Bone marrow mesenchymal stem cells with Nogo-66 receptor gene silencing for repair of spinal cord injury

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Abstract
We hypothesized that RNA interference to silence Nogo-66 receptor gene expression in bone marrow mesenchymal stem cells before transplantation might further improve neurological function in rats with spinal cord transaction injury. After 2 weeks, the number of neurons and BrdU-positive cells in the Nogo-66 receptor gene silencing group was higher than in the bone marrow mesenchymal stem cell group, and significantly greater compared with the model group. After 4 weeks, behavioral performance was significantly enhanced in the model group. After 8 weeks, the number of horseradish peroxidase-labeled nerve fibers was higher in the Nogo-66 receptor gene silencing group than in the bone marrow mesenchymal stem cell group, and significantly higher than in the model group. The newly formed nerve fibers and myelinated nerve fibers were detectable in the central transverse plane section in the bone marrow mesenchymal stem cell group and in the Nogo-66 receptor gene silencing group.

Key Words: nerve regeneration; spinal cord injury; bone marrow mesenchymal stem cells; Nogo-66 receptor; RNA interference; horseradish peroxidase; BrdU; gene silencing; neural regeneration

Introduction
Spinal cord injuries result in complete or partial loss of sensation and/or mobility and affect the quality of life of patients¹⁴-⁵⁰. Severe spinal cord injury often causes paralysis and loss of sensation and reflex function below the site of injury, as well as impairing autonomic activity, such as breathing, and other functions, such as bowel and bladder control. Stem cell therapy has considerable therapeutic potential in spinal cord repair⁶-⁸. A recent study has shown that canine bone marrow mesenchymal stem cells exhibit various features in vitro that make them suitable for spinal cord reconstruction, such as their ability to proliferate as undifferentiated spheres, and under appropriate stimuli, to differentiate into neurons, astrocytes and oligodendrocytes. Because these cells can form neurosphere-like clumps and differentiate into neuron-like cells expressing neuronal markers⁹-¹², they hold great potential for nerve repair. However, mesenchymal stem cell transplantation alone is not sufficient for spinal cord repair because the majority of the mesenchymal stem cells engrafted into the spinal cord phenotypically differentiate into glial lineages and rarely survive¹³-¹⁵. The microenvironment of the injured spinal cord is thought to play a crucial role in the differentiation and survival of engrafted mesenchymal stem cells¹⁶. The neurite growth inhibition mediated by the Nogo-66 receptor¹⁷ is a major factor affecting the efficacy of mesenchymal stem cell transplantation. In this study, we used RNA interference to silence Nogo-66 receptor gene expression in mesenchymal stem cells. Our findings demonstrate the effectiveness of this strategy for enhancing mesenchymal stem cell transplantation for spinal cord injury.

Results
Morphology of bone marrow mesenchymal stem cells
The numbers of bone marrow stromal cells and colonies were significantly increased at 5 days of culture. Cells at passages 1–3 proliferated actively, and the majority of cells were seen to adhere as a monolayer. These cells were either spindle-shaped, oval-shaped, flat-shaped, triangular or irregular, and very strongly refractive, with more than two processes, whereas some of which formed connections with each other. These cells had a visible nucleus and nucleolus, and when confluent, they grew in a parallel or spiral arrangement (Figure 1A–C). Flow cytometry showed that these cells were positive for CD29, CD44, CD105 and CD166, and negative for CD34, CD80 and CD86. The bone marrow mesenchymal stem cells were quite homogeneous, with a purity above 96%.

Nogo-66 receptor expression was reduced in siRNA-transfected bone marrow mesenchymal stem cells
RT-PCR and western blot assay showed that Nogo-66 receptor gene and protein expression in siRNA-transfected bone marrow mesenchymal stem cells were significantly decreased compared with cells transfected with a control siRNA (Figure 1D).

