The impact of bone marrow-derived mesenchymal stem cells on neovascularisation in rats with brain injury

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Abstract

Introduction: The impact of bone marrow-derived mesenchymal stem cells (BM-MSCs) on neovascularisation in rats with craniocerebral injury (CI) was investigated.

Material and methods: The rate CI model was established by the free-fall method, and then rats were randomly divided into a transplanted group (EG) and a control group (CG). Rats in the transplanted group were injected with BM-MSCs in the lateral ventricle, while those in the control group were injected with normal saline. Modified neurological severity scores (mNSSs) were compared. Furthermore, CD34+CD133 double-labelled cells (CD34+CD133+ cells) in the peripheral blood and expression of CD31 and neuron-specific enolase in injured tissues were detected.

Results: There were significant intra- and intergroup differences in modified neurological severity scores at different time points. The number of CD34+CD133+ cells in the peripheral blood of CG initially decreased, but it increased later and peaked 6 h after injury, then gradually decreased and returned to normal 24 h after injury. Cells in the peripheral blood of EG continued to increase until 24 h after injury, reaching a number higher than that in CG (p < 0.05). The postoperative expression of neuron-specific enolase in EG was higher than that in CG (p < 0.05). The positive expression of CD31 was lower in the two groups before surgery, but the expression in EG was higher than that in CG after surgery (p < 0.05).

Conclusions: Bone marrow-derived mesenchymal stem cells transplantation can increase the number of endothelial progenitor cells in the peripheral blood of rats with traumatic brain injury and increase the expression of peripheral angiogenetic markers and neuronal markers. The neurological function in EG improved significantly compared to that of CG.

Key words: bone marrow mesenchymal stem cells, rat, traumatic brain injury, neovascularisation.

Introduction

The incidence of craniocerebral injury (CI) is high, and its pathological changes can be divided into primary injury and secondary injury [22,23]; the former is neuronal damage directly caused by external violence, while the latter is caused by a variety of factors, and thus its pathophysiological responses are very complex [7,19], making CI one of the main causes of human disability and death. Presently, there are a variety of treatment methods against secondary injury, which can reduce mortality and improve prog-
nosis, but many patients still suffer from neurological dysfunction and life-long disability, causing a heavy burden to individuals, their families, and societies.

As described above, numerous studies have examined neurological repair. Recent studies have shown that the cerebrovascular system that remains stable after traumatic brain injury can be activated to initiate vessel remodelling [1,11], and the newly formed vasculature can rapidly promote nerve recovery, including nerve regeneration and synaptic regeneration, to promote the recovery of neurological function [13].

Transplanting neural stem cells (NSCs) or bone marrow-derived mesenchymal stem cells (BM-MSCs) can promote angiogenesis after ischaemic brain injury [5,18,20]. However, whether endogenous or transplanted NSCs after trauma are involved in neovascularisation after traumatic brain injury remains unclear. In this study, rats with brain injury were divided into an experiment group (EG) and a control group (CG) to analyse and compare their modified neurological severity scores (mNSS), changes in CD34+CD133+ cells in the peripheral blood, and expression of CD31 and neuron-specific enolase (NSE) in injured tissues, to analyse the neovascularisation conditions in peri-injury brain tissue and the impact of BM-MSC transplantation on nerve repair and functional recovery.

Material and methods

Animals

A total of 50 adult male healthy SD rats (120-300 g) were provided by the Experimental Animal Centre of the Department of Medicine, Nanchang University.

This study was carried out in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health. The animal use protocol was reviewed and approved by the Institutional Animal Care and Use Committee (IACUC) of the First People's Hospital of Jiujiang.

Preparation of rat craniocerebral injury model

After anaesthetisation by intraperitoneally injecting 10% chloral hydrate (1-2 ml/kg), each rat was fixed onto a brain stereotaxic apparatus, sterilised, the scalp was cut open, and one 3-mm round bone window (in diameter) with a point 3 mm posterior to the anterior fontanelle and 2 mm left of the sagittal line as the centre was drilled. One self-made strike tube was disinfected and then tightly fixed at the edge of the bone window. One 20-g weight was placed 15 cm down the strike tube to strike the dura mater, striking the right parietal lobe of the dura mater with a striking force of 300 g × cm to induce brain laceration. The bone window was sealed with bone wax and the rat was masked for oxygen supply after suturing the scalp [19].

