber scaffold. The Basso–Beattie–Bresnahan (BBB) locomotor rating scale was used to evaluate the locomotor ability of the hind limbs of the rats, and the amplitude and latency of motor evoked potentials (MEP) were recorded by neurophysiological testing at 12 w postoperatively. The levels of growth-associated protein 43 (GAP43) and neurofilament protein 200 (NF200) in the spinal cord tissue of the injury site were determined using Western Blot at 12 w after surgery. Spinal cord tissues of 2 cm within the injury site, the thoracic segment above the injury site, and the lumbar segment below the injury site were collected from the measurement of axonal transport using fluorescent retrograde tracer fluorogold, and the integrated absorbance (IA) values of FC-positive cells were calculated.

Results. After treatment, the negative control rats showed normal locomotion function of the hind limb with the highest BBB scores, while the positive control rats had the lowest BBB scores and showed paraplegia. The scaffold groups exhibited better locomotion function of the hind limb and higher BBB scores than the positive controls, with greater improvement observed in the CBD-NGF/PCL/GE scaffold group (P < 0.05). Compared with the positive controls, the PCL/GE scaffold group and CBD-NGF/PCL/GE scaffold group exhibited significantly shorter latency and increased amplitude of MEP, with more significant changes observed in the CBD-NGF/PCL/GE scaffold group (P < 0.05). Compared with the positive control group, the GAP43 and NF200 levels of spinal cord tissue were significantly elevated in both the PCL/GE scaffold group and the CBD-NGF/PCL/GE scaffold group, and the changes were more pronounced in the CBD-NGF/PCL/GE scaffold group (P < 0.05). The differences in the IA values of FC-positive cells in the spinal cord tissue of the lumbar segment below the injury site among the four groups did not come up to the statistical standard (P > 0.05). Compared with the positive control group, the FC-positive cell IA values of spinal cord tissue in the thoracic segment above the injury area were markedly increased in the PCL/GE scaffold group and the CBD-NGF/PCL/GE scaffold group, and the alterations were more significant in the CBD-NGF/PCL/GE scaffold group (P < 0.05).

Conclusion. PCL/GE composite fiber scaffold with NGF significantly improves motor and neurological functions in the hind limbs of SCI rats and promotes the recovery of axonal transport, and the mechanism may be associated with the upregulation of GAP43 and NF200 levels in spinal cord injury site tissues.
1. Introduction

Spinal cord injury (SCI) is a common complication after spinal fracture and/or dislocation [1], which mainly manifests as urinary dysfunction due to disorders of the nervous system within the spinal cord and loss of control of the bladder sphincter and may cause paraplegia in severe cases [2]. The main pathological manifestations are axonal degeneration and neuronal apoptosis of spinal cord neurons. Axons function in transporting proteins and nutrients and are an important part of neurons. The present clinical treatment for SCI consists mostly of pharmacological therapy, surgery, and rehabilitation training, while their therapeutic benefits are mediocre. With ongoing research into stem cell treatment, stem cell transplantation holds significant promise for regenerating neurons and restoring neurological abilities in SCI cases [3]. However, stem cells exhibit a low survival profile and lack effective support for directed growth [4]. Repair and functional reconstruction of spinal cord injury is a major challenge in spinal surgery. Natural materials are weak in mechanical support when used directly for injury repair and fail to guide the regenerated nerve fibers through the injury site. Thus, synthetic material scaffolds provide other available alternatives. Biomaterial scaffolds are important sites for seed cells to perform their biological functions. Recent research has discovered that biomaterial scaffolds with good biocompatibility, degradability, and mechanical strength provide effective support for stem cells and boost the adhesion, migration, differentiation, and regeneration of nerve axons in vivo [5], and polycaprolactone/gelatin (PCL/GE) composite fiber scaffolds are the most commonly used in clinical practice. The combination of PCL and GE has also been proven to increase cell adhesion and degradation, which are considered contributory factors of SCI recovery [6]. Nerve growth factor (NGF) is a neurotrophic factor that is closely related to SCI [7]. NGF belongs to the important bioactive substances in the central nervous system and plays a role in promoting the growth, development, and regeneration of nerve cells in the spinal cord. It has been shown that NGF improved spinal cord injury by inhibiting neuronal apoptosis after 30-60 μg NGF intervention in t-SCI rats. With reference to the report of Chen et al. [8], our research team developed the collagen-binding domain-Nerve growth factor (CBD-NGF) for the SCI rat models to investigate the role of PCL/GE composite fiber scaffold and CBD-NGF/PCL/GE scaffold in SCI recovery, aiming to provide more references for clinical treatment.