Transplantation of Nogo-66 receptor-silenced bone marrow mesenchymal stem cells improved the morphology of the injured spinal cord
At 4 weeks after transection injury, spinal cord tissue...
increase in vehicle use, the incidence to migrate
ecrotic dysfunction. Hind limb movement recovered at 2
hind limb or tail, and urinary dysfunction, but without de
Neurological function
performance in rats with spinal cord injury
stem cell transplantation improved behavioral
delay in the bone marrow mesenchymal stem
cell group, astrocytes aggregated at the edge of the affect
ed site and formed scars at the junction of the intact and
damaged spinal cord. The cavity was smaller than in the
model group, but larger than in the Nogo-66 receptor gene
silencing group (Figure 2B). In the Nogo-66 receptor gene
silencing group, astrocytes exhibited reactive hypertrophy,
aggregated and formed scars at the edge of the affected site.
Some cells were spindle-shaped, forming a dense network
with their processes, but the cavity disappeared (Figure 2C).
Immunohistochemical staining showed that the number of BrdU-positive cells increased in rats transplanted with
Nogo-66 receptor-silenced bone marrow mesenchymal
stem cells (Figure 3), indicating improved survival of the
transplanted bone marrow mesenchymal stem cells at the
site of injury.

Transplantation of Nogo-66 receptor-silenced bone
marrow mesenchymal stem cells promoted the growth of
nerve fibers after spinal cord injury
By horseradish peroxidase retrograde nerve tracing, only a
few horseradish peroxidase-positive nerve fibers were visible
at the T1 and higher segments in the model group (Figure
4A). The number of horseradish peroxidase-positive nerve
fibers in the bone marrow mesenchymal stem cell group was
less than in the Nogo-66 receptor gene silencing group, but
more than in the model group (Figure 4B). The Nogo-66 re
ceptor gene silencing group showed a large number of horse
radish peroxidase-positive nerve fibers in the spinal cord
(Figure 4C). The number of horseradish peroxidase-positive
nerve fiber bundles is shown in Figure 4D, demonstrating
significant differences among the three groups at 8 weeks
post-injury (P < 0.01).

Effects of Nogo-66 receptor-silenced bone marrow
mesenchymal stem cell transplantation on tissue
ultrastructure in the injured spinal cord
Transmission electron microscopy showed glial scarring and
a small number of myelinated nerve fibers, macrophage
phagocytosis, degeneration and necrotic myelinated nerve
fibers in the model group (Figure 5A). Massive myelinated
nerve fibers and non-myelinated nerve fibers could be seen
in the Nogo-66 receptor gene silencing group, with more
axons and intact myelin (Figure 5B). The numbers of my
elinated nerve fibers and non-myelinated nerve fibers at the
injury site in the bone marrow mesenchymal stem cell group
were greater than in the model group, but less than in the
Nogo-66 receptor gene silencing group (Figure 5C).

Nogo-66 receptor-silenced bone marrow mesenchymal
stem cell transplantation improved behavioral
performance in rats with spinal cord injury

Neurological function
After injury, rats manifested paraplegia, no activity of the hind limb or tail, and urinary dysfunction, but without def
ecatory dysfunction. Hind limb movement recovered at 2
and 4 weeks postinjury and became more coordinated at 6
weeks, and urinary function was partially restored, but re
sidual urine was still visible in the bladder. The three groups
exhibited similar changes after injury. Basso, Beattie and
Bresnahan scores in the bone marrow mesenchymal stem
cell and Nogo-66 receptor gene silencing groups were high
er than in the model group. Moreover, Basso, Beattie and
Bresnahan scores were higher in the Nogo-66 receptor gene
silencing group than in the bone marrow mesenchymal stem
cell group (P < 0.01, P < 0.05; Figure 6A).

Inclined plate test
At 4 weeks post-injury, scores in the inclined plate test were
higher in the bone marrow mesenchymal stem cell and No
go-66 receptor gene silencing groups than in the model
group. Moreover, scores in the inclined plate test were higher
in the Nogo-66 receptor gene silencing group than in the
bone marrow mesenchymal stem cell group (P < 0.01, P < 0.05; Figure 6B).

Nogo-66 receptor-silenced bone marrow mesenchymal
stem cell transplantation reduced mortality in rats with
spinal cord injury
At 8 weeks after injury, mortality was substantially lower in
the bone marrow mesenchymal stem cell and Nogo-66 recep
tor gene silencing groups than in the model group (P < 0.05).
Mortality was lower in the Nogo-66 receptor gene silencing
group than in the bone marrow mesenchymal stem cell
group (P < 0.05; Figure 6C).

