Investigation for the correlation between brain injury and injured ipsilateral sciatic nerve regeneration in a rat model

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Previous studies showed that brain trauma promotes repair of peripheral nerve injury by reducing scar in nerve endings. The effect of brain injury at different locations on ipsilateral and contralateral sciatic nerve regeneration in Sprague-Dawley rats was found to promote the repair of ipsilateral sciatic nerve injury, thus providing further experimental evidence for the unveiling of traumatic brain injury promoting peripheral nerve regeneration.

Keywords
Brain injury; neurotrophic factor; peripheral nerves; nerve regeneration; gastrocnemius motor endplate; rat model

1. Introduction
Traumatic brain injury (TBI) is one of the most common clinical traumas, incidence of which is also on rise (Kulbe and Geddes, 2016; Reis et al., 2015). Many studies (Eggers et al., 2016; Li and Cao, 2000; Liu et al., 2013; Yang et al., 2015) have shown that traumatic brain injury could up-regulate the neurotrophic factors, some of which could promote regeneration of peripheral nerve damage, such as neurotrophin-4/5, brain-derived neurotrophic factor. The previous experimental findings indicate that accompanied by the brain injury, regeneration of sciatic nerve injury in rats could be promoted (Zhou et al., 2018). This study aimed to explore whether there are differences in effects of brain injury at different locations on repair and regeneration of sciatic nerve.

Gibson (1960) reported the phenomenon of sciatic nerve repair associated brain injury and fracture healing for the first time. They discovered that femoral fracture associated craniocerebral injury demonstrates more osteophyte formation at the fracture end. Roberts (1968) reported that heterotopic ossification occurs after the brain injury. Subsequently, a large number of experimental studies and clinical observations confirmed that craniocerebral injury could accelerate healing of fractures (Hofman et al., 2015; Wildburger et al., 1994). Meanwhile, the neurotrophins, basic fibroblast growth factors, and the other substances have been proven to play important roles in the pathological processes (Hofman et al., 2015; Wildburger et al., 1994).

A few studies (Pu et al., 1999; Santos et al., 2016) also clarified the effects of rat brain injury on regeneration of a sciatic nerve post the injury. The results demonstrated that sciatic nerve injury with a brain injury demonstrates better functional recovery and neuro-morphology compared to that with sciatic nerve injury alone (Pu et al., 1999; Santos et al., 2016). The other studies also showed that when cerebral cortical injury and its corresponding inferior neuronal axon damage at the same time, the expression of some related growth factors are increased and the nerve conduction pathway is also changed (Acosta et al., 2013).

A former study (Lykissas et al., 2007) for traumatic brain injury reported that a major mechanism promoting neuronal repair after brain injury is the up-regulation and release of endogenous neurotrophic factors, such as neurotrophin-4/5, nerve growth factor, basic fibroblast growth factors and brain-derived neurotrophic factors. Peripheral nerve regeneration is a process of axon regeneration and remyelination. This process is affected by many factors, such as microtubule structure. The microtubule structure of basement membrane is composed of extracellular matrix, which could provide molecular guidance signals for regeneration of axons. Schwann cells and neurotrophic factors, such as cell adhesion molecules, could also provide materials and nutritional supports for the axon regeneration. The abundant vascular system could provide sufficient nutrients and oxygen and transport the metabolites. Appropriate physical stimulation can also facilitate activation of neuronal cells and promote growth and development of axon growth cones (Gonzalez-Perez et al., 2013; Homes et al., 2011; Sebben et al., 2011). Then, when the inferior neuron had a certain change due to damage of the superior neuron, its axon regeneration may be affected accordingly. Moreover, Zhou et al. (2018) also reported that accompanied by brain injury, regeneration of sciatic nerve injury in rats could also be promoted. Therefore, this study experimentally designed the sciatic nerve injury model group with ipsilateral and contralateral brain injury, which was used to observe whether there is any difference in effect on functional recovery and histo-morphology of damaged sciatic nerve. Moreover, this study also attempted to explore the relevant mechanisms for the regeneration of peripheral nerve damage in brain-injured rats.
2. Material and methods
2.1 Animals and reagents
This study mainly included the following reagents and equipment: fluoro-gold (Cat No.: 39286-10MG-F, Sigma-Aldrich, St Louis, MI), acetylcholinesterase staining of gastrocnemius motor plates, according to the previous study recorded (Das et al., 2017). Briefly, anesthesia was conducted with a volume fraction of 10% chloral hydrate (3.5 mL/kg) intraperitoneally. Postanesthesia, skin preparations were done by disinfecting with alcohol and iodine. Then, the skin was opened in right hip, and the biceps femoris was bluntly separated to expose sciatic nerve. The sciatic nerve was cut approximately 5 mm below the lower hole of piri-formis, and then the sciatic nerve was sutured under a microscope with a 10-0 non-invasive suture. Finally, the skin was sutured to close the wound.