2. Materials and Methods

2.1. Materials. The trial was conducted according to Good Clinical Practice guidelines developed by the International Council for Harmonisation and in compliance with the trial protocol. The protocol was approved by the institutional review boards or independent ethics committees at each site. Ethics No.: NU-20200102. All patients provided written informed consent per Declaration of Helsinki principles. An independent data monitoring committee monitored safety and efficacy data.

2.1.1. Main Reagents and Instruments. PCL (No. 704105) and GE (No. G9382): Sigma, U.S.A. Trifluoroethanol (Item No. TCL-T0435): Hangzhou Haoxin Biotechnology. CBD-NGF: prepared by our research team. RIPA solution (Item No. R0020), PMSF solution (Item No. P8340), and BCA microprotein concentration assay kit (Item No. PC0050): Beijing Solaobao Technology. ECL (Item No. ASW2010): Shanghai Jizhi Biochemical Technology. Mouse anti-GAP43 monoclonal antibody (Item No. ab277627): Abcam, U.S.A. Mouse anti-NF200 monoclonal antibody (item number P63-L-GE): Leica, Germany. Fluoro-gold (FG): Fluoro-chrom, U.S.A. Accumeter conductivity meter: Thermo Fisher, USA. Injection pump: Harvard Bioscience, U.S.A. Electromyography/evoked potential meter: Natus, U.S.A.

2.1.2. Laboratory Animals. Sixty female Sprague-Dawley (SD) rats (specific pathogen-free), 200-250 (220.83 ± 5.34) g, were purchased from Hubei Bainite Biotechnology Co. The animals were fed in separate cages, supplied with a daily ration of animal feed and sufficient water, and the environmental conditions were set at a temperature of 22-25°C and relative humidity of 40%-70%. The experiment strictly followed the relevant guidelines in the “Guideline on the Good Treatment of Laboratory Animals” and was approved by the hospital ethics committee.

2.2. Treatment Methods

2.2.1. Scaffold Preparation [9]. PCL and GE were dissolved in trifluoroethanol, added with 1% glacial acetic acid, and stirred at room temperature for 12 h to obtain 10% PCL/GE mixed solution. The combined solution was poured into a 10 mL syringe with a Teflon plastic hose attached at the nipple and a 20 gauge needle affixed at the end of the hose after the conductivity was measured using the Accumet conductivity meter. The syringe was clamped to the syringe pump, and a flat plate lined with aluminum foil was used as the electrospun fiber receiver, with the front end of the needle 20 cm from the receiver, and the experimental environment was set with a temperature of 20°C, humidity of 30%, voltage of 20 kV, and syringe pump rate of 1 mL/h. The PCL/GE composite fiber scaffolds were collected and dried in a drying oven under vacuum for 24 h. CBD-NGF was dissolved in sterile PBS solution and configured at a concentration of 0.25 μg/μL, and a portion of the scaffold was incubated in the solution for 2-3 min before implantation to obtain a PCL/GE composite fiber scaffold with NGF.

2.2.2. SCI Rat Model and Scaffold Implantation [10]. The rats were randomly divided into negative control group, positive control group, PCL/GE scaffold group, and CBD-NGF/PCL/GE scaffold group, with 15 rats in each group. A 4 cm longitudinal incision was made along the spinous process, concentrating on the bony landmarks of segments T9 to T10, followed by a longitudinal incision at the paravertebral muscles on both sides. The spinous processes of segments T7 to T9 were removed with bone-biting forceps, and the vertebral plates were scraped with mosquito hemostatic forceps to expose the extradural area. Using micro-biting forceps, the
spinal cord was totally exposed by biting off the vertebral plate on both sides of the pedicle. The spinal cord tissue of the negative control group received no treatment, and in the other three groups, the dura mater and spinal cord were quickly severed at the T10 segment using a scalpel, and the two severed ends of the spinal cord were gently lifted with micro-forceps to ensure complete transection of the spinal cord. After hemostasis by gelatin sponge compression, the PCL/GE scaffold group and CBD-NGF/PCL/GE scaffold group were transplanted with a 4 mm PCL/GE scaffold and a CBD-NGF/PCL/GE scaffold at the injury site, and the remaining 2 groups were left untreated. Layers of muscle, fascia, and skin were sutured, and 0.2 mL penicillin was administered intramuscularly into the leg twice daily for 10 days following surgery to avoid infections. The bladder was massaged once every 8 h until spontaneous urination resumed.

