Fate of Transplanted Bone Marrow Derived Mesenchymal Stem Cells Following Spinal Cord Injury in Rats by Transplantation Routes

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INTRODUCTION

Cell transplantation for the regeneration of an injured spinal cord would be one of the promising regenerative trials. Cumulative research has demonstrated its feasibility and various stem cells have been tried to protect against the secondary damage and to enhance the regeneration of a damaged spinal cord (1-6). As one trial of this cell therapy, mesenchymal stem cells (MSCs) have been highlighted because they can not only be easily harvested, expanded and transplanted, but they can also be directly harvested and transplanted, which obviates the ethical and immune rejection problems (7-10). However, several issues need to be addressed in order to establish a successful cell therapeutic strategy. Selection of the ideal cell, transplantation method, dosage, and the timing of transplantation are still the questions that should be addressed. Moreover, knowledge about the survival, migration, proliferation and differentiation of the transplanted cells in the injured site is also essential for successful cell therapy for spinal cord injury (SCI).

MSCs are known to have a homing effect and to be neuroprotective following SCI when they are injected in the early stage of SCI (8, 11-13). The suggested neuroprotective effects of MSCs for SCI are that they act as an inducer of neurotrophic factor, a modulator of inflammation. Moreover, they are suggested to be able to replace damaged cells through trans-differentiation (9, 14-16). However, there is little research regarding the fate of the transplanted cells and their neuroprotective effects in different transplantation conditions (17, 18). In the present study, MSCs were delivered via different transplantation routes in a contusive SCI animal model to investigate whether there are any different in terms of the fate of transplanted MSCs and the alteration of microenvironment of the injured site.

MATERIALS AND METHODS

Animal Model and group allocation

All of the surgical interventions and the pre-surgical and post-surgical animal care were provided in accordance with the Lab-
oratory Animal Welfare Act and the Guidelines and Policies for Rodent Survival Surgery, as provided by the Animal Studies Committee of the Catholic University of Korea (IACUC approval No.2011-0174-01). A total of 36 of adult male Sprague-Dawley rats (body weight: 250-300 g each) were used in this study. They were kept under standardized conditions (4 rat/cage, 20-24°C, 45%-65% humidity, and a 12 hr of daily light) and given free access to food and water throughout the study. Rats were randomly assigned to one of the following three groups before operation: the control group (n = 12, SCI only), the intravenous (IV) group (n = 12, SCI + IV administration of MSCs), intraslesional (IL) group (n = 12, SCI + IL administration of MSCs).

After random allocation, the rats were anesthetized with ketamine (50 mg/kg) and xylazine (2 mg/kg, intraperitoneal). Spinal cord injury was made as described previously (19, 20). Briefly, their backs were shaved and then sterilized with antiseptic betadine. Lamincotomies were performed at T9 after exposure of the paravertebral muscles from T8-10. All the spinal contusions were induced by a 25 g-cm contusion using the MASCIS (Multicenter Animal Spinal Cord Injury Study) impactor (a rod weighing 10 g and dropped from a height of 2.5 cm). The 25 g-cm lesion was chosen to evaluate the neuroprotective effect of the experimental trials in severe SCI. Postoperatively, 5 mg gentamicin was administrated intramuscularly. The postoperative care procedures involved manual emptying of the bladder twice a day during the experiment.

Preparation of allogenic mesenchymal stem cells
The femoral bone was used to obtain bone marrow. After anesthesia, the femoral bone was harvested and both ends of the femoral bone were cut. Bone marrow was aspirated with an 18-gauge needle and then, diluted to 25 mL with Dulbecco’ Eagles medium (DMEM) (Sigma, St. Louis, MO, USA) supplemented with 10% heat inactivated fetal bovine serum (FBS) (GibcoBRL, Grand island, NY, USA), 2 mM L-glutamate (Sigma), 100 U/mL penicillin and, 0.1 mg/mL streptomycin (Sigma). The bone marrow aspirates were plated and then incubated in a humidified atmosphere of 5% CO2 at 37°C. For selecting the MSCs, the nonadherent cells were eliminated by replacing the medium 48 hr after cell seeding. For each passage, the cells were plated at about 8,000 cells/cm² and they were grown to confluency.

