Safety of Multiple Intrathecal Administrations of Cultured Human Autologous Adipose-Derived Stem Cells in the Patients with Neurological Disorders Including Parkinson’s Disease

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Abstract

Although autologous adipose stem cells have much therapeutic potential for incurable diseases, safety concerns have been constantly raised for clinical use. An intrathecal injection of the stem cells to improve therapeutic efficiency in central nervous system diseases is also relevant due to concerns of adverse reaction. To investigate the safety of multiple intrathecal administrations of autologous adipose-derived mesenchymal stem cells (hAdMSCs), adverse reactions were analyzed by questionnaire survey and blood chemistry examination in 70 patients with Parkinson’s disease and other neurodegenerative diseases who received hAdMSCs multiple times in China and Japan, retrospectively. As results, multiple IT administrations of cultured autologous hAdMSCs were safe, although mild back pain at the injection site in 4 patients and headache, dizziness, fever and asleep legs in other 4 patients. Adverse reactions were disappeared completely without specific treatment within a day. There were no serious adverse reactions during the cell treatment period. No remarkable atypical clinical change was observed in the blood analysis before and after the cell administrations. This study demonstrates that multiple IT administrations of hAdMSCs are safe for clinical use.

Keywords: Autologous adipose-derived mesenchymal stem cells; Intrathecal injection; Safety

Abbreviations: AD: Alzheimer’s Disease; ALT: Alanine Aminotransaminase; AST: Aminotransferase; BMSCs: Bone Marrow Stem Cells; BUN: Blood Urea Nitrogen; CNS: Central Nervous System; DMEM: Dulbecco’s Modified Eagle’s Medium; PBS: Fatal Bovine Serum; hAdMSCs: Human Adipose-Derived Mesenchymal Stem Cell; HbA1c: Glycated Hemoglobin; HBsAg: Antigen of the Hepatitis B Virus; HCV: Hepatitis C Virus; HIV: Human Immunodeficiency Virus; IT: Intrathecal; IV: Intravenous; ND: Neurodegenerative Diseases; PBS: Phosphate-Buffered Saline; PD: Parkinson’s Disease; VDRL: Venereal Disease Research Laboratory; γ-GTP: γ-Glutamyl Transpeptidase

Introduction

Parkinson’s disease (PD) is a neurodegenerative disease (ND) showing the progressive abnormal motor symptoms such as rigidity, tremor, bradykinesia, akinesia and the dopaminergic neuronal loss accompanied by the formation of Lewy body that is primarily composed of aggregated α-synuclein in the brain [1]. Long-term loss of dopaminergic neurons in the nigrostriatal dopaminergic pathway leading to dopamine depletion is closely related to the major motor symptoms in the PD patients [2,3].

Several available PD medications to compensate dopamine levels in the brain improve symptoms at early stage of the diseases but do not stop or slow down the progression of the disease [4]. Moreover, long-term treatment of dopaminergic drugs causes undesirable side effects such as dyskinesia and wearing off [4,5]. Although many more studies are required to clear the cause of dyskinesia, therapeutic drug capable of disease modification would make possible to retard or stop severe and irreversible behavioral impairments without these adverse effects at late stage of the disease [6].

Stem cell-based therapy have been expected as a hopeful disease-modifying approach for the ND including PD and Alzheimer’s disease (AD) [7-9]. Because stem cell transplantation can rescue neuronal loss in the brain by substitution of lost cells, cell combination, release of neurotrophic factors, proliferation of endogenous stem cells, and transdifferentiation [10,11]. Preclinical and clinical studies for stem cells have proven to have therapeutic potential for PD. Neural precursor cells from embryonic stem cells, neural stem cells and bone marrow stem cells (BMSCs) provided therapeutic abilities when they were grafted in animal models of PD [12-14].

