Transplantation of Oligodendrocyte Precursor Cells Improves Locomotion Deficits in Rats with Spinal Cord Irradiation Injury

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Abstract

Demyelination contributes to the functional impairment of irradiation injured spinal cord. One potential therapeutic strategy involves replacing the myelin-forming cells. Here, we asked whether transplantation of Olig2+GFP+ oligodendrocyte precursor cells (OPCs), which are derived from Olig2-GFP-mouse embryonic stem cells (mESCs), could enhance remyelination and functional recovery after spinal cord irradiation injury. We differentiated Olig2-GFP-mESCs into purified Olig2+GFP+OPCs and transplanted them into the rats’ cervical 4–5 dorsal spinal cord level at 4 months after irradiation injury. Eight weeks after transplantation, the Olig2+GFP+OPCs survived and integrated into the injured spinal cord. Immunofluorescence analysis showed that the grafted Olig2+GFP+OPCs primarily differentiated into adenosomatous polyposis coli (APC+) oligodendrocytes (54.6±10.5%). The staining with luxol fast blue, hematoxylin & eosin (LFB/H&E) and electron microscopy demonstrated that the engrafted Olig2+GFP+OPCs attenuated the demyelination resulted from the irradiation. More importantly, the recovery of forelimb locomotor function was enhanced in animals receiving grafts of Olig2+GFP+OPCs. We concluded that OPC transplantation is a feasible therapy to repair the irradiated lesions in the central nervous system (CNS).

Introduction

The spinal cord is one of the important dose-limiting normal tissues in clinical radiotherapy. Excessive doses of radiation to the spinal cord result in radiation injury, which is a rare but serious complication of radiotherapy for cancer [1,2]. The pathophysiology involved in irradiation-induced spinal cord injury is demyelination caused by death of oligodendrocytes [3,4]. Reduction in the number of oligodendrocytes was observed as early as 24 hours after X-ray irradiation [5,6]. There is a persistent decline in the number of OPCs from two weeks to three months after X-ray irradiation [7,8]. Remyelination often fails because of primary deficiency in the precursor cells, failure of precursor cells’ recruitment, or incompetence of differentiation and maturation of precursor cells [9]. Persistent demyelination may result in further axonal loss. Such changes could be associated with permanent motor and sensory deficits [10], which become fatal if the damage occurs at the upper cervical level [11]. There are currently no therapeutic approaches that promote remyelination available in a clinical setting.

Cell therapy offers an attractive option as grafted cells may replace the lost ones as well as provide neurotrophic benefits to surrounding tissue [9,12]. To date, primary cultured adult neural stem cells, Schwann cells and olfactory ensheathing cells have been transplanted into the radiation-injured spinal cord [13,14,15,16,17]. However, these cells have limited capacity in producing oligodendrocytes, the primary cell type that is damaged in radiation injury. Oligodendrocyte precursor cells (OPCs), which can be isolated from brain tissues [18,19], offer an alternative source. Nevertheless, they need to be derived from brain tissues. Embryonic stem cells (ESCs) may become a suitable candidate because they are genetically normal, pluripotent, and capable of indefinite replication, and they can be differentiated to all the cell types in the body, including OPCs [20,21,22]. Although the induced differentiation of ESCs is a well-developed approach, there are still problems, especially the purity of target cells [23]. Therefore, it is important to develop strategies for the directed differentiation of ESCs into specialized functional cell types and/or purification of the target cells in vitro before transplantation.

In the present study, we employed a mESCs line that carries a GFP reporter in the locus of Olig2, a transcription factor that is critical for OPCs development [24]. Following transplantation of Olig2+GFP+OPCs into the spinal cord of rat that underwent irradiation injury, the grafted OPCs survived, differentiated to myelinating oligodendrocytes and improved locomotor function of the injured rats.
Materials and Methods

Ethics Statement

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 protocol was approved by the Committee on the Ethics of Animal Experiments of Fudan University (Permit Number: SCXK 2009-0019). All efforts were made to minimize suffering.