Isolation, amplification, culture, and identification of bone marrow-derived mesenchymal stem cells

The tibia and femur were sampled from each rat killed by cervical dislocation (body weight ~120 g) and then rinsed with DMEM (Hyclone, Logan, UT, USA). The obtained cell suspension was then centrifuged at 1500 r/min for 5 min. The cell precipitate was mixed with 10% foetal bovine serum (SJQ Co., Hangzhou, China)-containing DMEM and cultured at 37°C and 5% CO₂; the cells were digested and passaged once every three days (1 : 2), and P5-generation cells were harvested for analysis.

P3 generation BMSCs were extracted by flow cell phenotype identification and added to three tubes after digestion and suspension. Next, 0.02 ml CD90-fluorescein isothiocyanate (FITC) and CD34-PE (Wuhan Boster Co., Ltd., Wuhan, China) were added to the tubes, and the tubes were incubated in the dark for 30 min. Marked single-cell suspensions were obtained for flow cytometry after washing the cells with PBS.

Transplantation of bone marrow-derived mesenchymal stem cells and grouping

The rats were divided into the EG and CG groups by using a simplified randomisation method, with 25 rats in each group. Rats in the EG were transplanted with BM-MSCs via intraventricular injection 12 h after surgery; before transplantation, BrdU (Medchem Express, Monmouth Junction, NJ, USA) (final concentration: 10 mM) was added to the medium for 2–4 h of co-incubation with the cells. The cells were then digested and adjusted to a density of 1 × 10⁶ cells/ml, and then each rat in the EG was anesthetised and fixed onto the stereotaxic for injection. Needling point: AP = −1 mm, ML = −1.0-1.5 mm, 3.5-4.0 mm dorsoventral to the skull plane. The needle
was vertically inserted into the left ventricle and labelled cells (10 μl, 1 × 10^4/rat) were slowly injected into the ventricle by one micro-injector. The needle was maintained intracerebrally for 3 min before suturing the wound. Rats in the CG group were injected with saline intraperitoneally.

**Determination of motor function**

Six rats were sampled from each group on day 1 (D1), D3, D7, D14, and D21 after surgery, and motor function was determined by mNSS (Table I) and a double-blind method.

**Detection of CD34+CD133+ cells in peripheral blood**

Twelve rats in each group were randomly selected, and 0.60 ml of venous blood was sampled from the venous plexus of the medial canthus on H3 (3 h after injury), H6, D1, D3, D7, and D14 after surgery using a clean hard glass capillary (1 mm in diameter), which was then placed into an EDTA-anticoagulated tube to isolate mononuclear cells form the peripheral blood by Ficoll density gradient centrifugation.

Ten microliters of goat anti-rat CD133 monoclonal antibody (primary antibody, Wuhan Boster Co., Ltd.) and 2 μl of FITC-labeled rabbit anti-goat IgG monoclonal antibody (secondary antibody, Wuhan Boster Co., Ltd.) were mixed and incubated at 4°C in the dark to prepare FITC-labelled CD133 (CD133-FITC); CD34-TRITC (Wuhan Boster Co., Ltd.) was prepared using the same method. The rat mononuclear cells were first added to 100 μl of 0.50% foetal bovine serum buffer (v/v), mixed evenly, and then 10 μl of CD34-TRITC and 10 μl of CD133-FITC were mixed, followed by low-speed agitation, full rinsing, 10-min incubation at room temperature in the dark, rinsing,
and centrifugation to discard the supernatant and prepare the mononuclear cell suspension. The number of CD34+CD133+ cells in the mononuclear cell suspension was detected by flow cytometry.

**Detection of neuron-specific enolase and CD31 in peri-injury brain tissue**

Two rats from each group were randomly selected and killed at different time points (before surgery, H6, D1, D3, D7, D14, and D21 after surgery), followed by 4% paraformaldehyde perfusion, fixation and dehydration in neutral formaldehyde solution, wax embedding, slicing of the site with obvious injury, haematoxylin and eosin (HE) staining, and detection of CD31 and NSE by the SP immunohistochemical method. Four sites were selected from each slice under a low-magnification microscope (×100), and each site was photographed five times using a high-magnification microscope (×400). The images were analysed using the Image Pro plus 6.0 colour pathological image analysis system; the mean optical density and relative area (%) of the positive area of each image were measured, and the product of these two was used to determine the relative content.

