Transplantation of human amniotic mesenchymal stem cells in the treatment of focal cerebral ischemia

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Received February 6, 2012; Accepted June 15, 2012

DOI: 10.3892/mmr.2012.968

Abstract. Cerebrovascular injury is one of the three major causes of death and is the leading cause of adult disability. Despite the increasing progress in emergency treatment and early rehabilitation in patients with cerebrovascular injury, treatment options for neurological dysfunction that presents at a later stage are lacking. This study examined the potential of human amniotic mesenchymal stem cell (hAMSC) transplantation in the repair of neurological deficits in an experimental focal cerebral ischemia model. Following the isolation of hAMSCs, growth characteristics and surface antigen expression were observed. Butylated hydroxyanisole (BHA) was used to induce the cultured cells into neuron-like cells, which were identified by immunocytochemistry. The suture model was used to induce focal cerebral ischemia in rats, which were subsequently randomly divided into experimental and control groups for treatment with BrdU-labeled hAMSCs or PBS, respectively. Neurological deficits were assessed following transplantation using the neurological severity score, beam balance test and elevated body swing test. Eight weeks later, rat brain tissue was analyzed with H&E staining and BrdU immunohistochemistry, and the survival and spatial distribution of the transplanted hAMSCs were determined. The hAMSCs proliferated in vitro, and it was found that neuron-specific enolase (NSE) was expressed in neurons, whereas glial fibrillary acidic protein (GFAP) was expressed in astrocytes. The focal ischemia model caused varying degrees of left hemiplegia accompanied by right sided Horner's Syndrome. When examined 1, 3, 6 and 8 weeks later, significant recovery in neurological behavior was detected in the rats treated with hAMSC transplantation compared with the control (P<0.01). BrdU-labeled hAMSCs were concentrated near the graft site and surrounding areas, in certain cases migrating towards the ischemic lesion. Local gliosis and lymphocytic infiltration were not detected. hAMSCs exhibit great potential for proliferation and are induced to differentiate into NSE-expressing neuron-like cells following treatment with BHA. Moreover, hAMSC transplantation may improve neurological symptoms following focal cerebral ischemia.

Introduction

Cerebrovascular injury is one of the three major causes of death and is the leading cause of adult disability. The annual incidence rate in China is 130-300 million, with 60-100 million deaths and 75% of survivors suffering disabilities of various degrees. Despite the increasing progress in emergency treatment and early rehabilitation in patients with cerebrovascular injury, treatment options for neurological dysfunction that presents at a later stage are lacking.

Regenerative medicine and stem cell research have progressed significantly in the 21st century, offering novel routes for the treatment of neurological disorders. Mesenchymal stem cells (MSCs), unlike hematopoietic stem cells, are present in bone marrow. Bone mesenchymal stem cells (BMSCs) have become a progressive research field in modern biology and medicine. MSCs are derived from the mesoderm early in development and may be exploited as an ideal source of seed cells, which exhibit the potential to be induced into osteogenic, chondrogenic and adipogenic cells, or even tendon and adipose tissues (1-4). MSCs are easy to obtain, culture and expand in vitro, and are readily induced into designated tissues. Currently, BMSCs are widely used. However, MSCs are present in extremely low amounts in bone marrow, accounting for 0.01-0.001% of the bone marrow
derived cells (5). Increasing evidence indicates that MSCs with osteogenic potential may be isolated from a diverse range of tissues, including adipose (6) and perinatal tissues, such as umbilical cord (7), placenta (8,9), umbilical cord blood (10,11) and amniotic fluid (12,13), or even fetal blood, bone marrow and liver (14-17).

Placenta, a temporary organ, is important for maintaining maternal and fetal oxygen and nutrients during embryonic development. The full-term placenta comprises amnion and chorion, and our previous findings (18) indicate that MSCs may be obtained and expanded from the amnion (amnion-derived MSCs; AMSCs) and chorion (chorionic MSCs; CMSCs) of placenta (placental MSCs; PMSCs) in vitro; their biological characteristics remain well maintained, similar to those of BMSCs. In addition, a cell bank of PMSCs may be set up in advance for clinical trials, suggesting that PMSCs have a wide application prospect (18).

In this study, we aimed to establish a stable and reliable method for the isolation and amplification of human amniotic mesenchymal stem cells (hAMSCs) in vitro. Following induction into neural cells, the hAMSCs were transplanted into the ischemic tissue of rats subjected to focal cerebral ischemia by middle cerebral artery occlusion (MCAO). The survival, migration and differentiation of the implanted cells and the recovery of neurological function were assessed in rats 1-8 weeks later to examine the potential therapeutic benefit of the hAMSC-derived neuron-like cell transplantation in the treatment of focal cerebral ischemia.

