Original

Differentiation of Dental Pulp-Derived MSCs into Hepatocyte-Like Cells and Their Therapeutic Use for Chemical Liver Injuries of Rats

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Abstract: Mesenchymal stem cells (MSCs) derived from dental tissues have gained attention in the field of regenerative medicine, in part because they can be obtained from deciduous or extracted teeth. This study aims to investigate the potential of dental pulp-derived MSCs (DP-MSCs) to differentiate into cells with hepatocyte function and to examine the therapeutic effects of these cells on acute chemical liver injuries of rats. MSC fractions from dental pulp were cultured using specific reagents containing activin A and hepatocyte growth factor (HGF). Albumin, fibrinogen, and urea production were assessed and their specific mRNAs were detected by reverse transcription (RT)-PCR. Therapeutic effects of DP-MSCs on rats with acute chemical injuries induced by concanavalin A (ConA) and D-galactosamine (D-gal) were also investigated. DP-MSCs differentiated into polygonal hepatocyte-like cells (HLCs) that produced albumin and converted ammonium to urea. Importantly, HNF4α, which is a liver-specific transcription factor, was expressed in HLCs, confirming the liver-specific properties of HLCs. Administration of HLCs induced significant improvements in liver function following hepatic injury in rats. Thus, DP-MSCs could differentiate into cells with hepatic function. The potential contribution of these cells in regenerative medicine for refractory liver diseases is expected.

Key words: Mesenchymal stem cell, Dental pulp, Hepatocyte, Fulminant hepatitis, Regenerative medicine

Introduction

Mesenchymal stem cells (MSCs) have attracted substantial clinical attention as one of the important cellular resources of regenerative medicine1, 2. They reside in many kinds of tissues and differentiate into various cell types. Because MSCs do not require the introduction of foreign genes or proteins for differentiation, unlike induced pluripotent stem (iPS) cells, they may be less tumorigenic. They also cause low ethical concern compared with embryonic stem (ES) cells because they do not involve germ cells. MSCs have immunomodulatory properties, resulting in low risk of immune rejection3. Recently, cytotherapy using bone marrow-derived MSCs for acute graft-versus-host disease has been approved4.

Gronthos et al. first succeeded in cloning cells derived from dental pulp and showed that such cells rapidly proliferated and could differentiate into a dentin/pulp-like complex under certain culture conditions, indicating the presence of MSCs in dental pulp5. Since then, dental pulp-derived MSCs (DP-MSCs) have been reported to differentiate into bone, cartilage, adipose tissue, and neural cells6. Because dental pulp is covered by enamel, making it anatomically similar to bone marrow, DP-MSCs have similar cell surface markers to bone marrow-derived MSCs (BM-MSCs). It has even been reported that DP-MSCs were more proliferative than BM-MSCs7.

BM-MSCs were reported to differentiate into cells with hepatocyte characteristics8, 9. Adipose tissue (AT)- and umbilical cord-derived MSCs were also reported to differentiate into a hepatic lineage10-11. Ishkitiev et al. first reported the differentiation of DP-MSCs into cells with a hepatocyte function12. They cultured dental pulp cells obtained at the time of extraction of deciduous or wisdom teeth under the presence of insulin-transferrin-selenium-X and oncostatin M and found that dental pulp cells differentiated into cells with a polygonal hepatocytic morphology and produced albumin. However, thus far, few reports have shown differentiation of DP-MSC into hepatocytes13-15, and the results must be verified by further studies.

Many studies have attempted to treat severe liver disease with MSCs16-19. Clinical trials have been reported, including those using BM- or AT-MSCs15-19. However, there have been no clinical trials using DP-MSCs for decompensated liver cirrhosis or fulminant hepatitis. Because dental-pulp cells are obtained using less invasive methods from extracted deciduous or wisdom teeth, they are considered to be one of the best cellular resources of regenerative medicine, for dental applications as well as for other organs20, 21. In fact, some dental-pulp cell banks are currently in operation. Re-utilization of extracted teeth for treatment of severe diseases is an attractive approach for future clinical application15.