**Microvessel identification and density peri-injury brain tissue**

Immunohistochemical staining was used to detect the positive expression of CD31 in peri-injury brain tissue before and 3, 7, and 14 days after surgery. Endothelial cells stained as brownish yellow particles were defined as positive. Low-magnification scans of the whole slice were acquired to find the “hotspot” with maximum vascular density and determine the average number of blood vessels stained from three high magnifications (×400) to represent microvessel identification and density (MVD). The microvessel was defined from the separation of endothelial cells and nearby tissue, but not by the vascular lumen or erythrocyte, and excluded lumen with a diameter more than 60 μm.

**BrdU-positive cells in the experimental peri-injury brain tissue group**

Fourteen days after brain injury, the expression of BrdU was detected by immunohistochemistry using the primary antibody 5-bromodeoxyuridine. Positive expression appeared as brown particles in the nucleus. The inspected area was defined by low magnification (×100), which was the edge of injury grey matter and white matter, while morphological changes were observed at high magnification (×200 and ×400).

**Statistical analysis**

SPSS 16.0 (SPSS, Inc., Chicago, IL, USA) was used for statistical analysis. The measurement data are expressed as x ± s; intergroup comparison was conducted by the paired t test or independent-sample t test, and intragroup comparison was conducted by analysis of variance (ANOVA), with the test level α as 0.05.

**Results**

**Amplification, culture, and identification of bone marrow-derived mesenchymal stem cells**

After seeding, the cells gradually showed flask wall-adherent growth, the number of third-generation BM-MSCs was low, and the cells were scattered and had clustered spindle or triangular shapes. With the rapid proliferation of BMSCs and natural selection of wall-adherent predominance, P3-generation cells were purified to only wall-adherent cells (Fig. 1). The phenotype was tested based on P3 cells by flow cytometry: the expression of CD90 was positive (68.49%) while the expression of CD34 was negative (2.85%), as shown in Figure 2.

**CD34+CD133+ cells in peripheral blood**

Mononuclear cells in the peripheral blood were isolated at different time points, and the number of CD34+CD133+ cells in mononuclear cells was detected by flow cytometry. CD34+CD133+ cells in the circulation of peripheral blood were approximately 40-52 cells/200,000 mononuclear cells, with the average being 47/200,000 mononuclear cells. The number of CD34+CD133+ cells in the peripheral blood in group CG decreased until 3 h after injury and then increased to normal. The average number of CD34+CD133+ cells reached the highest level 6 h after injury, with an average of 66 cells/200,000 mononuclear cells, which gradually decreased and returned to normal 24 h after injury. The EG group exhibited the same trend; however, the number of CD34+CD133+ cells continued to increase until 24 h after injury, and
the number was significantly higher than that in CG group (t test, $p < 0.05$, Table II; Figs. 3, 4).

**Expressions of neuron-specific enolase and CD31 in peri-injury brain tissue**

The relative contents of NSE and CD31 were detected with the Image pro plus 6.0 colour pathological image analysis system. The results are shown in Figures 3 and 6.

Positive expression of CD31 in the two groups before surgery was low, but began increasing slightly at postoperative 6 h and 1 day; CD31 was highly expressed on postoperative day 3, reaching a peak on postoperative D7, and began decreasing on postoperative D14. There were significant differences in the expression of CD31 between the EG and CG groups on postoperative D1, D3, and D7 ($p < 0.05$).
The two groups exhibited minor positive expression of NSE before surgery, but NSE expression first decreased at postoperative 6 h and then increased at postoperative three days, which peaked on postoperative D7 and began decreasing on postoperative D14. There were significant differences in the expression of NSE between the EG and CG groups on postoperative D3 and D7 ($p < 0.05$, Table III).

In the normal group, CD31 expression was very low in the normal group (Fig. 5A), which was increased in the experiment group seven days after surgery (Fig. 5B) and decreased significantly by 14 days after surgery (Fig. 5C); expression was high in the control group seven days after surgery (Fig. 5D) and decreased by 14 days after surgery (Fig. 5E).

The expression of NSE in the normal group, shown in Figure 6A, was decreased at 6 h after injury (Fig. 6B) and increased seven days after injury (Fig. 6C). Expression was even higher 14 days after injury (Fig. 6D) in the experimental group, and the neurons and cytoplasm were nearly normal with a clear nucleolus. NSE was decreased 6 h after injury (Fig. 6E) and increased seven days after injury (Fig. 6F). Expression was even higher 14 days after injury (Fig. 6G) in the control group. Some neurons showed atrophy with reduced cytoplasm and an unclear nucleolus.