Discussion
With the continued increase in vehicle use, the incidence of spinal cord injury has risen in tandem. Spinal cord in
jury has become one of the main causes of morbidity and mortality. Current treatment methods including surgery,
medication and physiotherapy have limited efficacy. In recent years, mesenchymal stem cell transplantation for the
treatment of neurological diseases has shown considerable therapeutic potential. Mesenchymal stem cells have
many advantages, including easy collection, advanced meth
ods for separation, culture, amplification and exogenous
gene transfection, feasibility of autologous implantation
following in vitro amplification or genetic modification, low
risk of immune rejection, and fewer ethical considerations.
Mesenchymal stem cell transplantation has been shown in
a variety of experimental studies to be able to treat nervous
system injury. The mechanisms of action are complex. Mes
enchymal stem cells show a high expansion potential, genetic
stability, and stable phenotype, they can be easily collected
and shipped from the laboratory to the bedside, and they
are compatible with different delivery methods and formul
ations. In addition, mesenchymal stem cells have two
other extraordinary characteristics; they are able to migrate
to sites of tissue injury and have strong immunosuppressive
properties that can be exploited for successful autologous
as well as heterologous transplantation without requiring pharma
cological immunosuppression. Furthermore, mesenchymal
stem cells are capable of differentiating into neurons
and astrocytes in vitro and in vivo\[^{36}\]. Recently, mesenchymal stem cell injection has shown promise for amyotrophic lateral sclerosis treatment in humans\[^{37}\]. They are able to improve neurological deficits and promote the restoration of functional synaptic transmission when transplanted into animal models of neurological disorders\[^{38}\]. Mesenchymal stem cells have been observed to migrate to the injured tissues and mediate functional recovery following brain, spinal cord and peripheral nerve lesions. However, bone marrow mesenchymal stem cell transplantation alone is not sufficient for spinal cord repair, because the majority of the mesenchymal stem cells implanted into the spinal cord have been shown to differentiate into a phenotype that is restricted to glial lineages and they rarely survive. The microenvironment of the injured spinal cord is believed to play a crucial role in inducing the differentiation and survival of the grafted mesenchymal stem cells.

Growth inhibitory factors associated with the myelin
sheath limiting axonal regeneration are a major impediment to adult mammalian regeneration in the central nervous system. A variety of factors isolated from central nervous system myelin have been shown to inhibit nerve regeneration, including the protein Nogo, one of the most important growth inhibitory factors. Guo et al.\textsuperscript{39} reported that Nogo protein gene expression was elevated after central nervous system injury in rats. Jiang et al.\textsuperscript{40} showed that Nogo protein, Nogo-66 receptor and RhoA expression in the brain tissues of rats with focal cerebral infarction began to increase at 6 hours, reached its peak at 24 hours, and lowered back to normal levels at 96 hours. Nogo protein may lead
to growth cone collapse and inhibit neurite extension. The Nogo monoclonal antibody can neutralize the inhibitory activity of the protein. In vitro cultured oligodendrocytes also exhibit an inhibitory effect on axonal extension. Similarly, Yang et al.\textsuperscript{[45]} showed that Nogo neutralizing anti-body could promote recovery in rats with spinal cord injury. The inhibitory effect of Nogo protein is mediated by the No-go-66 receptor\textsuperscript{[42–47]}. In this study, gene silencing was used to suppress Nogo-66 receptor expression in mesenchymal stem cells, thereby promoting neurite growth following mesenchymal stem cell differentiation, and improving the efficacy of mesenchymal stem cell transplantation for repairing damage following central nervous system injury. Nogo-66 receptor gene knockout can result in permanent Nogo-66 receptor gene silencing, but the physiological function of the Nogo-66 receptor gene remains unclear, and permanent Nogo-66 receptor gene silencing may lead to unexpected consequences. RNAi is a very convenient and effective method of gene silencing, which is usually maintained for 3–5 days. This method is ideal for our study because high Nogo-66 receptor expression is observed following traumatic brain injury\textsuperscript{[48–49]}. To avoid the drawbacks of permanent gene silencing, RNAi silencing combined with mesenchymal stem cell transplantation is the most promising treatment strategy for spinal cord injury.