2.2.3. Behavioral Observations in Rats. The hind limb locomotor function of rats was evaluated before and after surgery at 1 d, 3 d, 1 w, 2 w, 4 w, 8 w, and 12 w using the Basso-Beattie-Bresnahan (BBB) locomotor rating scale. The rats were put in an open environment, and the movement and coordination of both hind limbs were evaluated for 5 minutes, with a score of 0 representing no visible hind limb motor function and a score of 21 representing typical hind limb locomotor performance. The score is related to the motor function of the hind limb.

2.2.4. Neurophysiological Testing. The stimulating electrode was placed at the intersection of the coronal and sagittal sutures in the cortical motor area, and the recording electrode was placed at the posterior tibial nerve. The amplitude and latency of motor evoked potential (MEP) were recorded with a single stimulation of 45 mA and 46 V, with a pulse width of 0.2 ms and a frequency of 1 Hz.

2.2.5. Determination of Spinal Cord Tissue Protein Expression by Western Bolt. Five rats were randomly selected in each group at 12 w postoperatively and sacrificed by cervical dislocation. Their spinal cord tissues were collected 1.5 cm above and below the surgical cut ends, cleaned with sterile PBS solution, dried, weighed, lysed on ice for 20 minutes with a combination of RIPA and PMSF, and crushed using an ultrasonic pulverizer. The protein was quantified using the bicinchoninic acid (BCA) test, followed by SDS-PAGE electrophoresis. The protein samples were wet transferred to PVDF membranes and blocked for 1 hour at room temperature with 5% skim milk. Then, growth-associated protein 43 (GAP43) and neuroprotein 200 (NF200) primary antibodies were added at a ratio of 1:200 and incubated overnight at 4°C in the refrigerator. The membrane was rinsed 3 times with PBST solution, an HRP-labeled secondary antibody was added at a dilution ratio of 1:500 and incubated for 90 min at room temperature, and the membrane was rinsed 3 times with PBST solution. The electrochemiluminescence (ECL) method was used for color development, the gel imaging system was used for observation, and the Image J software was used to analyze the grayscale and optical density values of the bands and calculate the relative expression of the target protein.

2.2.6. Axonal Function Measurement by Fluorescent Retrograde Tracer Fluorogold. With the posterior median vein of the L2 spinal cord as the center, one needle was inserted at 1 mm to the left and right of the spinal cord, and fluorescent gold (FG) 0.1 μL was injected at the depths of 0.5 mm, 1.0 mm, and 1.5 mm, and the needle was left in each place for 5 min. After the injection, the needle was slowly withdrawn, and the wound was sutured layer by layer and disinfected. After 1 w, the heart of the rats was perfused with 4% paraformaldehyde, and spinal cord tissues of approximately 2 cm in length, including the injury site, and spinal cord tissues of the thoracic segment above the injury site and the lumbar segment below the injury site, were cut into small segments of 4 mm in length and fixed in 4% paraformaldehyde, to prepare frozen serial coronal sections with a thickness of 30 μm. Each rat had five coronal slices of the brain and spinal cord taken, and the pictures were viewed using a laser scanning confocal microscope with an excitation wavelength of 405 nm. Positive cells were detected as brownish-yellow in the cytoplasm and/or nucleus staining using Image Pro Plus 6.0 software, and five high magnification fields were randomly selected for each sample for observation, and integral absorbance (IA) values were computed for FC-positive cells.

2.3. Statistical Analysis. The mean difference between the two groups were tested using student’s t-test for normally distributed variables and Mann–Whitney U test for nonnormal variables.