In this study, we attempted to induce differentiation of dental pulp cells into hepatocytes and examined whether administration of these cells could ameliorate hepatic injury and improve survival in a rat model of fulminant hepatitis.
Materials and Methods

Ethical statements

The study was conducted in accordance with the ethical standards for research involving human and animal subjects described in the Helsinki Declaration (The World Medical Association, 2004). All protocols for present study were approved by the Research Ethics Committee of Nippon dental University (Approved No. EC-NG-R-291). Animal experimental protocol was approved by the animal experiment ethics committee of School of Life Dentistry at Niigata, The Nippon Dental University (Approved No. H30-200).

Isolation of primary cell cultures and differentiation into hepatocyte-like cells

Pulps of wisdom teeth were reached through resorbed root canals and extracted using a sterile barbed broach. The pulp was then digested for 30 min at 37°C in 0.2 mg/ml collagenase type I (Wako Pure Chemical Industries Ltd. Osaka, Japan), and 1 mg/ml dispase (Invitrogen, Grand Island, NY, USA). All cell suspensions were seeded into 25-cm² flasks (TPP, Trasadingen, Switzerland) and cultured in growth medium containing DMEM/F12 (Gibco, Grand Island, NY, USA), supplemented with 10% fetal bovine serum (FBS; Sigma-Aldrich, St. Louis, MO, USA), 100 μM glutamax (Sigma-Aldrich), 0.1% minimum essential medium (MEM)-nonessential amino acids (Gibco), 50 U/ml penicillin (Sigma-Aldrich), 50 μg/ml streptomycin (Sigma-Aldrich), and 0.25 μg/ml fungizone® (Gibco). MSC fractions were purified using the Friedenstein method, by which colonies with the greatest proliferating activity were selected and grown. Colonies were differentiated into endodermal cells under the presence of 4 nM human activin A (SBI 800-01, Shinanoki, Japan) and subjected to reverse transcription (RT)-PCR analysis. The study was conducted in accordance with the ethical standards for research involving human and animal subjects described in the Helsinki Declaration (The World Medical Association, 2004). All protocols for present study were approved by the Research Ethics Committee of Nippon dental University (Approved No. EC-NG-R-291). Animal experimental protocol was approved by the animal experiment ethics committee of School of Life Dentistry at Niigata, The Nippon Dental University (Approved No. H30-200).

Transmission electron microscopy

HLCs were fixed in 2.5% glutaraldehyde solution for 2 h at 4°C, washed in 0.1 M phosphate buffered saline, fixed in 1% osmium tetroxide for 30 min at 4°C, dehydrated in ethanol, penetrated in propylene oxide, embedded in Epon and cut into 90-nm sections. Sections were then stained with 3% uranyl acetate and lead citrate and observed under a Hitachi-12 A transmission electron microscope (Hitachi, Co., Ltd. Hitachi, Japan).

Hepatocyte function assays

The presence of the urea cycle may be one of the most characteristic features of mature hepatocytes; thus, we first assayed changes of urea concentrations in HLC culture medium after 24-h exposure of cells to 5 mM/L NH₄Cl (Sigma-Aldrich) in order to clarify the time course of urea production. Urea concentrations were measured colorimetrically (QuantiChrom Urea Assay Kit, Bioassay Systems, Hayward, CA, USA). Nephelometry, followed by the bromocresol green assay, was employed to detect human albumin in the culture medium. Fibrinogen and alanine aminotransferase concentration were commercially measured (SRL, Tokyo, Japan). Periodic acid-Schiff (PAS) staining was performed to detect glycogen storage of cells as follows. After 4% formaldehyde fixation, slides were oxidized in 1% periodic acid for 5 min and rinsed in pure water. Slides were then treated with Schiff’s reagent (Sigma-Aldrich) for 15 min.

RT-PCR

Total RNA from 1 × 10⁴ HLCs or DP-MSCs before differentiation into HLCs was extracted using the RNeasy mini kit (QIAGEN, Hilden, Germany) and subjected to reverse transcription (RT)-PCR analysis. Human liver and lung total RNA (BioChain, Newark, CA, USA) were obtained commercially. A total of 0.5 μg RNA was subjected to RT-PCR using the SuperScript™ One-Step RT-PCR System (Invitrogen) using a thermal cycler (StepOne PCR System, Applied Biosystems, Foster City, CA, USA). The cycle parameters were 94°C for 2 min, followed by 40 cycles of 95°C for 15 sec, 55°C for 30 sec, and 72°C for 1 min. Primers used are listed in Table 1.

