Cartilage repair and inhibition of the progression of cartilage degeneration after transplantation of allogeneic chondrocyte sheets in a nontraumatic early arthritis model

Naoki Takatori 1, Masato Sato*, Eriko Toyoda, Takumi Takahashi, Eri Okada, Miki Maehara, Masahiko Watanabe

Department of Orthopaedic Surgery, Surgical Science, Tokai University School of Medicine, 143 Shimokasuya, Isehara, Kanagawa 259-1193, Japan

Abstract

Introduction: Using a rat model of nontraumatic early arthritis induced by intra-articular administration of low-dose monoiodoacetic acid (MIA), we transplanted allogeneic chondrocyte sheets and examined the effects on tissue repair.

Methods: MIA (0.2 mg/50 μl) was injected into the right knee of 20 male Wistar rats. Four weeks later, rats were randomly allocated into three groups: Group A was examined 4 weeks after administration of MIA; Group B, 8 weeks after MIA injection and chondrocyte sheet transplantation, and Group C, 8 weeks after MIA injection but without chondrocyte sheet transplantation. Allogeneic chondrocyte sheets were transplanted into the right knee of Group B rats. Pain was assessed as the weight distribution ratio of the damaged to undamaged limb. The OARSI score was used for histological scoring.

Results: The limb weight distribution ratio indicated significantly less pain in Group B. Histological scoring showed significant differences in cartilage repair and inhibition of the progression of cartilage degeneration between Groups B and C, but not between Groups A and B, or Groups A and C.

Conclusions: These findings suggest that, in this rat model of nontraumatic early arthritis induced by low-dose MIA injection, allogeneic chondrocyte sheet transplantation induces cartilage repair and suppresses cartilage degeneration.

1. Introduction

Osteoarthritis (OA) is one of the most prevalent joint diseases, especially in mature women [1]. The condition is characterized by degenerative changes in articular cartilage, which cause pain and malfunctions as it progresses [2,3]. Structural changes include loss of articular cartilage, subchondral bone sclerosis, and formation of cysts and osteophytes [4]. The low cellularity and avascular nature of articular cartilage means that its capacity to self-regenerate after injury or degeneration is limited [5]. Mild OA is treated conservatively with oral administration of nonsteroidal anti-inflammatory drugs, intra-articular injection of corticosteroid [6] and/or hyaluronic acid [7], and exercise therapy [8]. However, surgical treatment may be required as the disease progresses. The main treatments are high tibial osteotomy and total knee arthroplasty.

Various arthritis models, such as surgical intervention and intra-articular drug injection, have been used to study cartilage regeneration. The anterior cruciate ligament resection model is the surgical intervention model used most frequently. In this model, anterior cruciate ligament resection triggers cartilage degeneration, subchondral bone sclerosis, and osteophyte formation, which mimic the pathological changes observed in human OA [9–11].
Intra-articular injection of monooiodoacetic acid (MIA) is one method for inducing OA experimentally. MIA inhibits glyceraldehyde-3-phosphate dehydrogenase activity in chondrocytes and causes cell death and decreased proteoglycan content. The reaction is simple, rapid, and reproducible [12–15]. Some reports indicate that the OA caused by MIA-induced chemical induction is different from human OA [16,17]. On the other hand, Sangeetha et al. and others have reported histological and morphological changes in the MIA model similar to those in human OA [18–23]. Udo et al. [24] reported that the changes caused by MIA depend on the dose and timing, and that low-dose administration is useful for evaluating the pathology of OA, pain mechanisms, and therapeutic effects. Mohan et al. [25] reported that the low-dose MIA-induced arthritis model is suitable for studying nontraumatic OA and the therapeutic effects on cartilage and bone.

We have transplanted chondrocyte sheets into areas showing traumatic changes in an articular cartilage damage model in various animals and have confirmed its effectiveness [26–28]. The purpose of this study was to determine whether the repair or inhibition of the degeneration of damaged cartilage can be obtained by transplanting allogeneic chondrocyte sheets in a nontraumatic arthritis model of OA induced by intra-articular injection of low-dose MIA.

2. Materials and methods

All procedures using animals in this study were performed in accordance with the Guide for the Care and Use of Laboratory Animals (NIH Publication No. 8023, revised 1978) published by the National Institutes of Health, USA, and the Guidelines of Tokai University on Animal Use. This animal experiment was approved by the Animal Committee of Tokai University (permission number 171059).