Materials and methods

Cell culture. hAMSCs were isolated from normal post-partum placenta. The amnion and villus layer were bluntly separated and repeatedly washed with D-Hank's solution, including double resistant (100 U/ml penicillin and 100 µg/ml streptomycin). After rinsing, the amnion was cut into several 1x1-mm sections with ophthalmic scissors and digested at 37°C in a water bath for ~30 min with the action of 2.5 µg/ml trypsin (Gibco-BRL, Carlsbad, CA, USA). The digestion of the amnion was terminated with Dulbecco's modified Eagle's medium (DMEM) containing 5% calf serum and filtered through a 200 mesh cell sieve. The filtered amnion products were digested again in a 37°C water bath for ~0.5 h with the addition of 1.0 g/l collagenase II (Sigma-Aldrich, St. Louis, MO, USA). Subsequent termination and filtrations were performed as described above. Finally, the harvested cell suspensions were centrifuged at 1,000 rpm for 5 min and the cell pellet was resuspended in low-glucose DMEM (L-DMEM; Hyclone Laboratories, Inc., South Logan, UT, USA) supplemented with 10% fetal bovine serum (FBS; Hyclone) and 1% penicillin-streptomycin (Invitrogen Life Technologies, Carlsbad, CA, USA). The cells were then plated in 25-cm² culture flasks at a density of 1x10⁶ cells/ml and incubated at 37°C with 5% carbon dioxide. The medium was changed every 2 days. When the established adherent cell colonies reached 70% confluence, they were detached with 2.5 µg/ml trypsin and replated at a ratio of 1:2 in 25-cm² flasks.

Differentiation of hAMSCs. Second or third generation hAMSCs were plated onto 6-well plates. When 60% confluence was achieved, the harvested cells were washed with phosphate-buffered saline (PBS). To induce neural differentiation, the hAMSCs were incubated with serum-free medium containing DMSO (2%) and butylated hydroxyanisole (BHA) (100 µM). The media were changed every 3 days and culturing was continued for 14-21 days. The neural induced cells were then confirmed by neuron-specific enolase (NSE) and glial fibrillary acidic protein (GFAP) immunofluorescence staining.

Immunofluorescence. Immunofluorescence was performed on hAMSCs cultured for 24 h. The cells were grown to 60% confluence on 6-well plates, washed with PBS three times, fixed in 4% paraformaldehyde for 30 min, washed as previously described, permeabilized in 0.3% Triton X-100 for 20 min and then rinsed with PBS three times. The cells were then blocked with goat serum for 20 min, incubated with the appropriate primary antibody in PBS for 2 h at 37°C, washed with PBS three times, incubated with secondary antibodies in PBS for 30 min at 37°C (in the dark) and then viewed under a fluorescence microscope. The following primary antibodies were used: rabbit anti-human NSE (1:500) and rabbit anti-human GFAP (1:500), both from Boster Biological Technology, Ltd). The secondary antibody for immunofluorescence was goat anti-rabbit IgG (1:500; Sigma).

BrdU labeling and preparation of cell suspension for transplantation. Third generation cells from the AMSCs were collected and plated in 25-cm² culture flasks and 6-well plates at a density of 1x10⁶ cells/ml. The cell pellet was resuspended in L-DMEM supplemented with 10% FBS, penicillin-streptomycin (100 µg/ml), 5 ng/ml bFGF and 10 µg/ml BrdU, and then incubated at 37°C with 5% carbon dioxide for 48 h. BrdU-labeled AMSCs were centrifuged at 1,000 rpm for 10 min and the cell pellet was resuspended in PBS at 1x10⁶ cells/µl. Finally, 5-µl cell suspensions were used for cell transplantation.

Animal model. Healthy male Wistar rats, aged 3-4 months and weighing 250-300 g, were obtained from the Schistosomiasis Prevention and Control Center of Jiangsu Province, China. Briefly, the rats were placed in a supine position on an operating table following the intraperitoneal injection of 10% chloral hydrate (4 ml/kg) anesthetic. A blunt dissection of the sternocleidomastoid was made through the middle line neck incision, and the carotid artery (CCA) was isolated and then separated into the right external carotid artery (ECA), internal carotid artery (ICA) and the wing jaw artery. A slipknot was left under the ECA and the wing jaw artery, after threading deeply into all the arteries. The CCA was clamped, a small incision was made in the proximal sidewall of the CCA and a nylon suture filament (0.24 mm) was inserted and advanced to a depth of ~18.5±0.5 mm away from the CCA bifurcation. The suture was removed following a 2-h right MCAO, the ECA was ligated and the skin was sutured.

Animal grouping. Out of 45 rats subjected to focal cerebral ischemia, 13 died and 8 did not exhibit paralysis of the limbs. The remaining 24 rats were randomly divided into two groups (n=12 per group).

hAMSC transplantation. Two weeks after MCAO, the rats were placed in a stereotactic apparatus and the bregma was exposed through a median head scalp incision. Coordinates
Table I. Neurological severity scores.