Animal experiment protocols

Four-week-old male Sprague-Dawley rats (body weight, 100 g; CLEA Japan Inc., Tokyo) were used. Animals received a single injection of concanavalin A (ConA, Sigma-Aldrich) diluted with phosphate-buffered saline via the tail vein at 20 mg/kg and D-galactosamine (D-gal, Sigma-Aldrich) dissolved in physiological saline and adjusted to pH 7.3 with 1 N NaOH intraperitoneally at 1.2 g/kg. Immediately after administration, HLCs were administered into tail veins. Jaundice and sickness levels of animals were carefully observed. Blood samples were obtained from the tail veins of animals at 24, 48, and 72 h, at which time they were sacrificed.

Immunohistochemistry

Rat liver and lung tissues were fixed in 4% buffered formaldehyde, dehydrated and embedded in paraffin. Sections were stained with hematoxylin-eosin (HE). For immunohistochemistry, sections were treated with the monoclonal antibody to human albumin (ab137885, Abcam, Cambridge, UK) or anti-human mitochondria antibody (clone 113-1, EMD Millipore, CA, USA) at 4°C overnight. Slides were then incubated with peroxidase-labeled anti-rabbit or mouse IgG (Nichirei Bioscience, To-
kyo, Japan), respectively. The color was developed using diaminobenzidine (Nichirei Bioscience).

Methods

Statistical analyses

Significant differences between control and experimental animals were evaluated using Kaplan-Meier survival analysis with the log-rank test. Two-way repeated measures ANOVA was used to compare changes in laboratory test values between control and experimental animals. All statistical procedures were carried out using IBM SPSS Statistics 23 (Chicago, IL, USA). P values <0.05 were considered to be statistically significant.

Results

Histological characterization of HLCs

Differentiation of DP-MSCs into polygonal HLCs occurred within 4 weeks. Phase-contrast microscopic images of DP-MSCs and HLCs showed cells with spindle-shaped fibroblast-like and polygonal hepatocyte-like morphology, respectively (Fig. 1a, b). Transmission electron microscopy showed the presence of desmosome-like structures at the cell membrane contacting adjacent cells and glycogen-like granules in the cytoplasm (arrows) (Fig. 1c, d). PAS staining showed the accumulation of glycogen in cells (Fig. 1e).

Hepatocyte function assays of HLCs

We examined changes of urea concentrations in the culture medium of HLCs containing 5 mM ammonium chloride (Fig. 2Aa). Urea concentrations increased during the time course and peaked at 40 min, suggesting the presence of the urea cycle as a characteristic feature of mature hepatocytes. Along with urea production in the culture medium, human albumin, alanine aminotransferase, fibrinogen, and heparplasmin activities were also detected (Fig. 2A b, c, d, e), confirming the hepatogenic characteristics of DP-MSC-derived HLCs.

Confirmation of mRNA expression of HLCs

In concordance with these hepatocyte-specific biological activities, i.e. mRNA expression of albumin, fibrinogen, and genes responsible for urea production such as glutamine synthetase (GS), arginase 1 (ARG1), and carbamoyl phosphatase synthetase 1 (CPS1) were confirmed by RT-PCR (Fig. 2B b, c, d, e). We especially compared the expression level of albumin mRNA from original DP-MSCs with that from HLCs (Fig 2Bb, lane 3 and 4). Albumin mRNA was detected more strongly in HLCs than in DP-MSCs. In addition, the expression of hepatocyte nuclear factor-4α (HNF4α) mRNA, which regulates many hepatocyte-specific genes, was confirmed in HLCs (Fig. 2Bc).
Figure 3. A. Experimental protocol of HLC transplantation. Immediately following injection of Con A and D-Gal, HLCs or control medium (DMEM/F12) were administered via the tail. Blood samples were taken at 24, 48, 72 h after administration. B. Histological images of liver samples obtained at 72 h showed massive necrosis with lymphocytes infiltration (left: lower magnification, scale bar = 400 μm; right: higher magnification, scale bar = 100 μm. * indicates the area magnified. Areas of massive hepatic necrosis were observed throughout the slides (left). The severe area are roughly indicated by black arrows, and white arrows indicate lymphocytes infiltrations. C. Kaplan-Meier survival analysis of control and HLC-treated animals. D. Changes in mean AST, ALT, LDH, and T.Bil values. Blue and orange lines/bars indicate HLC and control groups, respectively. Data of standard deviation were omitted. Decreases in AST, ALT and LDH from 24 to 48 h were significantly higher in HLC-administered animals.