Brain injury model: The brain injury model was generated according to the previously published study (Akyol et al., 2018). In brief, the Feeney method was used to establish brain injury models in groups B and C at the same time. Group B received the right brain injury, and group C received a left-brain injury. All rats were given a volume fraction of 10% chloral hydrate (0.35 mL/kg) intraperitoneal injection for anesthesia. The skin was prepared by disinfecting with alcohol and iodine. Sagittal cutting along the head was made to expose the parietal bone. At 1.5 mm behind the coronal line, and 2.5 mm lateral to the midline, a dental drill was used to make a 5 mm diameter bone window, while keeping the dura intact. A 20 g hammer was used to hit the impact rod from a height of 30 cm along the peripheral catheter. After the striker hit the brain tissue, the bone window was closed with bone wax, and the skin was closed by staples.

2.4 Measurements after surgery
All animals were survived after modelling, and no wound infection was found. After the model was established, the right lower limbs of the rats were limited in active flexion. The right foot was deformed, and the foot was healed when walking.

2.5 Measurement and calculation of sciatic nerve function index
At 4, 6, 8, 10, and 12 weeks after modelling, 8 rats in each group were randomly selected to measure and calculate the sciatic nerve function index. The length, breadth, height, and height of wooden channel was 70 cm, 9 cm, 10 cm, respectively. One end was connected to a homemade cassette, and a long strip of white paper of 70 cm × 9 cm was placed at the bottom of the channel. Rats’ feet were evenly stained with red ink. From one end, the testing rat walked through the channel into the cassette, and step on the white paper to leave the footprints. After drying, the value was measured. Each rat was measured 3 times at a time, and the average value was taken. The parameters to be measured were listed as the followings: 1 print length (PL): the longest distance from the footprint, i.e., from the heel to the toe; 2 toe spread (TS): from the first to the fifth toe Connection distance; 3 intermediate toe spread (IT): the distance between the 2-4 toe.

Sciatic function index = 38.3 × (EPL - NPL)/NPL + 109.5 × (ETS - NTS)/NTS + 13.2 × (EIT - NIT)/NIT - 8.8 (Yue et al., 2019), where E is the experimental side (right side), N is the healthy side (left side), and the nerve function index S is 0 as the normal value, and -100 is the index of complete nerve dissociation.

2.6 Wet mass of gastrocnemius muscle and calculation of the wet mass ratio
At the 4th, 8th, and 12th week after modelling, 8 rats in each group were randomly selected and anesthetized by intraperitoneal injection of chloral hydrate at a volume fraction of 10%. The calf muscle was incised at the posterior edge of the calf by retrograding from calcaneal tubercle to the femoral condyle. Then, the gastrocnemius muscle was completely removed. The electronic balance was used to weight the wet mass of gastrocnemius muscle. Then, the wet mass ratio was calculated as the following formula: Wet mass ratio = ipsilateral wet mass of muscle/contralateral wet mass muscle. Post weighting the gastrocnemius muscle mass, the isolated muscle was stored at -80 °C for the following analyses.

2.7 Acetylcholinesterase staining of gastrocnemius motor endplates
Preparation of incubation solution (Kamorsky-Roots solution): iodine acetylcholine 12.5 mg (Sigma-Aldrich), 0.1 mol/L sodium dihydrogen phosphate 7 mL, 0.1 mol/L disodium hydrogen phosphate 9 mL, 0.1 mol/L 1 mL of sodium citrate, 2.5 mL of copper sulfate (30 mmol/L), 2.5 mL of 5 mmol/L potassium ferricyanide, and 2 mL of distilled water were added in order and mixed together. The fresh liquid was appeared to be bright green.