Irradiation Injury of the Spinal Cord

Female Wistar adult rats (100–200 g, Shanghai Laboratory Animal, China), were anesthetized by an intraperitoneal injection of 7.5 mg/kg ketamine, 60 mg/kg xylazine and immobilized during irradiation. The rats’ cervical spinal cords were irradiated using a 6 Mev Electron Beam Linear Accelerator (Elekta AB, Stockholm, Sweden). A 22 Gy radiation dose was delivered to a 2 cm × 2 cm radiation field, maintaining cervical 4–5 as the center of the electron-irradiated zone. Radiographs were taken before and after X-irradiation to confirm that all the animals were in the same position. After irradiation, the animals were given subcutaneous saline (5 ml) and housed in a 25°C warm room.

Derivation of Olig2-GFP-OPCs from Olig2-GFP-mESCs

Olig2-GFP-mESCs were the gift of Dr. Su-Chun Zhang (University of Wisconsin, USA) and have been previously described [25]. Olig2-GFP-mESCs were differentiated according to a modified protocol [25]. Briefly, mESCs were trypsinized and placed onto low attachment flasks (Greiner, Germany) in a neural differentiation medium containing DMEM/F12, N2 supplement, leukemia inhibitory factor (LIF), nonessential amino acids, L-glutamine, 2-mercaptoethanol and 10% knockout serum replacement (all from Invitrogen, Carlsbad, CA, USA). Under this condition for 2 days, mESCs formed small aggregates. Retinoic acid (RA, 0.5 μM, Sigma, St Louis, MO, USA) and Purmorphamine (Pur, 0.5 μM, Calbiochem, San Diego, CA, USA) were added at days 2–6 to induce neural progenitors. The aggregates were cultured in suspension in the neural differentiation medium with bFGF (20 ng/ml, R&D systems) and heparin (2 μg/ml, Sigma) for another 6 days. On day 12, the differentiated aggregates were dissociated with trypsin/EDTA (0.05%, Gibco) and plated onto flask coated with Matrigel in modified Bottenstein-Sato medium. The modified Bottenstein-Sato medium contained insulin (10 μg/ml), BSA (100 μg/ml), human transferrin (100 μg/ml), progesterone (60 μg/ml), sodium selenite (40 μg/ml), N-acetyl-cysteine (60 μg/ml), putrescine (16 μg/ml), biotin (10 ng/ml) and cAMP (5 μM) (all reagents from Sigma). T3 (40 ng/ml, Sigma), PDGF-AA (10 ng/ml) and NT3 (5 ng/ml) (all from R&D systems) were added to promote the proliferation of the OPCs.

For immunofluorescence, dissociated spheres (2×10⁴ cells/μl) were plated on coverslips coated with poly-L-lysine and human laminin (Sigma).

For fluorescence activated cell sorting (FACS), cells were harvested at day 12, gently dissociated to single cells and washed with a FACS buffer (PBS, 1%N2, 200 mM L-glutamine, 55 mM β-ME, 50 ng/ml NAC). Cells were analyzed by a Becton Dickinson FACSCaliber with CellQuest Pro (BD Biosciences, San Diego, CA, USA).

For transplantation, FACS-sorted Olig2-GFP-OPCs were resuspended in a fresh culture medium and cultured for 2 days. At day 14, cells were dissociated with accutase (Invitrogen) and prepared at approximately 100,000 cells/μl in artificial cerebrospinal fluid (aCSF). Trypan blue exclusion testing indicated that this preparation consisted of 95% viable cells at time of transplantation.

Cell Transplantation

Transplantation was performed as described [26]. The animals were immunosuppressed using cyclosporine A (10 mg/kg/d, s.c.; Novartis Pharma Schweiz AG, Switzerland) 1 day prior to transplantation and then every day for the duration of the study. Cell transplantation occurred 4 months after irradiation surgery. The animals were anesthetized as above, and the spinal cord was exposed by laminectomy at the cervical 4–5 vertebral level. After immobilization of the spinal cord, a 10 μl Hamilton syringe (Hamilton, Reno, NV) was lowered into the spinal cord using a stereotactic manipulator arm. Cell suspension (2 μl) was injected into two sites along the midline at a depth of 1.2 mm, 4 mm cranial and 4 mm caudal to the lesion epicenter, at a rate of 1.0 μl/min. The needle was removed after 5 min. The males in which the injected solution was observed to exit from the needle track during injection or after withdrawal of the needle were omitted from the experiment. Animals receiving the same surgery and injection of 2 μl aCSF solution (without cells) were served as controls.