In recent animal studies, intravenous (IV) transplantation of human adipose-derived mesenchymal stem cell (hAdMSCs) isolated from adipose tissue showed therapeutic effects in 6-hydroxydopa-induced PD mouse by the recovery of dopaminergic neurons and the restoration of cytosolic mitochondria [15]. Both intracerebral and IV injection of hAdMSCs improved motor abilities by returning dopaminergic cell numbers to normal in 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine-induced PD mouse [16].

Stem cell therapy for the patients is limited because of concerns about safety, in particular carcinogenicity rather than efficacy of stem cells [17,18]. However, our previous retrospective research clearly demonstrated that no seriously concerned adverse effects-such as tumorigenicity were observed after multiple IV administrations of cultured hAdMSCs to normal subjects and patients for 3 years [19].

IT injection of therapeutic drugs has tried to improve therapeutic efficiency in CNS diseases despite the inconvenience of the patient [20]. The problem remains to be solved as an administration route for stem cell therapy. Multiple IT administrations of hAdMSCs would induce unexpected side effects due to direct exposure of cultured stem cells into cerebrospinal fluid. Therefore, it is the most important to confirm safety...
of repeated IT administrations of hAdMSCs before entering main clinical studies.

To identify safety of IT administration further, we analyzed side effects observed in 70 patients who suffered from PD or other ND during repeated stem cell treatments. They received hAdMSCs via IT treatment route from minimum once to maximum 25 times in China and Japan from 2009 to 2017.

Materials and Methods

The subjects participated in the study

Seventy patients who have Parkinson’s disease, dementia or other neurological disorders were recruited in Japan and China. All subjects were Asian, and their age was widely distributed from 52 to 89 years and the average was 71.53 ± 7.73 (Table 1). The study consisted of 38 men and 32 women patients. The subjects were selected according to inclusion and exclusion criteria as below. The case inclusion criteria were as follows:

1. Age 50 years or older.
2. Patients who received hAdMSCs minimum one time.
3. Able to provide informed consent.
4. Clinical diagnosis of Parkinson disease, dementia or other neurological disorders.
5. Patients are healthy enough to undergo the research protocol.
6. Patients living at home or in the community.
7. Able to read and speak Korean.

The case exclusion criteria were as follows:

1. Uncontrolled medical condition requiring immediate treatment that would make a walking trial unsafe for the subject.
2. Current or recent orthopedic disorder that severely limits gait.
3. Any current acute psychosis, alcohol abuse or drug abuse.
4. Clinical trial intervention wishing the last 6 months.
5. Patients living in a continuous care nursing facility.

This study was approved by the institutional review boards at the Biostar Stem Cell Research Institute, Ltd., Seoul, Korea.

Cell preparation from the patients

Human adipose tissues of the patients were gained from abdominal subcutaneous fats. The tissues were gradually stirred with 1 mg/ml of collagenase I for 60 min at 37°C for digesting. After 60 min, the tissues were filtered out through a 100 mM nylon sieve to eliminate cellular debris and centrifuged at 470 g for 5 min to gain a pellet. The pellet was suspended in Dulbecco’s modified Eagle’s medium (DMEM; Invitrogen)-based media containing 0.2 mM ascorbic acid and 10% fetal bovine serum (FBS) which was gained from bovine spongiform encephalopathy free hered and centrifuged again. The Cell pellet was assembled and the supernatant was removed. The part of cell was cultured at 37°C/5% CO in DMEM overnight. After 24h, the cell adhesion strength was inspected by an inverted microscope and removed non-adherent cell by washing with phosphate-buffered saline (PBS). The cell medium was altered as Keratinocyte-SFM (Invitrogen)-based media containing 0.2 mM ascorbic acid, 0.09 mM calcium, 5 ng/ml rEGF and 5% FBS. The cells retained around 5 days for confluence. They were subculture-expanded in Keratinocyte-SFM-based media containing 0.2 mM ascorbic acid, 0.09 mM calcium, 5 ng/ml rEGF and 5% FBS when the confluence arrived 90%. The cells were washed with PBS and thereby FBS was completely removed and demonstrated the albumin measurement limit using a bovine albumin ELISA quantitation kit (Bethyl Laboratories). The procedure for the cell preparation was performed under Good manufacturing practices in the Biostar Stem Cell Research Center.