Recent in vivo and in vitro studies in non-neuronal and neuronal tissues have shown that different pathways of macrophage activation result in cells with different properties. Interleukin-6 triggers the classically activated inflammatory macrophages (M1 phenotype), whereas the alternatively activated macrophages (M2 phenotype) are anti-inflammatory. In this study, mesenchymal stem cells were subjected to Nogo-66 receptor gene silencing before transplantation, which may result in better repair of the damaged brain tissue, and promote mesenchymal stem cell proliferation and differentiation in the grafted area after injury. Our results showed that mesenchymal stem cell transplantation, after Nogo-66 receptor gene silencing, is greatly superior to simple mesenchymal stem cell transplantation in the treatment of spinal cord injury in rats, in terms of histological and functional outcomes. Tissue repair was better in the Nogo-66 receptor gene silencing group compared with the model and bone marrow mesenchymal stem cell groups. Immunohistochemical staining demonstrated a significant difference in the number of BrdU-positive cells and horseradish peroxidase-positive nerve fibers at the site of spinal cord injury, as follows: Nogo-66 receptor gene silencing group \textgt; bone marrow mesenchymal stem cell group \textgt; model group.

In summary, the Nogo-66 receptor gene in mesenchymal stem cells can be silenced using the RNAi approach. The mesenchymal stem cells can be transplanted into the site of spinal cord injury via the tail vein. The transplanted mesenchymal stem cells better survive, proliferate, differentiate and migrate at the site of injury, and promote the recovery of nerve function after spinal cord injury. Our findings provide support for the use of this novel approach for the clinical treatment of spinal cord injury.

### Materials and Methods

#### Design

A randomized, controlled, animal experiment.

#### Time and setting

This experiment was performed at Hebei Medical University in China from May 2010 to May 2011.

#### Materials

Sixty-four clean, healthy Wistar rats of both genders, aged 2 months and weighing 250–300 g, were purchased from the Chinese Academy of Medical Sciences Animal Laboratory (license No. SCXK20060008). All experimental procedures were performed in accordance with Chinese National Natural Science Foundation animal research regulations and the animal care guidelines of the National Institutes of Health.

#### Methods

**Bone marrow mesenchymal stem cell isolation and culture**

Bone marrow was harvested aseptically from the tibias of rats at the age of approximately 2 months. Nucleated cells were isolated by density gradient centrifugation using Percoll (1.073 g/mL) and were plated in growth medium consisting of Dulbecco’s modified Eagle’s medium/F12 (HyClone, Logan, UT, USA) supplemented with 20% fetal bovine serum (Sigma, St Louis, MO, USA) and benzylpenicillin (1 × 10^5 U/mL).\textsuperscript{[49]} The mesenchymal stem cells were isolated in the medium by their tendency to adhere to plastic.\textsuperscript{[50–52]} After 3 days, the dishes were washed twice with PBS to remove nonadherent cells. The remaining cells were fed every third day. Nonadherent cells were removed and adherent cells were expanded until subconfluence and processed through sequential passages. Most contaminating hematopoietic stem cells were progressively lost, and after the second passage, cultures contained a morphologically homogenous cell population designated bone marrow mesenchymal stem cells. This was confirmed by fluorescence-activated cell-sorting analysis showing a lack of expression of the typical hematopoietic cell surface markers, including CD45, CD34 and CD14, and positivity for CD71, CD105 and CD44. Cells between passages 3 and 6 were used for our experiments. They were labeled using medium containing BrdU.\textsuperscript{[49–52]}

**RNAi-transfected Nogo-66 receptor-silenced bone marrow mesenchymal stem cells**

The two target sequences of the rat Nogo-66 receptor mRNA were as follows: 5’-UGG AGU ACC UCU ACC UAG ACA A-3’, 5’-UUU UGU UGU AGU AGU UAG UAC UGC A-3’. The siRNA template was synthesized by Shanghai Sangon Biological Engineering Technology and Service Co., Ltd. in Shanghai, China. According to the in vitro transription kit (Silencer\textsuperscript{TM} siRNA Construction Kit, Ambion) instructions, 1 mL of culture medium containing 1 × 10^5 mesenchymal stem cells was added into a centrifuge tube and centrifuged at 800 r/min (radius = 16 cm) for 5 minutes. After discarding the supernatant, 900 μL culture medium without
antibiotics was added, and cells were resuspended. Fifty pmol siRNA was diluted with 50 μL Opti-MEM, and 1 μL Lipo-fectamine 2000 was diluted with 50 μL Opti-MEM, mixed and incubated at room temperature for 15 minutes. The two solutions were then gently mixed and incubated for 15 minutes at room temperature. The mixture was added into the mesenchymal stem cell suspension, placed at 37°C in a 5% CO₂, saturated humidity incubator, and 72 hours later, Nogo-66 receptor expression was assessed.