<table>
<thead>
<tr>
<th>Grading</th>
<th>Score (normal, 0; maximum, 5)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal walk</td>
<td>0</td>
</tr>
<tr>
<td>Flexion of forelimb (raising the rat by the tail)</td>
<td>1</td>
</tr>
<tr>
<td>Circling toward the paretic side (walking)</td>
<td>2</td>
</tr>
<tr>
<td>Falling down to the paretic side (walking)</td>
<td>3</td>
</tr>
<tr>
<td>No spontaneous walking, decreased consciousness</td>
<td>4</td>
</tr>
<tr>
<td>Ischemia-related deaths</td>
<td>5</td>
</tr>
</tbody>
</table>

Neurological function in the rats was assessed using the neurological severity scores (NSS) test. The higher the NSS score, the more severe the ischemia.

Table II. Beam balance test.

<table>
<thead>
<tr>
<th>Grading</th>
<th>Score (normal, 0; maximum, 6)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Balances with steady posture</td>
<td>0</td>
</tr>
<tr>
<td>Grasps side of beam</td>
<td>1</td>
</tr>
<tr>
<td>Hugs the beam and one limb</td>
<td>2</td>
</tr>
<tr>
<td>falls down from the beam</td>
<td>3</td>
</tr>
<tr>
<td>Hugs the beam and two limbs fall down from the beam, or spins on beam (&gt;60 sec)</td>
<td>4</td>
</tr>
<tr>
<td>Attempts to balance on the beam, but falls off (&gt;40 sec)</td>
<td>5</td>
</tr>
<tr>
<td>Attempts to balance on the beam, but falls off (&gt;20 sec)</td>
<td>6</td>
</tr>
<tr>
<td>Falls off: no attempt to balance or hang on to the beam (&gt;20 sec)</td>
<td>6</td>
</tr>
</tbody>
</table>

Neurological function in the rats was assessed using the beam balance test (BBT). The higher the BBT score, the more severe the ischemia.

Preparation of paraffin and frozen sections. Eight weeks after MCAO, the rats were anesthetized intraperitoneally with 400 mg/kg chloral hydrate, perfused transcardially with 4% paraformaldehyde in PBS, and their brains were quickly extracted. An ~2 cm ischemic area of brain tissue, including the lateral ventricles and basal ganglia, striatum and hippocampus, was excised and post-fixed in 4% paraformaldehyde. Coronal brain slices (5-µm) were then consecutively sampled using paraffin sections or frozen sections.

Perl’s Prussian Blue stain for hemosiderin. Sections were transferred to distilled water with xylene and ethanol, placed into the working solution (an equal parts mixture of ferrocyanide and hydrochloric acid) for 15 min, rinsed with distilled water and then with tap water. Sections were then stained with neutral red for 1 min, rinsed well with tap water, dehydrated with ethanol and finally cleared with xylene. Out of 400 slices, every 20th slice was stained using this method to confirm the needle placement and injected sites.

Statistical analysis. Data were presented as the mean ± standard deviation. Comparisons of neurological scores were carried out by ANOVA (F test, q test), using SPSS 10.0. The paired t-test was used for the cell count. In the analysis, a value of P<0.05 indicated a statistically significant result.

Results

AMSC culture and cell phenotype. Adherent cells were observed 4 h after the cells were plated, and clone-like growth was observed 48 h later. The morphology of these cells was similar to that of BMSCs: spindle-shaped with fibroblast-like colonies adhering to the plastic surface. Flow analysis (Fig. 1) showed that the AMSCs expressed the typical MSC markers (CD73, CD105 and CD90), but were negative for hematopoietic markers (CD34 and CD45), the monocytic marker (CD14) and HLA-DR. A large number of BrdU-positive cells were observed using fluorescence microscopy, suggesting successful transplantation of the AMSCs.

Neural induction of AMSCs. Morphological changes, including condensed cell bodies with outgrowth in a few sites, were detected in some of the cells 2 h after incubation (Fig. 2A), with more cells showing these neural cell-like changes 1 h later (Fig. 2B). In addition to the morphological changes, differentiated cells expressed NSE, a marker for neural progenitor cells, and GFAP, a marker for astrocytes (Fig. 2C and D, respectively).

Neurological function score. The rats were tested for neurological function at different time-points using the NSS, BBT and EBST tests (Fig. 3A-C). In each test, the neurological behaviors were markedly improved, and there was a

Figure 1. Flow analysis of AMSCs. (A) Flow analysis of marrow cells and AMSCs. (B) Flow analysis of BrdU-positive cells in AMSC culture.
significant difference between the AMSC-transplanted and the PBS-injected groups.