History of hAdMSCs treatment

The IT injection was conducted in Japan and China according to the guidelines of the Declaration of Helsinki and Tokyo for humans and approved by the institutional review board. 5 × 10⁶ adipose-derived mesenchymal stem cells in 0.3 cc of injectable saline were given in once a month to the patients into the spinal cord over 5 minutes through lumbar spinal tapping. They received the stem cells from minimum once to maximum 25 times in China and Japan from 2009 to 2017.

Measurement

The normal daily dietary intake was recommended during the study. The patients maintained their daily life without any specific activity. The clinical pathology examination was conducted at the first visit to evaluate the physical health status of the subjects before the first IT injection of the cells and after the last IT injection of the cells and the last visits. Examination items included blood chemistry screen; glucose, total protein, albumin, aspartate aminotransferase (AST), alanine aminotransaminase (ALT), γ-glutamyl transpeptidase (γ-GTP), triglyceride (TG), blood urea nitrogen (BUN), creatinine (CRET), and total cholesterol. HbA1c, HbsAg, HCV, HIV, and VDRL were tested, too. Clinically significant abnormal changes between the first visit and last visit were recorded as the adverse reactions of the subjects.

Safety assessment

The investigator determined the severity of each adverse reaction observed during repeated treatments of hAdMSCs and until last visits after final IT administration of the cells. The information about the adverse reactions was identified by non-directive questions to the subjects when the patients visited the hospitals. Furthermore, voluntary reports from the subject were recorded in detail during the test period. They included the date of symptom onset and disappearance, and adverse reaction level.

The safety assessment in the patients was conducted in frequency and severity of adverse reaction and clinical laboratory examination (blood chemistry test).

Results

The subject enrollment and treatment

For safety assessment, we analyzed adverse reactions observed in 70 patients who have the central nervous system diseases including Parkinson’s disease and other neurological disorders (Table 1). They were 38 men and 32 women. Ages are widely distributed from 50 to 80 years (6 patients in 50s, 18 patients in 60s, 38 patients in 70s and 8 patients in 80s). All patients were classified according to the time elapsed since the last IT administration of hAdMSC (Figure 1). Twenty-two patients were less than 1 year after the last IT administration of hAdMSC. Seven patients were less than 2 years. Seven patients were less than 3 years and five patients were less than 4 years. Six patients were less than...
5 years and twelve patients were less than 6 years. Six patients were less than 7 years and one patient was less than 8 years. Four patients were more than 8 years.

Thirty-six patients were within 3 years after the last IT administration of hAdMSC. Thirty-four patients were between 4 years and 9 years. Two patients were over eight years.

Safety assessment

Fifty-four patients had Parkinson’s disease, nine patients had dementia and seven patients had other diseases such as AD, meningitis, depression, cognitive impairment, and brain damage (Figure 2).

Each patient received hAdMSCs multiple times from once to 25 times (once a month) for 9 years (Figure 3). 41 patients received 1~5 times, 18 patients received 6~10 times, 7 patients received 11~15 times, 3 patients received 16~20 times and 1 patient received 20~25 times. The minimum injection number was once, and the maximum injection number was 25 times. Mean injection numbers for all patients were 6 times.

Blood chemistry analysis results were not compared in all patients. The patients who did not agree on the post-treatment test were excluded in the blood chemistry test after IT treatment. The blood chemistry data for 19 patients who agreed on the post-treatment test were analyzed to identify whether IT treatment of the cells could induce adverse reactions.