Histology

For the histological analysis, the animals were deeply anesthetized with an overdose of 3% pentobarbital and perfused with 0.9% saline following by 4% ice-cold paraformaldehyde prepared with phosphate buffer (0.1 M, pH 7.4). The spinal cord was divided into eight 1 mm blocks that extended 4 mm cranial to and 4 mm caudal to the injury epicenter. Alternate blocks were processed to produce cryostat sections. Some of the cryostat sections were stained with LFB/H&E to examine general morphology and the extent of demyelination. Some of the cryostat sections were used for immunofluorescent analysis of grafted cells.

To quantify the demyelination of injured spinal cord, we used Image J [http://rsweb.nih.gov/ij/index.html] to measure the relative optical density of the dorsal funiculus from all groups. A pre-defined region of interest was selected to encompass the dorsal funiculus. Color images were acquired at the same exposure level, converted to 8-bit gray scale, and the mean density calculated from the threshold pixels excluding the background. Density was analyzed in four regions of the dorsal funiculus per section. Eight sections per spinal cord were analyzed. There were six animals in each group.

Immunofluorescence

Immunostaining was performed on cultured cells or cryosectioned spinal cord using standard protocols [27]. Primary antibodies used in present study included mouse anti-Oct4 (1:2000, Santa Cruz Biotechnology, Santa Cruz, CA), goat anti-Sox2 (1:1000, R&D Systems), rabbit anti-NG2 (1:500, Chemicon International Inc, Temecula, CA, USA), rabbit anti-O4 (1:50, Santa Cruz), rabbit anti-PDGFRα (1:500, Santa Cruz), rabbit anti-MBP (1:500, Abcam, Cambridge, MA), goat anti-GFP (1:200; Abcam), mouse anti-APC (1:1000; Abcam), mouse anti-p75 (1:1000; Chemicon), rabbit anti-GFAP (1:500; Chemicon), and mouse anti-NF (1:2000; Chemicon). Coverslip cultures or spinal cord sections were incubated in a blocking buffer (10% donkey serum and 0.2% Triton X-100 in phosphate buffer saline) for 60 min at room temperature before being incubated in the primary antibodies overnight at 4°C. Fluorescently conjugated secondary antibodies were used to reveal the binding of primary antibodies (1:2000, Invitrogen) and nuclei were stained with 4,6-diamidino-2-phenylindole (DAPI) (0.1 μg/ml, Sigma). The immu-
nofluorescent samples were visualized using a Nikon-Eclipse TE 2000-S fluorescence microscope (Nikon Instruments, Sterling Heights, MI, USA) or a Leica TCS-SP5 laser-scanning confocal microscope (Leica, Germany).

To quantify the differentiation pattern in vitro, the percentage of immunopositive cells was determined by dividing the total number of immunopositive cells by the total number of GFP/DAPI positive cells in each imaging field and then averaging the result from three fields per marker. These fixed fields were randomly selected from biological replicates using "Image J".

To assess the differentiation of grafted Olig2\(^+\)-GFP\(^+\)-OPCs in the injured spinal cord, four coronal sections, 200 \(\mu\)m apart, were taken spanning the lesion epicenter. The percentage of engrafted OPCs that co-expressed NG2, APC, MBP, p75 and GFAP was quantified for the grafted animals.

Electron Microscopy

For electron microscopic processing, the animals were sacrificed by aortic perfusion with isotonic, heparinized saline followed with 4% paraformaldehyde and 1% glutaraldehyde in 0.1 M phosphate buffer. The fixed spinal cords were washed in 0.1 M phosphate

![Figure 1. Radiation-induced demyelination in the dorsal funiculus of cervical spinal cord.](image)

(A) LFB/H&E staining showed normal spinal cord without irradiation. (B) Four months after irradiation, there was a focal demyelinated zone in the dorsal funiculus. (C) Six months following injury, focal necrosis was seen in the dorsal funiculus. (D) Immunostaining for MBP and NF, enlargement of framed area in A, depicted that non-irradiated myelin surrounding the axons (arrows). (E) Four months after irradiation, most of the axons lost myelin (arrows). (F) At six months, axons began to show necrosis. (G) Electron microscopy confirmed the normal structure of myelin. g. Higher magnification of the tissue in the box in G, indicated the compact myelin sheath. (H) Four months after irradiation, most of axons remained demyelinated (asterisk). (I) Six months following damage, axons necrosis was found (asterisk). Bar, 200 \(\mu\)m (A–C); bar, 50 \(\mu\)m (D–F); bar, 1 \(\mu\)m (G, H); bar, 100 nm (g); bar, 2 \(\mu\)m (I).