Materials acquisition, fixation, sectioning, and staining: The acetylcholinesterase staining was utilized to evaluate gastrocnemius motor plates, according to the previous study recorded (Liu et al., 2013), with a few modifications. Briefly, at 4 weeks, 8 weeks, and 12 weeks, 8 rats in each group were randomly selected (the same rats that were measured for the wet weight of the gastrocnemius muscles), and the volume fraction of chloral hydrate was 10%. The bilateral gastrocnemius muscles were weighed and recorded. Under the microscope, the medial nerve of the right gastrocnemius was taken from the muscle, and the 5 mm tissue section was cut out. The nerve tissue was fixed in 4% formaldehyde for 1 h and then rinsed twice with phosphate buffered saline (PBS) for 3 min each. OCT compound was used to embed the fixed tissue. Approximately 6 μm sections were cut and washed with double
Figure 1. Acetylcholinesterase staining on the rat gastrocnemius in each group. (A-C) The acetylcholinesterase staining of the right end of the sciatic nerve injury group in rats with acetylcholinesterase; (D-F) Sciatic nerve transaction plus ipsilateral brain injury group; (G-I) Sciatic nerve transaction plus contralateral brain injury group. The motor endplates are oval tan particles. As time goes on, the number of motor endplates gradually increases, and the staining increases. (J) Quantification of motor plates marked with acetylcholinesterase staining. * indicated $P < 0.05$ comparing to SNT only group. SNT = sciatic nerve transaction. Scale bar = 50 $\mu$m.

distilled water, dried, and sealed with neutral gum. Slides were then observed under the light microscope. Images of the endplate were taken and analyzed with Image-Pro Plus 6.0 to calculate the average absorbance density.

2.8 Fluoro-gold tracing

The fluoro-gold tracing was conducted according to the former study (Puigdellivol-Sanchez et al., 2002), with some modifications. In brief, at 4th, 8th, and 12th week after surgery, 3 rats in each group were randomly selected for the retrograde tracing of fluoro-gold. Fluoro-gold was used to trace L4-L5 vertebrae, and the number of fluoro-gold positive stained spinal cord anterior horn motor neurons was detected and calculated. Anesthesia was administrated in the abdominal cavity with a volume fraction of 10% chloral hydrate. The sciatic nerve was exposed again along the original incision of the right buttock. To ensure the integrity of epineurium, a Fluoro-gold solution (concentration: 4%, dose: 5 $\mu$L) was injected into the nerve trunk 5 mm distal to the anastomosis. The needle was kept for 2 min, and was slowly pushed to close the wound layer by layer. After 7 days, the chest of rats was opened to expose the heart under anesthesia. Place the puncture needle in the apex of the heart. Insert the ascending aorta and fix it with silk thread. Cut the right atrial appendage, quickly infuse the normal saline 200-300 mL, and observe the right auricle flowing out of the clear liquid. Then, followed with the injection of 400-500 mL of a 40 g/L paraformaldehyde solution containing 5% sucrose (faster and slower). Subsequently, the right sciatic nerve was exposed, and the spinal cord was opened layer by layer. The corresponding sciatic nerve was retrogradely positioned in the corresponding spinal segment. L4 and L5 spinal cords were cut out, and some nerve roots were left to identify the right and left. Total of 40 g/L paraformaldehyde solution was used to fix the tissue at 4 °C overnight. Then, tissue emerged in 10%, 20%, and 30% sequential sucrose solutions. After that, OCT was used to embed the tissue, and a cryotome was used to cut spinal cord cross-section with thickness of 20 $\mu$m. Sections were air-dried, sealed with anti-fluorescence quenching mounting medium, and stored in dark. Observation and counting of fluorescent gold-positive neurons were done under a fluorescence microscope.
2.9 Motor endplates examination

To identify motor endplate activities, the wholemount acetylcholinesterase (AChE) was employed to stain the muscles of rat models. The isolated and dissected muscles were stained with AChE to identify the distribution and localization for motor endplates of rat models. In brief, staining procedures for wholemount AChE were conducted depended on the previous study reported (Zhang et al., 2011).

2.10 Statistical analysis

All data were assigned as a mean ± standard deviation (SD) and analyzed with SPSS software 19.0 (SPSS Inc., Chicago, Ull, USA). The one-way analysis of variance (ANOVA) followed by post hoc pairwise comparison test was performed for comparing multiple groups. Student’s t-test was used to compare the mean of the two groups. P < 0.05 means a significant difference.

3. Results

3.1 Sciatic nerve function index

After a prolonged operation, the sciatic nerve function index of each group gradually recovered. In the 4th and 6th weeks, both group B and group C were superior to group A (Table 1, P < 0.05). Meanwhile, there was no significant difference between group B and group C (Table 1, P > 0.05). At 8 weeks, the recovery of sciatic nerve function index in group B was significantly better compared to that in groups A and C (Table 1, P < 0.05).