In blood chemistry analysis before IT administration of hAdMSC (Table 3), there was no abnormal value in total protein level and albumin level. 2 patients were abnormal values in AST levels, 1 patient was in ALT level, 3 patients were in r-GTP levels, 4 patients were in total cholesterol, 4 patients were in TG levels, 2 patients were in BUN levels, 3 patients were in CRTN levels, 2 patients were in HbA1c levels. Every patient was negative in HBsAg, HCV, HIV and VDRL. Average values for all analysis items were in normal range. Blood chemistry analysis after last IT administration of hAdMSC was carried out to check safety profile in the patients who received hAdMSC. 1 patient was abnormal in total protein level, 1 patient was in albumin level, 2 patients were in AST levels, 2 patients were in ALT levels, 4 patients were in r-GTP levels, 5 patients were in total cholesterol, 6 patients were in TG levels, 5

### Table 1: The number, sex, and age of patients enrolled for the safety assessment.

<table>
<thead>
<tr>
<th>Age (years)</th>
<th>50~60</th>
<th>60~70</th>
<th>70~80</th>
<th>80~90</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>4</td>
<td>10</td>
<td>21</td>
<td>3</td>
<td>38</td>
</tr>
<tr>
<td>Female</td>
<td>2</td>
<td>8</td>
<td>17</td>
<td>5</td>
<td>32</td>
</tr>
<tr>
<td>Total</td>
<td>6</td>
<td>18</td>
<td>38</td>
<td>8</td>
<td>70</td>
</tr>
</tbody>
</table>

### Table 2: Adverse reactions observed in the patients during multiple IT administrations of hAdMSCs.

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>None</th>
<th>1~5</th>
<th>6~10</th>
<th>11~15</th>
<th>16~20</th>
<th>21~25</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Headache</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Dizziness</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Fever</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Asleep legs</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>None</td>
<td>37</td>
<td>15</td>
<td>6</td>
<td>3</td>
<td>1</td>
<td>62</td>
<td></td>
</tr>
</tbody>
</table>

### Table 3: Number of IT administrations of hAdMSCs.

<table>
<thead>
<tr>
<th>Number of IT treatments</th>
<th>1~5</th>
<th>6~10</th>
<th>11~15</th>
<th>16~20</th>
<th>21~25</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>1~5</td>
<td>54</td>
<td>9</td>
<td>7</td>
<td>1</td>
<td>1</td>
<td>70</td>
</tr>
<tr>
<td>6~10</td>
<td>18</td>
<td>7</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>38</td>
</tr>
<tr>
<td>11~15</td>
<td>7</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>16</td>
</tr>
<tr>
<td>16~20</td>
<td>5</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>12</td>
</tr>
<tr>
<td>21~25</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>6</td>
</tr>
</tbody>
</table>
patients were in BUN levels, 3 patients were in CRTN levels, 6 patients were in HbA1c levels. Every patient was negative in HBsAg, HCV, HIV and VDRL. Average values for all analysis items were in normal range.

Discussion

Recently, many researchers have investigated the efficacy of MSCs to find new therapeutic way for the incurable, irreversible and fatal diseases including ND [21].

However, clinical use of MSCs has been limited because there have been suggested variable risk factors of stem cell-base therapy. Multiple risk factors resulting from cell characteristics (origin of cells, tumorigenic potential, proliferation capacity), manufacturing and handling (culture procedure, disease transmission, cell line contamination, neoplasm formation, storage and delivery condition), clinical characteristics (administration route, exposure duration, unwanted immune response, unwanted engraftment, toxicity) could contribute to the risks of stem cell therapy in patients [18].

Considering the low viability and retention time of the stem cell, many cells should be administered to obtain better clinical results. A smaller number of cells [23], although there is a need for special surgical procedures to make the patients uncomfortable or other risks.