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buffer and embedded in 3% agar. Transverse sections (40 μm thick) were obtained on a vibratome (Leica, VT1000S, Nussloch, Germany) and mounted onto slides. The sections were washed in PB and subjected to an additional fixation in 2% OsO4 in phosphate buffer for 2 h at room temperature. After sequential dehydration in increasing concentrations of ethanol (50–100%) and pure acetone, the sections were embedded in Epon 812. Ultrathin sections (80 nm) obtained using an ultramicrotome (UI-

Figure 2. In vitro differentiation of Olig2-GFP-mouse mESCs into Olig2\textsuperscript{+}-GFP\textsuperscript{+}-OPCs. (A) Schematic procedure for differentiation of Olig2-GFP-mESCs into Olig2\textsuperscript{+}-GFP\textsuperscript{+}-OPCs. (B) Phase contrast photograph of mESC colonies on mouse embryonic fibroblast. (C) mESCs remained undifferentiated and stained positive for Oct4/Sox2. (D) Phase contrast photograph of Olig2\textsuperscript{+}-GFP\textsuperscript{+} spheres at day 12. (E) GFP\textsuperscript{+} spheres expressed Olig2. (F) At day 12, the percentage of GFP\textsuperscript{+} cells was 76.79\%\pm1.35\%, sorted by FACS method. At day 14, all purified OPCs expressed GFP as well as the OPC markers NG2 (G) and PDGFR\textalpha (H). At day 18, plated cells adopted a typical oligodendrocyte morphology characterized by multiple branches and expressed GFP and MBP (I). (J) Quantification of immunostaining: 81.69\%\pm8.16\% of cells expressed NG2, 78\%\pm4.31\% of cells expressed PDGFR\textalpha, and 68.47\%\pm5.59\% of cells expressed O4. Data in J are expressed as the mean\pmSEM, n = 3 independent experiments, 6 total replicates. Bar, 200 μm (B–D); bar, 50 μm (E–H); bar, 30 μm (I). doi:10.1371/journal.pone.0057534.g002

buffer and embedded in 3% agar. Transverse sections (40 μm thick) were obtained on a vibratome (Leica, VT1000S, Nussloch, Germany) and mounted onto slides. The sections were washed in PB and subjected to an additional fixation in 2% OsO4 in phosphate buffer for 2 h at room temperature. After sequential dehydration in increasing concentrations of ethanol (50–100%) and pure acetone, the sections were embedded in Epon 812. Ultrathin sections (80 nm) obtained using an ultramicrotome (UI-
tracut E, Reicher-Jung, NY, USA) were stained with uranyl acetate and lead citrate, examined and photographed under a transmission electron microscope (Philips CM120, Amsterdam Netherlands).

Behavioral Assessments
The clinical degree of weakness was graded according to a scale developed by Ushio et al. [28,29]: grade 0 = normal; grade 1 = forelimb instability seen only when the animal jumps or runs; grade 2 = mild weakness, but able to run; grade 3 = moderate weakness, able to walk but not run; grade 4 = marked weakness, attempts to walk; grade 5 = severe weakness, purposeless movements of legs or complete paraplegia. All rats were allowed to live for 8 weeks after transplantation.

Statistical Analysis
All data are reported as Mean±SEM. Repeated measures ANOVA followed by the Tukey-Kramer test was used for

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Figure 3. Distribution and morphology of Olig2<sup>+</sup>-GFP<sup>+</sup>-OPCs in irradiated spinal cord at four weeks after transplantation. (A) GFP<sup>+</sup> cells were observed and distributed in the damaged spinal cord. (B) At higher magnification of the boxed area from (A), the GFP<sup>+</sup> cells exhibited typical bipolar OPC morphology. (C) Many grafted Olig2<sup>+</sup>-GFP<sup>+</sup>-OPCs co-expressed the OPCs marker NG2. Bar, 250 μm (A); bar, 100 μm (B); bar, 20 μm (C).