**Microvessel identification and density peri-injury brain tissue**

There was less cortical microvasculature in normal rats, and MVD was increased in other groups compared to the normal group, showing significant differences ($p < 0.05$); at three and seven days after surgery, MVD of experimental rats was significantly higher than in the control group (Table IV).

**BrdU-positive cells in the experiment group peri-injury brain tissue**

Multiple BrdU-positive cells (transplanted cells) are shown in Figure 7. Figures 7A and B shows the cortical regions; BrdU-positive cells were neuron (arrow)
Fig. 5. CD31 expression of peri-injury brain tissue. A) Normal group, B) 7 days after the injury in experiment group, C) 14 days after the injury in experiment group, D) 7 days after the injury in control group, E) 14 days after the injury in control group (OLYMPUS, ×400).

Table III. Expression of CD31 and neuron-specific enolase (NSE) in peripheral tissues of brain (n = 20)

<table>
<thead>
<tr>
<th>Index</th>
<th>Group</th>
<th>0 h</th>
<th>6 h</th>
<th>1 d</th>
<th>3 d</th>
<th>7 d</th>
<th>14 d</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CD31</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Control group</td>
<td>1.29 ± 0.04</td>
<td>9.22 ± 1.02</td>
<td>19.87 ± 5.03</td>
<td>27.89 ± 2.33</td>
<td>31.23 ± 2.38</td>
<td>6.02 ± 2.00</td>
</tr>
<tr>
<td></td>
<td>Experimental group</td>
<td>0.87 ± 0.06</td>
<td>12.98 ± 1.11</td>
<td>29.77 ± 6.29*</td>
<td>34.45 ± 3.02*</td>
<td>40.10 ± 3.00*</td>
<td>7.30 ± 1.19</td>
</tr>
<tr>
<td></td>
<td>NSE</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Control group</td>
<td>21.34 ± 3.42</td>
<td>14.92 ± 3.87</td>
<td>28.77 ± 2.21</td>
<td>31.67 ± 3.99</td>
<td>30.03 ± 4.88</td>
<td>22.22 ± 2.01</td>
</tr>
<tr>
<td></td>
<td>Experimental group</td>
<td>19.88 ± 2.77</td>
<td>18.18 ± 4.28</td>
<td>32.94 ± 3.02</td>
<td>38.63 ± 2.70*</td>
<td>44.29 ± 5.29*</td>
<td>23.20 ± 3.40</td>
</tr>
</tbody>
</table>

*p < 0.05
shaped, polygonal, had large cell bodies with a central nucleus, round, and the outline was clear with an obvious nucleolus. Some unclear nuclear shrinkage was observed. Figures 7C and D shows white matter; the cells had a glial cell morphology (arrow), were round, had a small cell body, and the nucleus was large and clearly with less cytoplasm observed (OLYMPUS, X400, Tokyo, Japan).

Comparison of modified neurological severity scores

There were significant differences in mNSS between the two groups (ANOVA of repeated-measurement, F = 5.997, p = 0.034), as well as significant differences in the intragroup comparison at different time points (ANOVA of repeated-measurement, F = 37.106, p = 0.000). Group CG showed significantly higher mNSSs than group CG on D7, D14, and D21 after surgery (t test, p < 0.05). The mNSS scores of the two groups first increased and then decreased over time, reaching a maximum on D7; the scores on D7 and D14 were significantly higher than on D3 (t test, p < 0.05, Table V).

Discussion

Craniocerebral injury is a common and frequently occurring neurological disease; although many treatment methods are available, such as surgery and drugs, many patients experience life-long disability. Therefore, researchers have examined the impact of stem cells on the recovery of neurological impairment.

Consistent with previous studies [3,15], CD90, a non-haematopoietic marker, showed high expression, confirming the purity of the cultured mesenchymal stem cells. Additionally, CD34, a haematopoietic marker, showed low expression. As shown in Figure 2, the CD90-positive rate of isolated BMSCs...
was 68.49%, while the CD34-positive rate was only 2.85%. Additionally, as a positive marker of haematopoietic stem cells, CD34 was not primarily expressed in BMSCs. These results reflect the high purity of third-generation BMSCs.