Multiple studies have been tried to discover the best administration route for improving therapeutic efficiency with a minimized dose amount in animal and human [24]. They have focused to expose the cells more to the target sites by changing carrier vehicles or administration route. Because the higher exposure of hAdMSCs to lesioned area with less cells in the brain compared to other treatment routes would compensate for inconvenience of IT administration. Many evidences suggested that IT injection does not induce noticeable adverse reaction in the animal disease models and human patients. The safety and feasibility of an IT injection of BMSCs were identified in multiple sclerosis patients [23]. The study also showed the efficacy can be seen at lower dose of hAdMSCs than with IV administration by changing of administration route into IT. An IT transplantation of bone-marrow-derived mononuclear cell was feasible and safe in a cerebral palsy patient for 2 years posttherapy assessment period, despite being a very young child [25]. Additional clinical case showed that multiple IT and IV administrations (total 7 times) did not induce any significant uncomfortableness except for a negligible fever during the treatment in 5-year-old girl suffering from cerebral palsy [26]. A retrospective study on cerebral palsy patients who received umbilical cord blood stem cells or BMNCs treatment intraheccally in China and Lebanon [27,28] supported that spinal cord injection might be safe and promising administration route. A large size of clinical study in 105 patients with cerebral palsy did not reveal any noticeable adverse effects by BMSCs transplantation into cerebrospinal cord during follow-up [29]. In case of spinal cord injury patients, several adverse events such as urinary tract infection, headache, nausea, and vomiting were observed during multiple IT administrations of autologous AdMSCs (1 × 10^7 cells × 3) but resolved without special treatment [30]. Bonab et al. demonstrated that IT transplantation of BMSCs could give a favor in terms of dose level compared to IV injection in Multiple Sclerosis patients [23]. Combined IV and IT injections of BMSCs were well tolerated and safe in epilepsy patients [31]. IT treatment of the stem cells was safe in clinical trial for traumatic brain injured patients [32].

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Normal range value</th>
<th>Before IT administration of hAdMSC</th>
<th>After IT administration of hAdMSC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total protein</td>
<td>6.7–8.3</td>
<td>19 0 7.53 ± 0.36 19</td>
<td>16 1 7.40 ± 0.51 19</td>
</tr>
<tr>
<td>Albumin</td>
<td>3.8–5.3</td>
<td>19 0 4.31 ± 0.22 19</td>
<td>16 1 4.26 ± 0.25 19</td>
</tr>
<tr>
<td>Aspartate aminotransferase (AST)</td>
<td>8–38</td>
<td>17 2 26.89 ± 9.31 19</td>
<td>17 2 26.16 ± 10.88 19</td>
</tr>
<tr>
<td>Alanine aminotransaminase (ALT)</td>
<td>4–44</td>
<td>18 1 23.37 ± 11.21 19</td>
<td>17 2 27.32 ± 21.43 19</td>
</tr>
<tr>
<td>Gamma-glutamyl transpeptidase (γ-GTP)</td>
<td>16–73</td>
<td>16 3 33.42 ± 20.88 19</td>
<td>15 4 33.79 ± 24.33 19</td>
</tr>
<tr>
<td>Total cholesterol</td>
<td>125–223</td>
<td>15 4 183.53 ± 37.94 19</td>
<td>14 5 196.74 ± 43.03 19</td>
</tr>
<tr>
<td>TG</td>
<td>45–150</td>
<td>15 4 110.16 ± 52.91 19</td>
<td>13 6 142.21 ± 134.25 19</td>
</tr>
<tr>
<td>BUN</td>
<td>8–20</td>
<td>17 2 16.22 ± 5.10 19</td>
<td>14 5 18.69 ± 4.86 19</td>
</tr>
<tr>
<td>CRTN</td>
<td>0.7–1.3</td>
<td>16 3 1.01 ± 0.24 19</td>
<td>16 3 1.04 ± 0.26 19</td>
</tr>
<tr>
<td>HbA1c</td>
<td>4.0–6.0</td>
<td>16 3 5.90 ± 0.92 19</td>
<td>13 6 5.93 ± 1.18 19</td>
</tr>
<tr>
<td>HBs Ag</td>
<td>Negative</td>
<td>19 0 - 19</td>
<td>19 0 - 19</td>
</tr>
<tr>
<td>HCV</td>
<td>Negative</td>
<td>19 0 - 19</td>
<td>19 0 - 19</td>
</tr>
<tr>
<td>HIV</td>
<td>Negative</td>
<td>19 0 - 19</td>
<td>19 0 - 19</td>
</tr>
<tr>
<td>VDRL</td>
<td>Negative</td>
<td>19 0 - 19</td>
<td>19 0 - 19</td>
</tr>
</tbody>
</table>

Data show patient number in normal or abnormal ranges of parameter values. Average shows mean value for all patients.
not show any abnormal symptoms or pathologic changes related to the IT treatments of human embryonic stem cells in the blood, brain, and spinal cord [34].