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comparison of clinical grade scores at each time point. The difference between the two was compared using Student’s tests. Statistical analysis was performed using SPSS 16.0. Differences were considered to be statistically significant at \( p < 0.05 \).

**Results**

**Demyelination Induced by Irradiation**

Non-irradiated animals (\( n = 6 \)) did not show any evidence of demyelination (Fig. 1A, D and G). Even though there was no histological evidence of demyelination observed from 1 to 3-month after irradiation, 80% of the animals (\( n = 60 \)) developed a focally demyelinated zone in the dorsal funiculus from the fourth month, as shown by LFB/H&E staining (Fig. 1B). This was confirmed by immunostaining for myelin basic protein (MBP) and neurofilament (NF), showing the presence of many axons but sharply reduced MBP staining (Fig. 1E). Electron microscopy showed many viable demyelinated axons in the dorsal funiculus (Fig. 1H). Six months after irradiation, focal areas of necrosis were apparent in the white matter (Fig. 1C), with MBP loss and axonal degeneration (Fig. 1F). Axon necrosis was also found by electron microscopy (Fig. 1I).

The clinical degree of weakness was consistent with the histological changes. The mean interval between irradiation and manifestation of the first neurological symptoms (grade 1 = forelimb instability seen only when the animal jumps or runs) was 112\( \pm 6.14 \) days. Four months after irradiation, the animals (\( n = 12 \)) that did not manifest neurological symptoms (grade 0 = normal) were omitted from the experiment.

**Generation and Purification of Olig2\(^+\)-GFP\(^+\)-OPCs from mESCs**

We used an established mESCs line (Olig2-GFP-mESCs) in which GFP was knocked into the locus of Olig2, a transcription factor expressed by ventral spinal cord progenitors [24]. In order to identify and purify OPCs, we used a modified protocol established in the Zhang Lab to differentiate mESCs into OPCs [25] (Fig. 2A). The Olig2-GFP-mESCs, expanding as discrete colonies on mouse embryonic fibroblast (Fig. 2B) and expressing Oct4/Sox2 (Fig. 2C), were detached from mouse embryonic fibroblasts to initiate differentiation and grown as free floating embryoid bodies (EB) for 2 days in the medium before being treated with RA (0.5 \( \mu \)M) and Pur (0.5 \( \mu \)M). At day 6, GFP\(^+\) spheres began to appear and by day 12 almost all the spheres were green (Fig. 2D). FACS analysis, confirmed by confocal microscopy, indicated that 76.79\( \pm 1.35 \)% of the total differentiated cells expressed Olig2 and GFP (Fig. 2E, F). To further purify the OPCs, we used FACS based on GFP expression. FACS isolated cells were plated in the presence of FGF2, PDGF-AA and NT3. The majority of GFP\(^+\) cells expressed NG2 (81.69\( \pm 4.31 \)% (Fig. 2G, J), PDGFR\(\alpha\) (78.4\( \pm 4.31 \)% (Fig. 2H, J) and O4 (68.47\( \pm 5.59 \)% (Fig. 2J). Further differentiation of the OPCs for 6 days resulted in generation of highly branched oligodendrocytes, as indicated by positive staining for MBP (Fig. 2I).

**Survival and Migration of Transplanted OPCs within the Spinal Cord**

A total of 24 injured animals received transplantation of Olig2\(^+\)-GFP\(^+\)-OPCs 4 months after irradiation. Four weeks after transplantation, grafted cells revealed by GFP, were found in the spinal cord of all transplanted animals. The animals (\( n = 24 \)) without cell transplantation did not exhibit such labeling. The GFP-expressing cells did not co-express GFAP (J-L) or P75 (M–O). (P) Quantification of GFP\(^+\) cell populations in the spinal cord. Data are expressed as the mean number of cells/spinal cord\( \pm \)SEM (n = 6). GFP\(^+\) cells expressed NG2 (15.1\( \pm 5.53 \)%), APC (54.6\( \pm 10.5 \)% and MBP (40.5\( \pm 3.8 \)%). Bar, 50 \( \mu \)m (A–C); bar, 35 \( \mu \)m (D–O).