Figure 2. A. a) A time-course of changes of urea concentrations in the culture medium of HLCs. b) Concentrations of albumin, alanine aminotransferase (ALT), fibrinogen, and hepaplastin obtained from 3rd day cultures. B. Results of RT-PCR analyses of (a) GAPDH, (b) albumin, (c) HNF4α, and (d) fibrinogen, and (e) GS, CPS1, and ARG1. a) Lane 1: HLC, b) Lane 1: buffer control, 2: human liver, 3: DP-MSC, 4: HLC. c) Lane 1: buffer control, 2: human liver, 3: human lung, 4: HLC. d) Lane 1: buffer control, 2: human liver, 3: human lung, 4: HLC. e) Lane 1: buffer control, 2: human lung, 3: human liver, 4: HLC.
Transplantation of HLCs to an animal model of Con A and D-gal-induced hepatitis

Immediately after administration of ConA and D-gal, $1.5 \times 10^6$ HLCs ($n = 14$, experimental group) and DMEM/F12 (Gibco) ($n = 15$, control group) were administered into tail veins (Fig. 3A). Blood samples were obtained from tail veins at 24, 48, and 72 h after transplantation at which point the animals were sacrificed. We first co-administered ConA and D-gal to one rat and it displayed severe hepatic injury (AST: 7,190 IU/l and jaundice with total bilirubin level of 6.73 mg/dl), which was deemed sufficient to evaluate the therapeutic effects of DP-MSC-derived hepatocytes on hepatic injury. Histological findings showed massive hepatic necrosis throughout the slides (severe parts are indicated by black arrows.) with lymphocyte infiltration (white arrows) (Fig. 3B).

Seven of 15 control and 3 of 14 HCLs-treated animals died during the 72 h observation period. Kaplan-Meier survival analysis showed no significant difference between control and experimental groups during this time period (Fig. 3C). Changes in AST, ALT, LDH, and total bilirubin (T.Bil, mg/dl) levels were observed (Fig. 3D). A significant reduction in AST, ALT and LDH were observed from 24 to 48 h in HCLs-treated animals compared with controls ($P$ values <0.05).

We then attempted to examine whether HLCs were detected in rat livers following administration of HLCs, by using antibodies specific for human albumin and mitochondria (Fig. 4A, B). No clearly-positive human cells were found using either antibody. In addition, there were no positive human cells in rat lungs following HLCs administration (Fig. 4C). Human liver sections as positive controls for albumin and mitochondria, lung section for mitochondria are also shown.

Discussion

Liver diseases have a variety of etiologies, including hepatitis viruses, metabolic or genetic disorders, alcohol and drugs. Although recent progress in treatment has contributed to improvement in the prognosis of liver disease, liver transplantation is currently the only cure for patients with irreversible decompensated liver cirrhosis or fatal fulminant hepatitis. Regenerative medicine represents new hope in the treatment of these intractable diseases. There have been many experimental and clinical trials for these diseases using tissue-derived MSCs. There are two main mechanisms for MSCs to exert their pharmacological effects. Many studies have attempted to clarify the ability of MSCs to differentiate and replace damaged resident cells. In most cases, however, engrafted MSCs tend to be short-lived, suggesting other mechanisms for them to exert therapeutic effects. Currently, the primary mechanism of MSCs appears to be mediated through its immunoregulatory properties.