We previously disclosed that the IV injection of hAdMSCs is permeable to blood brain barrier in mice models of AD and PD [15,35]. With expectation that IT administration can improve therapeutic efficiency by increasing exposure level in the brain, we have followed up on the patients who received IT treatment of the stem cells. To evaluate the safety and feasibility of multiple IT injections of hAdMSCs in the patients of neurologic disorder, we observed adverse reactions during and after repeated treatments of the cells in China and Japan from 2009 to 2017.

We already confirmed that the IV injection of hAdMSCs (~4×10^8 cells) to the patients suffering from a spinal cord injury did not induce any serious adverse reaction related to the stem cells in our previous clinical study [36]. The retrospective study also supported the safety of repeated IV injections of hAdMSCs (~1×10^6 cells/dose) used in our previous other clinical studies. Since the stem cells are expected to highly and directly be delivered to lesion site in the CNS by IT treatment [20,37], the effective dose will be determined at lower dose without any adverse reactions than the IV dose in the clinic.

Number of IT administrations of hAdMSCs to the patients was varied from 1 to 25 in this study. It is difficult to prove which amount and times of doses are the best option now. However, we identified that multiple IT treatments might be safe and feasible, based on the results that nobody appealed serious adverse symptoms and uncomfortableness during IT treatment.

Some mild adverse reactions such as back pain, headache, dizziness, and fever were observed during treatment period. The symptoms appeared immediately or within 1 day after IT treatment but disappeared spontaneously without specific treatment. Back pain may most likely result from penetration of the needle into the spine. Short-term adverse symptoms such as headache, dizziness, and fever were observed in other clinical trials for the stem cells commonly, but they were not serious enough to stop the studies [28,38] as in this study. Our study additionally supports safety of IT treatment of the stem cells by showing a suitable blood evaluation profile in the patients who received hAdMSCs. There was no statistical difference in blood chemistry analysis between before and after IT treatment. Three patients in the abnormal range were added than before the IT treatments of the cells. Because the patients are older, and the change is insignificant, it cannot be judged as an adverse reaction. These results indicate that multiple IT treatments of hAdMSCs might be safe in the clinic. There is a defect that the blood analysis results are not compared for all the patients who received IT treatment of the cells. Additional blood chemistry evaluation is required for the more robust evidence in the follow-up study.

Sixty-four patients were over 60-years-old. Fifty-four patients at early and late stage suffered from PD were involved in this study. Healthy condition should be maintained during the treatment period because long-term treatment is required to show therapeutic effects in the neurodegenerative diseases (ND). Considering most of ND patients are elderly people, the evidence that there are no adverse events in the PD patients would give a better opportunity for IT route for the administration.

We have continuously monitored adverse reactions from one year to over eight years after the last dose of the cells. However, none of the patients reported any unusual side effects or abnormal symptoms related to the cell treatment. IT treatment is expected in the CNS diseases that the less cells may be effective as blood brain barrier penetration is increased compared to IV and oral treatments. Moreover, IT route will save the cost and time required for treating the diseases because the therapeutic effect may be seen earlier than other treatment routes.

**Conclusion**

The IT treatment-induced inconvenience in our clinical study as well as in other studies was not serious or severe as predicted. Our study has a little limitation that ages and number of treatments in patients who received the cells were in wide range. However, it should be good and helpful evidence that there were no adverse reactions and inconveniences directly related to multiple IT injections of the cells to spine for long-term period.

In conclusion, this study strongly suggests that multiple IT administrations of the autologous adipose stem cells are safe for clinical use.

**References**