Phenotypic characterization of the MSC and transplantation
Flow cytometric analysis of the cultured MSC was performed. Briefly, the cells were detached with trypsin-EDTA solution (0.05%, 1 min, Sigma) and washed twice with PBS that containing 0.1% bovine serum albumin. For direct assays, aliquots of cells at a concentration of 1 × 10⁶ cells per milliliter were immunolabelled at 4°C for 30 min with the following antibodies: FITC-conjugated CD 45, PE-conjugated CD 29 and CD 73. MSCs are known to have the immunophenotype of CD 29 and CD 73 and they lack the CD 45 hematopoietic immunophenotype. All the monoclonal antibodies were purchased from Pharmingen/Becton Dickinson (Franklin Lakes, NJ, USA). As an isotype-matched control, mouse immunoglobulin G1-FITC or mouse immunoglobulin G1-PE was used. The labeled cells were analyzed by a FACS Calibur flow cytometer (Becton Dickinson) with the use of CellQuest software.

Before transplantation, cells were labeled with fluorescent membrane-intercalating dye PKH 26 (red fluorescence, 10⁻³ M, Sigma). PKH 26 is known to have the longest in vivo half-life and it is ideal for in vivo cell tracking and cell proliferation studies. For the IV transplantation, 1 × 10⁶ cells in a 0.5 ml total volume were injected through the tail vein 24 hr after SCI and PBS of same volume was injected through the tail vein for the control group. For the IL transplantation, at post-injury day one the injured sites were re-exposed and a concentration of 1 × 10⁶ cells in 10 µL was injected using a Hamilton needle. With the help of a microscope, even administration was done at the caudal and cephalad portions of the injured site. All rats received a daily injection of cyclosporine A (10 mg/kg) intra peritoneally for 5 days starting 2 days before surgery.

Immunohistochemical staining for identifying of the transplanted cells
To evaluate the survival, homing ability and proliferation of the PKH 26 labeled MSCs, 8 rats in each group were randomly assigned to undergo tissue harvesting. At 6 weeks post-injury, specimens that include spinal cord and spleen were obtained after transcardiac perfusion. Spleen was histologically evaluated to observe the transplanted cells’ entrapment by host immune system. To identify the cell type of the transplanted cells, double-labeling studies were performed with the use of primary antibodies to neurons (NeuN, 1:100, Chemicon, Pemecula, CA, USA), oligodendrocytes (CC-1, 1:50, Chemicon) and astrocytes (GFAP, 1:50, Chemicon). The details followed the previously described procedures (20). To quantify the differentiation of the transplanted cells in each condition, the cells that were exactly co-localized by expressing DAPI, PKH 26 and the cell markers NeuN, CC-1, and GFAP were considered as differentiated transplanted MSCs. Two tissue samples from each subject were used and on each slide, six fields were randomly selected and the high-powered images (× 400) were obtained using confocal microscope. The positive cells were counted and the mean numbers of each specimen were recorded and the mean numbers of each group were compared. We also performed staining for type II collagen to investigate the unintended mesenchymal differentiation of the transplanted cells.

Growth factor analysis
Recent studies have demonstrated that stem cell therapy, includ-
ing MSCs for SCI could alleviate secondary damage. The production of growth factors by the transplanted cells is suggested as one of the possible mechanisms of this. Therefore, we evaluated the expression of neurotrophic growth factors. Four rats in each group were used to assess the secretion of growth factors.

At 1 week post-injury, the spinal tissues were dissected and stored at -80°C. Subsequently, the samples were homogenized on ice in RIPA buffer (150 mM NaCl, 50 mM Tris-HCl pH 7.4, 2 mM EDTA, 1% NP-40, 0.1% Triton X-100, 0.1% SDS, 1 mM Na3VO4, 1 mM sodium deoxycholate, 1 mM PMSE, 10 mg/mL aprotinin, and 5 mg/mL leupeptin). The lysate was centrifuged at 16,000 rpm for 10 min at 4°C. The proteins were separated by SDS-polyacrylamide gel electrophoresis and they were transferred to a polyvinylidene difluoride membrane (Hybond-P, Amersham Pharmacia Biotech, Buckinghamshire, UK). The membrane was blocked with 5% fat-free dry milk for 1 hr in Tris-buffered saline (0.1% Tween-20, 20 mM Tris-HCl, 137 mM NaCl, pH 7.4) and then it was incubated overnight at 4°C with the primary antibodies. The antibodies used for immunoblotting were as follows: brain-derived neurotrophic factor (BDNF; 1:500, Santa Cruz, Santa Cruz, CA, USA), and neuronal growth factor (NGF; 1:500; Santa Cruz). After the membranes were washed, they were incubated with secondary peroxidase-conjugated anti-rabbit or anti-mouse antibodies (Amersham Pharmacia Biotech) that were diluted 1:2,000 in Tris-buffered saline with 0.01% Tween 20. An antibody detection system (ECL, Amersham Pharmacia Biotech) was used and the membranes were exposed to X-ray film. The protein band intensities were quantified with a VDS densitometer (Amersham Pharmacia Biotech).