Table 1. The rat sciatic functional index in each group at different time points after modeling.

| Time (week) | Group A | Group B | Group C |
|-------------|---------|---------|---------|
| 4           | -72.323 ± 5.203 | -69.541 ± 4.118<sup>a</sup> | -72.181 ± 5.634<sup>c</sup> |
| 6           | -72.936 ± 5.745 | -62.924 ± 3.248<sup>b</sup> | -64.729 ± 4.277<sup>a</sup> |
| 8           | -57.965 ± 6.498 | -47.406 ± 3.361<sup>ab</sup> | -56.106 ± 6.173 |
| 10          | -51.985 ± 5.659 | -37.752 ± 2.852<sup>ab</sup> | -49.602 ± 6.764 |
| 12          | -40.113 ± 4.469 | -29.794 ± 3.004<sup>ab</sup> | -36.051 ± 6.158 |

Group A was a simple right sciatic nerve transaction group. Group B was a right sciatic nerve transaction with a right-brain injury group. Group C was a right sciatic nerve transaction with a left-brain injury group. Compared with group A, <sup>a</sup>P < 0.05. Compared with group C, <sup>b</sup>P < 0.05. Compared with group A, <sup>c</sup>P < 0.05.

3.2 Gastrocnemius wet mass ratio

At 4 weeks after modelling, there was no significant difference in the wet weight ratio of the gastrocnemius muscle between 3 groups (Table 2, P > 0.05). From the 8th week, the B group was significantly higher compared to that in A and C groups (Table 2, P < 0.05).

3.3 Acetylcholinesterase staining of gastrocnemius motor endplate

Acetylcholinesterase staining was performed to detect the endplate of the gastrocnemius muscle. There was no significant difference in the absorbance of the endplate of the gastrocnemius muscle between three groups at 4 weeks after the surgery. From the 8th
Table 2. The wet weight ratio of rat gastrocnemius in each group at different time points after modeling.

| Time (week) | Group A          | Group B          | Group C          |
|-------------|------------------|------------------|------------------|
| 4           | 0.450 ± 0.031    | 0.528 ± 0.059    | 0.509 ± 0.068    |
| 8           | 0.507 ± 0.048    | 0.573 ± 0.047ab  | 0.537 ± 0.051    |
| 12          | 0.553 ± 0.051    | 0.682 ± 0.052ab  | 0.630 ± 0.031    |

Group A was a simple right sciatic nerve transaction group. Group B was a right sciatic nerve transaction with a right-brain injury group. Group C was a right sciatic nerve transaction with a left-brain injury group. Compared with group A, \(^{ab}P < 0.05\). Compared with group C, \(^bP < 0.05\).

Table 3. The absorbance density value of the motor endplate of gastrocnemius at different times after model establishment in rats.

| Time (week) | Group A          | Group B          | Group C          |
|-------------|------------------|------------------|------------------|
| 4           | 0.242 ± 0.056    | 0.215 ± 0.038    | 0.221 ± 0.064    |
| 8           | 0.287 ± 0.049    | 0.339 ± 0.052    | 0.290 ± 0.062    |
| 12          | 0.313 ± 0.070    | 0.403 ± 0.121ab  | 0.336 ± 0.058    |

Group A was a simple right sciatic nerve transaction group. Group B was a right sciatic nerve transaction with a right-brain injury group. Group C was a right sciatic nerve transaction with a left-brain injury group. Compared with group A, \(^{ab}P < 0.05\); compared with group C, \(^bP < 0.05\).

week, the absorbance density of group B was significantly higher compared that of group A and C (Table 3, \(P < 0.05\)). Also, the number of motor endplates gradually increased, the area gradually increased, and the color also gradually increased (Fig. 1). Quantification of motor endplate confirmed the impression of stained images. The number of motor endplates in group B and group C was significantly more compared to that in group A (Fig. 1).

3.4 Fluorogold tracing

As shown in Table 4 and Fig. 2, the results of Fluoro-gold retrograde tracing showed that the number of Fluoro-gold staining positive spinal anterior horn motor neurons in groups B and C was significantly higher compared to that in group A at 4 weeks after model establishment (\(P < 0.05\)). At 8 weeks, the number of Fluoro-gold staining positive neurons was significantly greater in group B compared to that of group A (\(P < 0.05\)). At 12 weeks, the number of fluorescent gold-positive neurons in group B was significantly higher compared to that in groups A and C (\(P < 0.05\)). At the same time, group C was more compared to that in group A (\(P < 0.05\)). Over time, the number of motor neurons marked in the anterior horn of the spinal cord gradually increased.