Stem cell transplantation can help in the recovery of neurological function after CI. Additionally, the brain vascular system can be activated after CI and begin vascular remodelling; this new vasculature is involved in the recovery of neurological function [1,11,13,18,20]. However, whether stem cells play a role in repairing injured nerves through neovascularisation remains unclear. This study investigated the mobilisation of CD34+CD133+ cells in the post-CI rat brain and vascular regeneration in the surrounding area by analysing neovascularisation in the surrounding area and the impact of BM-MSCs transplantation on nerve repair and functional recovery. The results showed that BM-MSC transplantation can increase the number of endothelial progenitor cells in the peripheral blood of rats with traumatic brain injury and increase the expression of peripheral vascular angiogenic markers and neuronal markers. Neurological function in the EG group significantly improved compared to that in the CG group.

CD34 is a surface marker of haematopoietic stem cells or mature endothelial cells [6]. However, endothelial progenitor cells (EPCs) only account for a small fraction of all CD34-positive cells in the bone marrow or peripheral blood; CD133 is a surface marker of neuroepithelial stem cells [8]. With the differentiation and mature of EPCs, the expression of CD133 gradually decreased, and mature vascular endothelial cells showed no expression of CD133, making it a good molecular marker for identifying mature endothelial cells from EPCs. Thus, the number of CD34+CD133+ cells reflects the number of endo-

Fig. 7. BrdU-positive cells expressed in the experimental peri-injury brain tissue group. A) and B) show the cortical regions, and BrdU-positive cells were neuron (arrow) shaped, polygonal, had large cell bodies with a central nucleus, round, and the outline was clear with an obvious nucleolus. Some unclear nuclear shrinkage was observed; C) and D) show white matter, and the cells had a glial cell morphology (arrow), were round, had a small cell body, and the nucleus was large and clear with less cytoplasm observed (OLYMPUS, ×400).
vascular progenitor cells [14]. EPCs are precursor cells of endothelial cells and are involved in the angiogenesis and repair of endothelial cells after injury. Mobilising and transplanting EPCs can improve the angiogenic ability of ischaemic tissues, thus promoting the repair and re-endothelialisation of damaged blood vessels. According to the results shown in Table II, Figures 3 and 4, CD34+CD133+ cells, namely EPCs, were present in the peripheral blood of rats under normal conditions, and the number of EPCs decreased rapidly in the superperic phase (3 h after CI) but increased until higher than the normal levels by 6 h after CI; the number of EPCs in the CG group returned to normal 24 h after surgery, but the number of EPCs in the EG group continued to increase until 24 h after injury. The numbers of EPCs in the EG group 6 and 24 h after injury were significantly higher than that in the CG group, suggesting that BM-MSC transplantation increases the number of EPCs in the peripheral blood of CI rats, with the effects persisting for approximately 24 h.

CD31, a platelet endothelial cell adhesion molecule, is mainly expressed in endothelial cells. As an endothelial cell junction molecule, CD31 is a marker of angiogenesis in the body [9]. Table III and Figure 5 show that the expression of CD31 in the two groups was low before injury, but expression began increasing at H6 and on D1 after injury, was highest on D7, and began decreasing on D14. The expression of CD31 in the EG group 6 and 24 h after injury were significantly higher than that in the CG group, suggesting that BM-MSC transplantation increases the number of EPCs in the peripheral blood of CI rats, with the effects persisting for approximately 24 h.

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Table IV. Microvessel density peri-injury brain tissue (vessels/400 field, \( \bar{x} \pm s \))

<table>
<thead>
<tr>
<th>Group</th>
<th>0 h</th>
<th>3 d</th>
<th>7 d</th>
<th>14 d</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control group</td>
<td>22.34 ± 1.72</td>
<td>53.57 ± 2.33</td>
<td>55.51 ± 3.89</td>
<td>35.53 ± 1.73</td>
</tr>
<tr>
<td>Experimental group</td>
<td>23.21 ± 1.18</td>
<td>65.45 ± 3.02*</td>
<td>79.38 ± 2.00*</td>
<td>37.30 ± 1.19</td>
</tr>
</tbody>
</table>

*Compare with control group, \( p < 0.05 \)

Table V. Comparison of modified neurological severity scores at different time points between two groups (\( \bar{x} \pm s, n = 6 \))