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Figure 4. Olig2\(^+\)-GFP\(^+\)-OPCs primarily differentiate along the oligodendrocyte lineage. At eight weeks after transplantation, a certain number of grafted GFP\(^+\) cells still expressed NG2 (A–C, arrows) and most of the grafted GFP\(^+\) cells became APC-positive mature oligodendrocytes (D–F, arrows). Double staining for GFP and MBP in cross-sections further confirmed that GFP-immunoreactive rings were composed of MBP\(^+\) myelin (G–I, arrows). The grafted GFP\(^+\) cells did not co-expressed GFAP (J–L) or P75 (M–O). (P) Quantification of GFP\(^+\) cell populations in the spinal cord. Data are expressed as the mean number of cells/spinal cord\( \pm \)SEM (n = 6). GFP\(^+\) cells expressed NG2 (15.1\( \pm 5.53 \)%), APC (54.6\( \pm 10.5 \)% and MBP (40.5\( \pm 3.8 \)%). Bar, 50 \( \mu \)m (A–C); bar, 35 \( \mu \)m (D–O).

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adopted a typical OPCs morphology, as characterized by bipolar branches (Fig. 3B), and expressed the OPCs marker NG2 (which marks undifferentiated OPCs) (Fig. 3C), indicating that these grafted cells retain the OPCs identity.

Transplanted OPCs Primarily Differentiate into Oligodendrocytes

Eight weeks after transplantation, $15.1 \pm 5.53\%$ of the GFP positive cells were double labeled with NG2, a specific protein for precursor cells (Fig. 4A–C, arrows). Around $54.6 \pm 10.5\%$ of GFP$^+$ cells expressed APC (Fig. 4D–F, arrows), indicating that the majority of grafted OPCs differentiated into mature oligodendrocytes. About $40.5 \pm 3.8\%$ of the grafted cells also expressed MBP, a major constituent of myelin (Fig. 4G–I, arrows). Although several previous studies [30,31] showed that adult OPCs differentiated into Schwann cells after transplantation into the chemically demyelinated spinal cord, our transplanted OPCs did not express p75 (Fig. 4M–O). Other studies suggest that the microenvironment in the injured spinal cord induces grafted neural stem cells to differentiate primarily into astrocytes [14,32]. In the present study, the transplanted OPCs did not differentiate into astrocytes marked by GFAP (Fig. 4J–L). In summation, the staining results revealed that Olig2$^+$-GFP$^+$-OPCs mostly differentiate into mature oligodendrocytes in the injured spinal cord. Over the course of the experiment, tumors, teratomas, or non-neuronal tissue formation were not observed in the transplant recipients.

Figure 5. Transplantation of Olig2$^+$-GFP$^+$-OPCs results in a delayed demyelination and cavitation. (A) LFB/H&E staining showed smaller cystic cavitations in the transplanted cord than those in the sham and the control group in the dorsal funiculus. (B) Optical densities of dorsal funiculus in spinal cord was significantly increased in rats grafted with Olig2$^+$-GFP$^+$-OPCs, as compared to the sham and the control group (*p < 0.05, n = 12). (C) Enlargement of framed area in part a, illustrated immunostaining of MBP and NF, showing axons (green arrow) with or without myelin (red arrow). (D) Electron microscopy confirmed the structure of axons (asterisk). Bar, 200 µm (A); bar, 35 µm (C); bar, 0.5 µm (D).

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Figure 6. Forelimb locomotion is improved after transplantation of Olig2$^+$-GFP$^+$-OPCs. Locomotion function, as defined using the clinical grade score, was significantly reduced in Olig2$^+$-GFP$^+$-OPCs grafted animals at six to eight weeks after transplantation, as compared to the control group (*p < 0.05, n = 12).

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OPCs Transplantation Attenuated Demyelination after Radiation

Eight weeks after transplantation, irradiated spinal cord exhibited obvious demyelination and cavitation (Fig. 5A). However, the degree of demyelination was significantly decreased by OPCs transplantation as compared to the sham and the control group (Fig. 5B). Immunostaining for MBP and NF indicated numerous clusters of axons that maintained integrity (Fig. 5C). Electron microscopic analysis showed myelinated axons (Fig. 5D).

OPCs Grafts Alleviated the Deterioration of Neurologic Function Following Injury

To determine whether transplantation of OPCs improved recovery of function after injury, locomotion was assessed by using the clinical scoring. Six to eight weeks after transplantation, Olig2-GFP+OPCs grafted animals exhibited significantly lower clinical scores as compared to the control group (Fig. 6) (all $p<0.05$). The behavioral results showed that transplantation of Olig2-GFP+OPCs significantly improved forelimb functions.