PSS 21.0 software was used for data analyses. Normally distributed measurement data were expressed as (mean ± standard deviation), one-way ANOVA was used for multiple group comparisons, and the Snk-q test was used for two-way comparisons. Count data were expressed as frequencies or composition ratios and analyzed using the chi-square (χ2) test. A difference was defined as statistically significant at P < 0.05.

3. Results

3.1. Behavioral Comparison. After treatment, negative control rats had normal hind limb locomotion function; positive control rats had paraplegia, and their hind limb locomotion capacity showed no significant changes over time. When compared to the positive controls, the rats in the PCL/GE scaffold group and the CBD-NGF/PCL/GE scaffold group regained their hind limb function gradually, with the CBD-NGF/PCL/GE scaffold group recovering faster. (P < 0.05) (Figure 1).

3.2. Neurophysiological Results. At 12 w postoperatively, MEP potentials were found in all four groups. The positive control group showed longer latency and lower amplitude of MEP than the negative control group (P < 0.05). Compared to the positive controls, the PCL/GE scaffold group and CBD-NGF/PCL/GE scaffold group had considerably reduced latency and greater amplitude of MEP, with the
CBD-NGF/PCL/GE scaffold group having more significant modifications \((P < 0.05)\) (Figure 2).

3.3. GAP43 and NF200 Protein Levels in Spinal Cord Tissue. The level of GAP43 and NF200 in spinal cord tissue was significantly lower in the positive control group compared with the negative control group \((P < 0.05)\). GAP43 and NF200 levels in spinal cord tissue were considerably higher in both the PCL/GE scaffold group and the CBD-NGF/PCL/GE scaffold group as compared to the positive control group, with the alterations being more evident in the CBD-NGF/PCL/GE scaffold group \((P < 0.05)\) (Figure 3).

3.4. Axonal Transport. The differences in the IA values of FC-positive cells in the spinal cord tissue of the lumbar segment below the injury site among the four groups did not come up to the statistical standard \((P > 0.05)\). Significantly reduced IA values of FC-positive cells in the spinal cord tissue of the thoracic segment above the injury area were found in the positive control rats versus negative control rats \((P < 0.05)\). The FC-positive cell IA values of spinal cord tissue in the thoracic segment above the injury area were significantly increased in the PCL/GE scaffold group and the CBD-NGF/PCL/GE scaffold group when compared to the positive control group, and the alterations were more significant in the CBD-NGF/PCL/GE scaffold group \((P < 0.05)\) (Figure 4).

4. Discussion

SCI is a dysfunction of the spinal cord at sites below the level of injury. Axonal regeneration and remodeling of damaged neurons are inhibited after spinal cord damage, leading to disruption of the microenvironment and causing secondary injury. Bladder dysfunction is a common SCI complication, mainly characterized by clinical features such as urinary retention, urinary incontinence, and increased residual urine volume, and in severe cases, urinary tract infection and renal impairment [12]. It is well established that axonal loss in the injured site is the main pathological change after SCI, and the restoration of axonal transport after SCI is the priority of current treatment [13]. NGF is a protein with regulatory effects on nerve cell growth and development, differentiation, regeneration, and functioning, and it has been shown to promote the repair of axonal injury and improve motor function of both lower limbs in SCI rats [14]. It has been shown that NGF in the central nervous system regulates the growth, maturation, and survival of nerve cells and is essential in neuronal development. Research has demonstrated that the level of NGF expression increases with spinal cord injury, but its low expression level fails to sustainably protect the injured neurons. NGF is a neuroprotective and regenerative agent that is commonly used for clinical symptoms of neurological injury and is available for the treatment of SCI. Therefore, exogenous NGF is required to promote motor as well as central nervous system function recovery in SCI patients.