<table>
<thead>
<tr>
<th>Group</th>
<th>1 d</th>
<th>3 d</th>
<th>7 d</th>
<th>14 d</th>
<th>21 d</th>
<th>F</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control group</td>
<td>5.00 ± 0.63</td>
<td>6.83 ± 0.75</td>
<td>8.17 ± 0.75**</td>
<td>7.00 ± 0.89**</td>
<td>5.33 ± 0.52**</td>
<td>5.997</td>
<td>0.034</td>
</tr>
<tr>
<td>Experimental group</td>
<td>5.33 ± 1.03</td>
<td>6.00 ± 0.89</td>
<td>7.50 ± 1.05*</td>
<td>6.17 ± 0.41*</td>
<td>4.17 ± 0.41*</td>
<td>5.997</td>
<td>0.034</td>
</tr>
<tr>
<td>Sum</td>
<td>5.17 ± 0.87</td>
<td>6.42 ± 0.90</td>
<td>7.83 ± 0.94</td>
<td>6.58 ± 0.79</td>
<td>4.75 ± 0.75</td>
<td>5.997</td>
<td>0.034</td>
</tr>
</tbody>
</table>

*Intragroup comparison among D7, D14, and D3; \( p < 0.05 \); intergroup comparison, \( p < 0.05 \)
were round, had a small cell body and clear nuclei, as well as less cytoplasm. Combined with the regional expression of CD31 in the pia mater in cortical areas, transplanted cells underwent directed migration to the injury site and integrated the damaged region, which was coordinated with vascular regeneration, to repair the nerve. However, the mechanism of directed migration to the lesion is unknown.

Table V shows that there were significant differences in mNSS between the two groups (ANOVA of repeated measurement, F = 5.997, p = 0.034), and mNSS in the CG group on D7, D14, and D 21 was significantly higher than in the EG group (t test, p < 0.05), indicating that BM-MSC transplantation significantly improved the neurological function of CI rats.

EPCs are a class of cells that have been widely examined in recent years. In 1997, Asahara et al. [2] isolated CD34+ cells from adult peripheral blood mononuclear cells for the first time. Because these cells can differentiate into endothelial cells in vitro, express endothelial markers, and participate in angiogenesis, they were named EPCs. EPCs, also known as vascular cells, are pluripotent stem cells that can circulate, proliferate, and differentiate into precursor cells of mature vascular endothelial cells, but lack the characteristic phenotypes of mature endothelial cells.

Under ischaemic and hypoxic conditions after injury, angiogenesis and vascularisation can occur [12]. Angiogenesis refers to the process by which EPCs differentiate into endothelial cells and form the original vascular network. Vascularisation refers to the process of new capillary growth from existing blood vessels. Previous studies of adult neovascularisation consider that this process is controlled by an angiogenic mechanism, and current studies of EPCs in the peripheral blood have shown that adult angiogenesis also includes an angiogenesis mechanism [16]. During the disease process of local tissue ischaemia or endothelial injury, EPCs can be mobilised from the bone marrow to the blood circulation, which then reach injured sites, differentiate into endothelial cells, and form new blood vessels [10,17]. Therefore, adults exhibit increased local blood flow and promoted tissue repair through the angiogenesis mechanism.

Mobilisation of EPCs after injury is very important. Under normal conditions, EPCs in the bone marrow are in a resting state and can be mobilised into the systemic circulation under many stimulating factors. An increase in the number of EPCs in peripheral circulation under many stimulating factors, thus forming endothelial cells and promoting angiogenesis. Factors that have been identified to mobilise EPCs include growth factors, such as vascular endothelial growth factor and basic fibroblast growth factor [21]. Mesenchymal stem cells (MSCs) are stem cells with self-proliferative capacity and multi-directional differentiation potential and are widely distributed in the connective tissue and organ stroma throughout the body. MSCs can be easily sampled, isolated, and largely amplified in vitro. Transplanted MSCs in vivo can secrete many vasoactive growth factors, including vascular endothelial growth factor, which is one of the main factors that promotes vascular growth [4]. MSCs can release cytokines to increase the number of EPCs in the peripheral blood circulation, which in turn promotes angiogenesis and the recovery of neurological function through angiogenesis.

The results of this study are consistent with those of previous studies. However, this study was conducted only at the molecular and behavioural levels, and thus there are some limitations. Additional studies should be conducted at the genetic level.

In conclusion, understanding the conditions of post-CI neovascularisation and the impact of BM-MSCs on neovascularisation in rats with traumatic brain injury provides a foundation for CI treatment by stem cell transplantation. Clinical treatment of severe traumatic brain injury is a long-term complex problem, and it is difficult to achieve breakthroughs and improvements in a short period or improve the treatment effects of patients with severe traumatic brain injury through one drug or one method. However, clinical and basic studies can improve the treatment effects of traumatic brain injury.

Disclosure

The authors report no conflict of interest.

References