Discussion

In this study, we have directed mESCs to a purified population of OPCs based on the GFP expression in the Olig2 locus, which differentiate into oligodendrocytes in vitro and in vivo. More importantly, the oligodendrocytes derived from the Olig2-GFP+OPCs express MBP, an essential component of myelin sheath, which corresponds to correction of locomotion deficits in rats with irradiation injury of spinal cord. These results raise the prospect of cell-based therapy as a potential treatment for irradiation spinal cord injury.

Cell transplantation has been attempted for the radiation injured spinal cord. Adult rat neural stem cells differentiate predominantly into Schwann-like cells and generated peripheral myelin after transplantation into the irradiation injured spinal cord [14]. Olfactory ensheathing cells appear to migrate within X-irradiated area, they behave similarly as Schwann cells, which survive poorly and are unable to migrate significant distances when transplanted into either intact or X-irradiated spinal cords [16,33]. OPCs are extremely motile and can differentiate into oligodendrocytes and produce myelin [34,35]. Differentiation of oligodendrocytes from mESCs was explored by Billon and colleagues [36]. One of the greatest challenges facing ESCs research is the derivation of high purity target cells from pluripotent ESCs [23]. In the present study, we employed a mESCs line that carries a GFP reporter in the Olig2 locus. The modified differentiation protocol that we used is unique in that it generates highly enriched OPCs. FACS sorting resulted in an enriched population of OPCs (76.79 ± 1.35%) (Fig. 2F). This provided us with a novel tool to investigate the ability of these cells to remyelinate demyelinated regions of spinal cord injury and determine whether remyelination confers functional benefit.

Most irradiation injury models are made using 40 Gy of X-irradiation which creates acute demyelination in the spinal cord [13,14,16,37,38]. However, the latent period of spinal cord irradiation in human has been reported to vary from 4 months to 4 years after cervical or thoracic irradiation [39]. In experimental animals, irradiation of the spinal cord leads to white matter necrosis within 3–8 months [3]. In the present study, we have used the dosage of 22 Gy to mimic the delayed demyelination of the spinal cord irradiation injury. The demyelination latent period was about 112 ± 14 days, while the white matter necrosis and paralysis that occurred within 6 months of radiation exposure. Therefore, our model is similar to the clinical pathology changes of spinal cord irradiation injury.

Promoting remyelination is an important strategy to treat spinal cord injury. Chronic demyelination predisposes axons to degeneration [40], an irreversible event that is thought to be the major cause of progressive functional decline [41]. Previous studies observed cells with the capacity to myelinate following spinal cord injury [32,34,35,42,43,44,45]. The functional behavioral improvement is correlated with the capacity of the transplanted cells to integrate, differentiate, and myelinate axons at sufficient number [9,46,47]. In our study, we have found that transplanted cells migrate to the injured region and approximately 40% of them differentiate into mature oligodendrocytes expressing MBP. More importantly, immunostaining and electron microscopy analysis reveal the presence of many axons in the areas with Olig2-GFP+OPCs transplantation, suggesting the protection of axonal integrity by transplanted cells. Further study is needed by immunoelectron microscopy to prove remyelination occurred by transplanted cells.

The increased recovery in the transplantation group may also be explained by mechanisms other than remyelination. CNS neurons require multiple signals for optical survival and maturation, and continued oligodendrocyte-derived signals are necessary to maintain neuronal integrity [40]. In addition to their role in myelinating axons, oligodendrocytes release soluble factors, including insulin-like growth factor (IGF-1), glial-derived neurotrophic factor (BDNF), and brain-derived neurotrophic factor (BDNF), that promote neuronal survival, maintain axonal structure, and support synaptic plasticity in surviving axons [49,50,51,52]. The expression and release of trophic factors offer a platform by which oligodendrocytes interact with neurons to form and maintain functional neural circuits in the injured spinal cord. Therefore, neural protection may be another mechanism by which grafted OPCs promote functional recovery.

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Author Contributions

Conceived and designed the experiments: YWP GPZ. Performed the experiments: YS CCX JL XYG. Analyzed the data: YS CCX JL XYG LG YWP. Wrote the paper: YS CCX GPZ.

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