**BBB locomotor rating scale**
The BBB locomotor rating scale was used to evaluate the neurological outcomes over the time course of 6 weeks after SCI (21). The behavior of each animal in an open field was observed and recorded by two researchers. Scores ranging from 0 to 21 were recorded every week after injury.

**Statistical analysis**
All the values in the figures and text are expressed as means ± S.E.Ms. The results were analyzed by one-way ANOVA followed by a Bonferroni post-hoc test for multiple comparisons. A P value less than 0.05 was considered to be statistically significant.

**Fig. 1.** Flow cytometric analysis of cultured cells with CD 45, 73, and 29. The positive expression of CD 73 and CD 29, and negative expression of CD 45 indicate its mesenchymal stem cell lineage.

**Fig. 2.** Type II collagen expression of the transplanted cells were observed. In both transplanted groups, cells with a colocalization of PKH 26 and collagen II were not detected. (A) The IV transplanted MSCs do not express type II collagen. (B) Some type II collagen expression are noted in the IL group, however, no colocalization are found with PKH26 expression (collagen was tagged with green fluorescence, magnification × 400, scale bar 20 µm).
RESULTS

Characterization of MSC

Before transplantation, the third passage cells were evaluated to confirm their phenotype as MSCs. Flow cytometry analysis of the MSCs showed positive cell surface markers in CD 29 and CD 73. The cultured MSCs lacked the expression of CD 45 (Fig. 1). This represents that the cultured cells have characteristics of MSCs.

Table 1. The mean numbers of engrafted and differentiated cells by transplantation routes

| Routes | PKH 26 positive (× 200) | Neuron differentiation (× 400, PKH26+NeuN) | Oligodendrocyte differentiation (× 400, PKH26+CC-1) | Astrocyte differentiation (× 400, PKH26+GFAP) | Other organ (spleen)* |
|--------|------------------------|----------------------------------------|---------------------------------------------|----------------------------------------|----------------------|
| IV     | 30 ± 6.8               | 9.4 ± 0.9                              | 8.2 ± 1.1                                   | 27 ± 1.0                               | ++                   |
| IL     | 47 ± 7.3               | 28.4 ± 2.4                             | 20.4 ± 1.3                                  | 20 ± 1.3                               | -                    |

*PKH26 positive cell in spleen. × 200 and × 400 represent the magnification rates which were used for cell counting. IV, intravenous transplantation; IL, intralesional transplantation.

Fig. 3. Various expressions of neural and glial cell makers of engrafted MSCs. PKH26 positive cells were mainly found at the injured sites. (A) Neuronal differentiation of the transplanted MSCs (n = 4, two tissue samples and six fields in each sample). (B) Oligodendrocyte differentiation of the transplanted MSCs. (C) Astrocyte differentiation of the transplanted MSCs. Arrow indicates co-localization of PKH and GFAP expression. IV transplanted MSCs were mainly expressed the astrocyte differentiation. The proportion of neuronal and oligodendrocyte differentiation were lower than that of IL transplanted MSCs (magnification × 200, scale bar 50 µm).
Identification of the transplanted cells in vivo
In both treated groups, the transplanted MSCs were found at the posterior portion (injured site) of the spinal cord and some scattered cells were also observed at the gray and white matter adjacent to the injured site. Under magnification (× 400), the cells expressing PKH staining coincident with DAPI were counted and quantified. In the control group, there were no cells expressing PKH 26. The mean numbers of PKH 26 positive cells in the IV and IL groups were 30.4 ± 6.9 and, 47 ± 7.3, respectively (Table 1). The numbers of implanted MSCs were greater in the IL group than that in the IV group with statistical significance (P < 0.05). Abundant MSCs were found in the spleen in the IV group but no MSCs were observed in the spleen of the IL group.