The liver is a central organ of metabolism. Albumin and clotting factors such as fibrinogen are synthesized in the liver. The typical function of the liver includes glycogen synthesis and urea production through the urea cycle. In this study, urea concentrations increased in the culture medium. Along with the detection of albumin, aminotransferase, fibrinogen, and heapaplastin activities, we confirmed the presence of mRNAs for enzymes involved in the urea cycle, as well as albumin and fibrinogen mRNAs, indicating that DP-MSCs acquired hepatic properties under culture conditions conductive to inducing hepatic differentiation. Notably, albumin mRNA was induced under the conditions where DP-MSCs differentiated into HLCs, showing that hepatocyte properties were obtained during the differentiation process from DP-MSCs.
MSCs have a gene expression profile that is similar to that of BM-MSCs\(^{27}\). Kumar et al. showed a higher expression of hepatic markers in DP-MSCs than BM-MSCs\(^{29}\). In this study, HNF4\(\alpha\) mRNA, which is essential for hepatic function and regulates many hepatocyte-specific genes, was detected, confirming the hepatocyte character of these cells\(^{29}\). Because HLCs derived from umbilical cord-MSCs did not express HNF4\(\alpha\)\(^{29}\), our finding might indicate the hepatocyte-prone nature of DP-MSCs\(^{28,30}\). In addition, HNF4\(\alpha\) was also reported to promote definitive endoderm differentiation from human iPS cells\(^{31}\).

Regarding the culture conditions, Lee et al. utilized a two-step induction method to differentiate BM-MSCs into HLCs by first using hepatocyte growth factor (HGF) and basic fibroblast growth factor (bFGF), followed by oncostatin M\(^{32}\). Banas et al. applied similar methods to differentiate adipose tissue-MSCs into HLCs\(^{11}\). These factors are commonly reported to be essential for differentiating MSCs into HLCs. We specifically used activin A and FGF in the first step and HGF in the second step. Activin A is a differentiation-inducing factor that belongs to the TGF\(\beta\) family. As a result, the phenotypes of hepatocytes became very stable, and were not lost even after storage in liquid nitrogen. However, the induced-cells showed characteristics of hepatocytes only in the presence of growth factors, and the phenotypes were lost after the withdrawal of growth factors when other culture methods were taken (personal experience). Han et al. also used Activin A to differentiate DP-MSCs into definitive endoderm, which is an interface during endodermal differentiation, bridging the final stage of hepatic differentiation\(^{33}\).

Shi et al. reported that human BM-MSCs infused via intrahepatic portal veins constituted only approximately 4.5% of pig hepatocytes with D-gal-induced fulminant hepatitis\(^{34}\). They also showed that trans-differentiation of implanted BM-MSCs was close to completion at day 7, concluding that paracrine effects played an important role in MSC-assisted liver restoration. In our study, although not significant, the survival rate of rats infused with HLCs was higher than that of control animals. ConA induces T-cell-mediated liver damage, and D-gal enhances TNF-\(\alpha\) levels of Kupffer cells, inhibiting mRNAs of anti-apoptotic genes\(^{35}\). Therefore, liver injury caused by these reagents are induced by immune reaction and are a good model to evaluate the immunomodulatory effects of MSCs-derived cells. In this study, the reduction of AST, ALT and LDH levels from 24 to 48 h was significantly higher in treated rats compared with control rats. In addition, we could not detect cells that were positive specifically for human albumin and mitochondria in livers of rats infused with HLCs. We speculated that the therapeutic effects of our treatment might be due to paracrine and immunomodulatory effects of DP-MSCs-derived HLCs, considering that a large number of cells infused intravenously might be trapped in the lungs\(^{36}\). However, we were unable to detect human cells in rat lungs; thus, this paracrine mechanisms could not be confirmed. The reason for this may be because the rats were harvested 72 h after cell administration, which might be too late to observe surviving HLCs in the liver and lung tissues. Although there have been no reports demonstrating paracrine function in MSC-derived hepatocytes to ameliorate disease, several reports showed that a paracrine-mechanism was involved in pathogenic improvements by growth-factor-treated MSCs, possibly supporting our hypothesis\(^{36-39}\). In addition, a recent report showed the importance of macrophage polarization in the therapeutic effects of DP-MSCs\(^{38}\).

Dental pulp cells obtained by less invasive methods can be stored at dental pulp-cell banks\(^{39}\). One possible implication of our study may be the utilization of DP-MSCs as an important cellular resource to treat refractory liver diseases.

In conclusions, DP-MSCs are a promising resource for regenerative medicine because of their differentiation potential, biosafety, and relatively non-invasive collection method. We found that DP-MSCs could differentiate into cells with hepatic function. We also found that administration of HLCs led to a possible improvement of liver functions in a rat model of fulminant hepatitis. The contribution of these cells in regenerative medicine for refractory liver diseases can be expected.

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### Conflict of Interest

The authors do not have any commercial affiliation or consultancy that could be construed as a conflict of interest.

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