Research has shown that tissue-engineered scaffolds could induce axonal rejuvenation by improving the internal microenvironment of the injured spinal cord, providing a new treatment alternative for SCI [15]. Because of its biodegradability, biocompatibility, and nontoxicity, PCL is commonly employed in biological tissue engineering scaffolds. Tissue-engineered scaffolds produced with PCL/GE have been shown to induce tissue remodeling and the production of neurotrophic factors to improve locomotor performance [16, 17], but their impact in encouraging nerve axon and neuron regeneration is marginal. [18]. It has been reported that trophic factors on top of tissue-engineered scaffold grafts contributed to inducing axonal regeneration at the
injury site [19]. Currently, studies on NGF plus PCL/GE scaffolds for SCI in rat models after spinal cord transection have been scarcely reported. To this end, this study was undertaken to investigate the effect of PCL/GE composite fiber scaffold with NGF for SCI and its mechanism.

In the present study, positive control rats had paraplegia, and the locomotion ability of their hind limbs showed no significant alterations with time. The latency and amplitude of MEP were reduced, as well as the IA values of FC-positive cells in the thoracic spinal cord tissue above the lesion location, which was consistent with prior study findings. Furthermore, rats implanted with PCL/GE scaffolds or CBD-NGF/PCL/GE scaffolds had better BBB scores, showing that scaffold implantation contributed to the restoration of hind limb locomotor capacity [20, 21]. The reduced latency and increased amplitude of MEP suggested that scaffold implantation could improve the abnormal nerve electrical impulses promoted by SCI and facilitate the integration of electrophysiological and synaptic signals in the neural circuit below the plane of injury [22].

In addition, the increased IA values of FC-positive cells in the spinal cord tissue of the thoracic segment above the injury area found in the rats treated with PCL/GE scaffolds or CBD-NGF/PCL/GE scaffolds suggested that axon regeneration and axonal transport restoration were associated with scaffolds implantation. GAP43 is a protein polymer that provides structural support for axons and is an essential regulator of neurite outgrowth, synapse formation, and neuronal regeneration. Both have been shown to have a role in the onset and progression of SCI. [23]. The results of the present study showed that the levels of GAP43 and NF200 were significantly increased in the spinal cord tissues of rats after scaffold implantation, suggesting that the mechanism by which scaffold transplantation improves motor and neurological functions in the hind limbs of rats with SCI and promotes axonal regeneration may be associated with the upregulation of GAP43 and NF200 levels in the tissues at the site of spinal cord injury [24, 25]. Furthermore, more pronounced alterations of GAP43 and NF200 levels in the CBD-NGF/PCL/GE scaffold group versus the PCL/GE scaffold group are indicative of the clinical value of NGF in the treatment of SCI.

The present study combined the use of traditional NGF and biomaterials to promote growth and fixation support and demonstrated that polycaprolactone-gelatin composite scaffolds carrying nerve growth factor could alleviate the local hypoxia within the early scaffold of tissue engineering materials, thus significantly promoting cell survival and

**Figure 2**: Neurophysiological results. Note: (a) indicates a significant difference ($P < 0.05$) when compared with negative controls; (b) indicates a significant difference ($P < 0.05$) when compared with positive controls; (c) indicates a significant difference ($P < 0.05$) when compared with the PCL/GE group.

**Figure 3**: GAP43 and NF200 protein levels in spinal cord tissue. Note: (a) indicates a significant difference ($P < 0.05$) when compared with negative controls; (b) indicates a significant difference ($P < 0.05$) when compared with positive controls; (c) indicates a significant difference ($P < 0.05$) when compared with the PCL/GE group.
proliferation and avoiding early apoptosis of cells due to ischemia and hypoxia, laying a certain experimental foundation for further in vivo studies using this composite material. This study explored the mechanisms of SCI and provided some insights into the treatment and diagnosis of SCI. However, no validated biomarkers were found in the present study, and the mechanism of polycaprolactone-gelatin composite scaffolds carrying nerve growth factor employed in this study was not investigated, which is a direction for future research.

5. Conclusion

PCL/GE composite fiber scaffold with NGF significantly improves motor and neurological functions in the hind limbs of SCI rats and promotes the recovery of axonal transport, and the mechanism may be associated with the upregulation of GAP43 and NF200 levels in spinal cord injury site port, and the mechanism may be associated with the upregulation of GAP43 and NF200 levels in spinal cord injury site.

Data Availability

All data generated or analyzed during this study are included in this published article.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Authors’ Contributions

Shengsen Yang and Na Zhang contributed equally to this work.

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