We isolated total RNA from the injured spinal cord (4 mm long) using the RNeasy Kit (Qiagen), and obtained cDNA using reverse transcription. Primers were as follows: Nogo-66 receptor sense, 5′-GGG CAA CCT CAC GCG CAT CT-3′ and anti-sense, 5′-GGT CCT GTA TGC CTC TGG TC-3′; β-actin sense, 5′-GTC CCT GTA TGC CTC TGG TC-3′ and anti-sense, 5′-GGT CTT TAC GGA TGT CAA CG-3′. All primers were synthesized by Sangon. The β-actin gene was amplified for 24 cycles, and the others were amplified for 28 cycles. The reverse transcription-PCR product was subjected to 1.3% agarose gel electrophoresis and processed with a gel imaging system (Beijing Seclaser Technology Co., Ltd., Beijing, China), and the ratio of Nogo-66 receptor absorbance to β-actin absorbance was used as an index of Nogo-66 receptor mRNA expression level.

Western blot assay for detecting the effectiveness of Nogo-66 receptor transfection

The undifferentiated mesenchymal stem cell suspension in the control group and the mesenchymal stem cell suspension at 72 hours after siRNA transfection were centrifuged at 800 r/min (radius = 16 cm) for 5 minutes. The cells were collected. Supernatant was discarded. Four hundred μL protein extraction solution was added, and proteins were extracted. Protein concentration was determined using the Bradford method. The samples were subjected to SDS-PAGE, blotted onto a membrane, and incubated with rabbit anti-mouse Nogo-66 receptor gene antibody (1:800; Sigma) at 37°C on a shaker for 2 hours. The membrane was washed with Tris-buffered saline containing Tween-20 for 5 minutes (four times). The blots were incubated with goat anti-rabbit antibody (1:700; Sigma) at 37°C for 1.5 hours, washed with Tris-buffered saline containing Tween-20 for 5 minutes (four times), and reacted with 3,3′-diaminobenzidine. The experiment was repeated three times. Quantity one image (BioRad, Hercules, CA, USA) analysis was conducted. The absorbance value ratios of the target and β-actin bands were assessed and expressed as protein expression levels.

Preparation of animal models of spinal cord injury

Sixty-three adult Wistar rats were anesthetized with a 4% chloral hydrate solution (360 mg/kg intraperitoneally). Body temperature was monitored and maintained using a thermal blanket. Using the aseptic surgical technique, a dorsal incision was made from the middle to lower thoracic regions. Using a microscope, T₉₋₁₀ laminectomy was performed, and the spinal cord was exposed. The spinal cord was transected at the level of T₉₋₁₀, leaving a 2-mm gap between the proximal and distal ends of the resected cord.

Transplantation of mesenchymal stem cells

Sixty-three model rats were equally and randomly divided into three groups. Six hours after injury, bone marrow mesenchymal stem cells pre-labeled with bromodeoxyuridine were transplanted via tail vein. The model group was injected with 1 mL of stem cell-free culture medium. The bone marrow mesenchymal stem cell group was injected with 1 mL of bone marrow mesenchymal stem cells (5 × 10⁷/L). The Nogo-66 receptor gene silencing group was injected with 1 mL (5 × 10⁷/L) Nogo-66 receptor gene-silenced bone marrow mesenchymal stem cells. Intraportal injection of gentamicin 2,000 U served as antibiotic treatment. Rats were fed in separate cages.

Histological examination

At 4 weeks after injury, the rats were randomly selected from each group and killed under anesthesia. Specimens were harvested for histological examination to determine the degree of recovery. The dissected spinal cord tissues were post-fixed for 3 hours in 4% paraformaldehyde, soaked overnight in 10% followed by 30% sucrose, and cut into 15-μm-thick sagittal and parasagittal sections using a cryostat. Hematoxylin-eosin staining was carried out for general histological examination.