4. Discussion

In this study, we selected the male rats to establish the traumatic brain injury due to the following reasons: (1) Male rats will be larger in size and weight and easier to operate. (2) The vitality of male rats is stronger. (3) The change of hormone level in male rats is more stable than that in female rats, such as estrogen, growth hormone, etc. (4) Male rats are more sensitive to chronic poisons. The current experimental results showed that the recovery of sciatic nerve function and morphological indexes in group with ipsilateral brain injury are superior to those with contralateral brain injury and sciatic nerve injury alone, with significant differences. The results of the contralateral brain injury group were also significantly better compared to those of the simple sciatic nerve group. In terms of functional performance, as measured by the sciatic nerve function index, which objectively reflected the overall functional recovery of the sciatic nerve, was the most commonly used evaluation for sciatic nerve injury regeneration (Shen and Zhu, 1995). Another parameter to reflect nerve function recovery is the gastrocnemius wet mass ratio. The endplate staining results indirectly reflected the recovery of motor nerve fiber function and the trophic effect on muscle tissue (Ma et al., 2002). Fluorogold tracer reflected the axon regeneration and the recovery of axon transport function, but also indirectly indicated the neuronal function recovery (Catapano et al., 2008). Combined with previous studies and current experimental results, it was presumed that a large number of neurotrophic factors produced by brain injury in two groups of rats with brain injury were the main reasons for the accelerated regeneration of sciatic nerve axons. However, the left cerebral cortex controlled the right sciatic nerve, therefore its injury and the secondary axonal injury adversely affected the secondary motor center, then reduced the promotion of neurotrophic factors and other substances on the sciatic nerve regeneration (Erturk et al., 2016; Kwon and Jang, 2014). Meanwhile, the left cerebral cortex was not damaged for the ipsilateral brain injury group, therefore, the production of neurotrophic factors was more than that in the contralateral brain injury group. Therefore, the contralateral brain injury group was slightly better than that of the sciatic nerve injury group, as well as the ipsilateral traumatic brain injury group was better than that of the contralateral brain injury group for promoting the regeneration of peripheral nerve injury.

In summary, an ipsilateral traumatic brain injury could promote the regeneration of peripheral nerve injury. The possible mechanism may be that the traumatic brain injury induces up-regulation of neurotrophic factors, which reversely improve the regeneration of peripheral nerve injury (Werner and Stevens, 2015). Therefore, our future studies will focus on the finding of the relevant neurotrophic factors or growth factors and the determination of the type of neurotrophic or growth factors that play a major role in the promotion of axon regeneration.

Table 4. The number of rat spinal cord anterior horn motor neurons positive for fluorogold in each group.

| Time (week) | A group          | B group          | C group          |
|-------------|------------------|------------------|------------------|
| 4           | 5.800 ± 2.305    | 6.800 ± 2.077ab  | 6.067 ± 2.052c   |
| 8           | 6.467 ± 2.066    | 8.733 ± 2.434ab  | 7.067 ± 2.017    |
| 12          | 7.733 ± 2.434    | 10.933 ± 2.219ab | 8.933 ± 2.017c   |

Group A was a simple right sciatic nerve transaction group; group B was a right sciatic nerve transaction with a right-brain injury group; group C was a right sciatic nerve transaction with a left-brain injury group. Compared with group A, \(^{ab}P < 0.05\); compared with group C, \(^bP < 0.05\); compared with group A, \(^{c}P < 0.05\).
role. The present study would provide a theoretical basis for further investigations revealing the therapeutic approaches of traumatic brain injury for treating and promoting the peripheral nerve injury.

Abbreviations
OTC, optimal cutting temperature; PL, print length; TS, toe spread; IT, intermediate toe spread; AChE, acetylcholinesterase; ANOVA, analysis of variance.

Ethics approval and consent to participate
The animal experiments were conducted according to Guidance of Care and Use of Laboratory Animals of National Institutes of Health (NIH). All of the animal experiments were approved by the Ethics Committee of the Affiliated Hospital of Chengde Medical University, Chengde, P. R. China. Approval number: LL037.

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Conflict of interest
The authors declare no conflict of interest.

Authors' contributions
HQW and PW designed this study. HQW, JJM, XZH, and QX performed the research. YTF made statistical analyses and conducted the literature review. HQW and PW wrote and revised the manuscript.

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