Aberrant differentiation of the transplanted cells
To identify an aberrant differentiation of the transplanted cells to mesenchymal lineage, type II collagen staining was performed. Primary antibody to type II collagen (1:100, Lab Vision, Fremont, CA, USA) was used. Alexa Fluor 594 was used for immunofluorescence detection of collagen. Neither of the transplanted groups showed collagen expressing transplanted cells (Fig. 2).

Differentiation of the transplanted cells
To observe any differences in the differentiation of transplanted cells depending on the transplantation routes, double staining was performed and the numbers of cells that were colocalized with DAPI, PHK 26 and other cell markers including NeuN, CC-1 and GFAP were counted. In the IV group, most of the PHK 26 positive cells were colocalized with GFAP labeling and the numbers of cells showing immunofluorescence colocalization with NeuN and CC-1 markers were very low. In the IL group, the transplanted MSCs were evenly colocalized with GFAP, NeuN and CC-1. Table 1 shows the numbers of colocalized expressions of the transplanted cells (Fig. 3).

Neurotrophic factors expression in the SCI lesion
BDNF and NGF levels in the spinal cord tissue (n = 4, each group) were measured. BDNF level in the IL group (mean relative optical density, 1.70 ± 0.2) were slightly increased compared to those in the control group (1.58 ± 0.22) and the IV group (1.39 ± 0.35). However, there was no statistical significance (Fig. 4). NGF level in the IL group (mean relative optical density, 2.4 ± 0.15) was significantly increased compared to the control (2.16 ± 0.04) and the IV group (1.7 ± 0.23) (P < 0.05) (Fig. 5).

Behavioral assessment
All the injured rats manifested complete hindlimb paraplegia immediately after the operation. In all the groups, the rats gradually recovered varying degrees of motor function over the time of observation (Fig. 6). At 6 weeks post-injury, the mean BBB motor scales in the control, IV and IL groups were 6.5 ± 1.8,
11.1 ± 2.1, and 8.5 ± 2.8, respectively. The functional recovery seen in the rats that underwent MSCs transplantation was significantly better than that in the control group (P < 0.05).

**DISCUSSION**

There is no doubt that cell based therapy, including stem cells, is an attractive and promising therapeutic strategy for many clinical conditions that currently lack efficacious treatment. However, there are many issues and concerns to be addressed before its clinical translation. As one of the efforts to address these topics, we tried different transplantation routes of stem cells for SCI. The efficacy and fate of the transplanted cells were observed according to different transplantation routes. In the present study, we used allogenic MSCs. MSCs have been demonstrated to have an ability to “home” into the injured site and trans-differentiate into neural lineage cells (13, 22-24). We were also able to observe this homing and trans-differentiation ability even when the MSCs were applied intravenously at the acute stage of SCI. Moreover, its results were connected to behavioral improvement in the both IV and IL groups compared to the control group.

Studies on applying MSCs application for various central nervous disorders have demonstrated that transplantation of MSCs alleviated further tissue damage and it yielded significant clinical improvement (7, 10, 12, 25, 26). These results were explained by the possibilities of a neuroprotective function and a tissue repair by the transplanted cells. In most of the reported studies, the differentiation of transplanted cells could be observed and to a limited extent, neuron and GFAP positive differentiation were reported (10, 25, 27). However, these studies did not present the differences depending on the transplantation route. A few studies that focused on comparing the efficacy following MSCs transplantation for SCI demonstrated more efficient engrafting of transplanted cells into lesion site when grafting by the intraselenal or lumbar puncture routes (14, 28, 29). These studies demonstrated this difference of efficacy only through examining the engrafting MSCs volume as counted by histological or radioisotope labeling examination. Clinical assessment or the differentiation of transplanted cells has not been addressed. As suggested in many studies, intravenous delivery has inherent concerns of its efficacy. Although the IV route has the advantages of easy and safe delivery, trapping of the transplanted cells in the other organs and the high chance of exposure to an immune reaction limits its clinical utility. In the present study, as predicted and suggested by other previous studies, IV delivery showed a lesser number of engrafted MSCs as compared to that of IL delivery. Many MSCs were observed in the spleen. However, IV delivery showed more effective clinical improvement as compared to that of the control group and the IL group. It is hard to conclude from our results that IV delivery could result in better clinical improvement in the early stage of transplanting MSCs for SCI. However, as other studies have reported, IV delivery could be an effective delivery route for early MSCs transplantation following SCI. Homing of the MSCs to the disrupted blood-spinal cord barrier tissue and avoidance of additional injury that can be caused by intradiscal delivery could account for these results (9, 12, 25).