Immunocytochemistry for BrdU expression in cells

Four weeks after operation, two rats were randomly taken from each group. Immunocytochemistry for the detection of BrdU requires a pretreatment of tissue sections to denature DNA. All staining was done on free-floating 40-μm sections. A mouseanti-BrdU monoclonal antibody (1:100; Boehringer Mannheim, Ingelheim am Rhein, Germany) was used in combination with avidin biotin complex and a horse anti-mouse IgG antibody conjugated with biotin (1:167; Vector Laboratories, Burlingame, CA, USA). Primary and secondary antibodies were incubated at 37°C. Ten fields were randomly selected from each slice under the light microscope at a magnification of ×200 (Sigma). The number of BrdU-positive cells was calculated in each field of vision, and the mean value was obtained.

Retrograde tracing with horseradish peroxidase

Eight weeks after operation, two rats were randomly taken from each group. After surgery, the spinal cord was exposed at T₁₀, and 1 μL aqueous suspension of 30% horseradish peroxidase (Sigma) was injected bilaterally 1 mm into the spinal dorsal vein. After injection, the incision was closed, and the animals were maintained for 36 hours before being perfused with buffered 1% paraformaldehyde and 1.25% glutaraldehyde. The spinal cord was removed and stored in 20% sucrose in 0.1 mol/L PBS at 4°C overnight. The spinal cord was dissected, and 10 fields were randomly selected from each slice under the light microscope at ×200. The number of horseradish peroxidase-la-
beled nerve fibers was calculated in each field, and the mean value was obtained.

**Electron microscopy**

Eight weeks after operation, two rats were randomly selected from those subjected to labeling from each group, and perfused intracardially with saline \(^{[17-20]}\), followed by 2% glutaraldehyde and 4% paraformaldehyde in 0.1 mol/L sodium cacodylate buffer, pH 7.4. Immediately after perfusion, the spinal cords were removed and postfixed in the same fixative overnight at 4°C. The spinal cord segment at the injury site was sliced into 1-mm pieces, postfixed for 2 hours in 1% OsO\(_4\) in 0.1 mol/L cacodylate buffer, dehydrated in graded ethanol solutions, and embedded in Epon-812. Semi-thin plastic sections (1 µm) were cut and stained with 1% toluidine blue before examination with a Nikon Eclipse TE300 microscope equipped with a Spot RT Color CCD camera. For electron microscopy, blocks were trimmed and sections were cut at 100-nm thickness, mounted on copper grids, stained with uranyl acetate and lead citrate, and viewed with a JEOL Jem 1200 EX transmission electron microscope (JEOL Ltd., Tokyo, Japan).

**Evaluation of functional recovery**

Two types of functional tests were used to assess functional recovery. Each test was observed by two independent investigators. The test was performed at 1, 2, 4, 6 and 8 weeks post-operation.

- **Basso, Beattie and Bresnahan:** the open-field locomotion test assesses movement, weight support and coordination. It was scored using the standardized Basso, Beattie and Bresnahan locomotor scoring system. Basso, Beattie and Bresnahan scores range from 0 (flaccid paralysis) to 21 (normal gait). Rats were acclimated to the testing environment (90-cm diameter plastic wading pool; 4-cm height) prior to testing. Basso, Beattie and Bresnahan scores were averaged for each group.

- **Inclined plane test:** this test evaluates the maximum angle on the inclined plane upon which each animal can maintain a stable position for 5 seconds. Rats were placed on a board that was incrementally raised to increasing angles \(^{[17-20]}\). Inclined plane scores were averaged for each group.

Mortality: the mortality of each group was evaluated at 8 weeks post-injury.

**Statistical analysis**

Data were expressed as mean ± SD, and analyzed using SPSS 17.0 software (SPSS, Chicago, IL, USA). Intergroup comparison was done using one-way analysis of variance. Paired comparison was done using Dunnett’s \(t\)-test. A value of \(P < 0.05\) was considered statistically significant.

**Author contributions:** All authors designed, implemented, evaluated the study, and approved the final version of the paper.

**Conflicts of interest:** None declared.

**Peer review:** This study used gene-silenced bone marrow mesenchymal stem cell transplantation in animal models of spinal cord injury, provided new ideas and experimental evidence for the treatment of spinal cord injury in the clinic. This method can restore neurological function in patients with spinal cord injury and improve their quality of life.

**References**