**TTC and H&E staining.** TTC staining is a standard for the measurement of infarct size and has previously been used for assessment of infarct size resulting from apoptosis and necrosis (6). Normal brain tissue was shown in gray, while the ischemic area was white in TTC staining (Fig. 4D). The ischemic region was observed in the dorsolateral striatum and was lateral to the ischemic hemisphere, which is consistent with the blood supply area of the middle cerebral artery (Fig. 4A). In the ischemic hemisphere, H&E staining indicated a large necrotic area in the 8th week after transplantation. Within this region, there was a significant loss of neurons, with only a few remaining astrocytes, and a marked interstitial edema. The surviving neurons had varying degrees of morphological
changes, with the most significant changes occurring in pyramidal cells, which showed a shrunken cell body, retracted processes and loss of Nissl bodies. Furthermore, the chromatin became cloudy and the nuclear membrane decreased in size (Fig. 4B). Fewer degenerated cells were observed in the AMSC-transplanted area (Fig. 4C).

**Determination of injection sites, needle passage and the BrdU-labeled transplanted AMSCs.** The injection sites and needle tracks were identified in each of the 24 rats using specific hemosiderin staining (Fig. 5A) and showed that 17 rats were injected into the striatum and the other 7 rats in the cerebral cortex. Small amounts of lymphocytic infiltration and glial cell proliferation were observed around the needle tracks (Fig. 5B and C). In the experimental group, BrdU-positive AMSCs near the ischemic lesion were found to be distributed around the needle passages (Fig. 5D), with some cells residing at a distance of 2 mm away. No BrdU-positive cells were observed around the needle passages in the control group.

**Discussion**

Cell replacement therapy has become a developing and promising approach for the treatment of central nervous system injury and disease. In this study, we used the focal cerebral ischemia model in rats and implanted hAMSCs in the ischemic hemisphere using stereotaxic targeting to the striatum or cortex. We observed that cell survival and differentiation of the hPMSCs in the cerebral ischemic rat brain was associated with recovery of neurological function. We found that PMSCs implanted into ischemic tissue in rats resulted in improved neurological function and balance beam test performances relative to the control group. Similarly, histological staining showed PMSC survival within the ischemic region.

Silva et al analyzed the gene expression of MSCs and found that MSCs, not only code the genes of mesenchymal tissue, but also the genes of endothelial and epithelial tissues (19). These results provide a theoretical basis for the potential differentiation of MSCs. MSCs may be used to replace a variety of cells due to their inherent plasticity of cross-system and even cross-germ layer differentiation. Deng et al showed that the bone marrow MSCs of rats spontaneously express neural-specific proteins, such as NSE, β-III tubulin, NF-M and S100-β. In this study, the expression of NSE and GFAP was detected following the induction of the hAMSCs by BHA. Furthermore, placental amnion develop from embryonic ectoderm, thus we speculated that the amniotic MSCs are more readily induced to differentiate into astrocytes and neuronal cells than are MSCs derived from other sources. Therefore, hAMSCs have broad application prospects in the treatment of nervous system damage and repair cell research.

When determining the best time-points for the transplantation of hAMSCs following ischemic injury, it is important to consider the release of toxic neurotransmitters and oxygen-free radicals at the early stage of transplantation, and the effect of scar formation on the growth and differentiation of the transplanted cells at chronic infarction. For example, Li et al found that when cells were transplanted 1 or 7 days after acute stroke, nerve toxins, free radicals and pro-inflammatory mediators led to the further development of ischemic injury and affected the transplanted cells which underwent apoptotic cell death in the ischemic penumbra (21). In addition, inflammation activates microglia and inhibits the growth and survival rate of endogenous neural cells. Fukunaga et al considered the best treatment window for BMSC transplantation to be at least 1 month after the patient experienced a stroke (22). In the present study, we transplanted cells 2 weeks after stroke and found that hAMSCs are dense within the ischemic lesion, suggesting that they migrate and/or proliferate within the injured tissue. Furthermore, we found that 8 weeks after cell transplantation, neurological function was improved compared with the control group.
In this study, we showed that the transplantation of hAMSCs markedly improves neurological recovery following MCAO through stereotaxic injection. Additionally, that the recovery was likely to be associated with the secretion function of the implanted MSCs, as it has been reported that the ratio of cell survival and differentiation reaches approximately 80% in vitro (23) and only 3-10% in vivo (24). However, the mechanism underlying recovery is unclear and should be investigated. However, the approach we have described in this study offers a promising new route for the treatment of neurological disorders, including ischemic stroke.

Acknowledgements

This study was supported by grants from the Major State Basic Research Development Program of China (973 Program: 2007CB512402), the National Natural Science Foundation of China (nos. 30930085 and 31000654). The authors especially thank Dr Shan Jiang and Qun Xue for the valuable suggestions and critical review of this manuscript.

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