Homing of MSC into the injured spinal cord is well known and stromal derived factor-1 (SDF-1)/CXCR-4 has recently been demonstrated to take part in the migration of MSCs (11, 30, 31). In our results, even though the absolute number of engrafted cells in the IV group was lower than that of the IL group, the degree of behavioral improvement was better in the IV group. This phenomenon suggests that the neuroprotective effects of early transplanted MSCs do not merely depend on the absolute number of the engrafted cells. Although it is controversial, some studies have suggested the possibility of replacement of damaged tissue by the transplanted MSCs (32, 33). However, the secretion of neurotrophic factors (BDNF, NGF, and VEGF), the modulation of inflammation and immune reactions and enhancement of axonal sprouting in the pathologic condition following SCI have been recently suggested as the primary effects of MSCs transplantation at the early stage, which is beyond their potential to differentiate to form glial and neural lineage cells (8, 27, 32).

And in terms of clinical results, our study has some limitations. IL transplantation needed the secondary surgery which might influence the final clinical results. Further research is needed to clarify the neuroprotective mechanism of MSCs transplantation for early stage of SCI.

In terms of the fate of the transplanted cells, numerous studies have reported that MSCs have the capability to differentiate
into neuronal cell lineages (5, 10, 24). Even though MSCs were intravenously transplanted, the transplanted MSCs expressed neuron or glial cell phenotypes. Our immunohistochemical data also demonstrated that the transplanted PKH26 positive cells were predominantly located at the damaged area. Surprisingly, the phenotype expressions of the transplanted cells are different according to the delivery routes. In the intrathecally transplanted group, the transplanted cells showed NeuN, CC-1, and GFAP positivity. It suggests that the transplanted cells expressed all the neuronal lineage phenotype differentiation. In contrast, the intravenously transplanted MSCs mainly expressed GFAP positivity. NeuN and CC-1 expressing MSCs were rarely observed. With our results, it is hard to conclude that one delivery route is superior to the other. However, the intravenous delivery route also showed effective neuroprotective results and this is supported by the histologic findings and the clinical improvement.

Preferential astrocytic differentiation of the transplanted MSCs has been reported several times (10, 25, 34). Some authors have suggested that differentiation into astrocytes is the default pathway for transplanted MSCs or neuronal stem cells through in vivo study (22, 35). However, of importance is that this predominant astrocyte differentiation of MSCs in models of central nervous system injuries has been associated with favorable functional results. To interpret these results, most authors agree with the hypothesis that differentiation into astrocytes is beneficial for early stage injury through the neuroprotective functions of astrocytes (34, 35). However, astrocytes are known to have a wide variety of biological activities following central nervous system injury (20, 36, 37). In the acute phase of injury, astrocytes have a major role to maintain and restore homeostasis in the injured site and this is also important to protect against further damage. On the other hand, in the chronic phase, reactive proliferation of astrocytes results in gliosis and this gliosis is a major obstacle to regeneration. Therefore, the long term effect of this predominant astrocyte differentiation remains to be established.

In terms of the neurotrophic factors expression, the IL group showed higher BDNF and NGF expression compared to those in the control and IV groups. As suggested by the previous studies, this might be related to the absolute number of the engrafted MSCs. However, in this study, the clinical improvement was not correlated to the absolute number of the engrafted MSCs, and the expression of BDNF and NGF. These findings might be also related to the additional injury during the transplantation in the intrathecally injections.

Before stem cell therapy for SCI can be a successful treatment, many issues need to be resolved including the optimal delivery route and the fate of the transplanted cells. According to our results, we suggest that early delivery of allogenic MSCs following SCI provided favorable behavioral improvement compared to the control group. And then the fate of the transplanted MSCs and expression of neuronal growth factors following MSCs transplantation are different along the transplantation route.

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