2.1. Animals

Twenty male Wistar rats (CLEA Japan Inc., Tokyo, Japan) at 8 weeks of age and three male Lewis rats (Charles River Japan Inc., Kanagawa, Japan) at 6 weeks of age were used for the study. The animals were housed under normal conditions for 2 weeks before the start of the experiments to acclimatize them to the environment. One to two rats were housed per cage in sterile conditions, and rodent chow and water were allowed ad libitum.

2.2. Chondrocyte sheets

2.2.1. Harvest of chondrocytes and fabrication of chondrocyte sheets

To prepare allogeneic cell sheets for transplantation, three 6-week-old male Lewis rats were used. The rats were sacrificed by an overdose of 50% isoflurane. A medial parapatellar incision was made in both legs, each patella was dislocated laterally, and the collateral ligaments and the anterior and posterior cruciate ligaments were severed. The joint capsule on the hip joint side was incised, and the femur was extracted. The articular cartilage tissue was collected with a scalpel from the bilateral femoral head and incised, and the femur was extracted. The joint capsule on the hip joint side was incised, and the femur was extracted. The collected cells were seeded at a density of 1 × 10^4 cells/cm^2 in a 150 cm^2 culture flask in DMEM/F12 supplemented with 20% FBS and 1% AB, and incubated at 37 °C. After 3 days, 100 μg/ml ascorbic acid (Nissin Pharmaceutical, Yamagata, Japan) was added to the medium, and the medium was replaced every 3 or 4 days. Cells were passaged once when they reached confluence and then cryopreserved.

To fabricate cell sheets, cells were thawed, passaged once, and then seeded onto UpCell® inserts (CellSeed Inc., Tokyo, Japan) at 1 × 10^5 cells/cm^2. Ascorbic acid was added, the medium was replaced every 3 or 4 days, and the cells were cultured for 14 days. To harvest the cells using the characteristics of the UpCell insert, the culture temperature was lowered to <25 °C, which caused the chondrocytes to detach, after which they were preserved for further analysis.

2.2.2. Cell counting

The cell sheets were washed in Dulbecco’s phosphate-buffered saline (PBS; Gibco). The sheets were then incubated in TripLE Express® (Gibco) at 37 °C for 40 min and centrifuged at 1500 rpm for 5 min. The cell sheets were resuspended in 0.25 mg/ml collagenase P (Roche, Basel, Switzerland) at 37 °C for up to 5 min and then centrifuged at 1500 rpm for 5 min. Finally, the isolated cells were resuspended in DMEM/F12, and the cells were counted using a trypan blue exclusion assay.

2.2.3. Histological staining

The collected cell sheets were fixed with 20% formalin (Wako Pure Chemical, Japan) and embedded in paraffin wax. The tissue was cut into 3 μm sections, which were then deparaffinized according to standard procedures, and stained with hematoxylin–eosin. The sections were stained with Safranin O in a 0.08% fast green aqueous solution and 0.1% Safranin O aqueous solution. The sections were also stained with a 0.05% toluidine blue solution.

2.2.4. Immunohistochemical staining

Paraffin-embedded tissue was sectioned at 3 μm and deparaffinized using standard procedures. The sections were stained for collagen type I with goat anti-type I collagen-UNLB (Southern Biotech, USA). The sections were washed with PBS and processed using an ImmPRESS Anti-goat Ig Reagent Kit (Vector, USA) and the slides were sealed. Sections for collagen type II staining were deparaffinized and then reacted with Anti-IκB (II) purified IgG (Daiich Fine Chemical, Japan). The sections were washed with PBS and processed using an ImmPRESS Anti-mouse Ig Reaction Kit (Vector), and the slides were sealed.

2.2.5. RNA isolation, cDNA synthesis, and quantitative real-time PCR

Three samples of passage 1 chondrocytes and three samples of cultured cartilage cell sheets were analyzed. The recovered chondrocyte sheets were crushed at 1500 rpm for 3 min using SHAKE Master Neo (Bio Medical Science, Japan). Total RNA was extracted using an RNeasy Mini Kit (Qiagen, Germany) according to the manufacturer’s instructions. First, first-strand cDNA was synthesized using 1 μg of total RNA and the enzyme, buffer, and primer of a Quantitect Reverse Transcription Kit (Qiagen) and elongated using a GeneAmp PCR system 9600 (Thermo Fisher Scientific, USA) at 42 °C for 15 min and 95 °C for 3 min. Quantitative real-time polymerase chain reaction (qPCR) was performed using an Applied Biosystems 7300 Real-Time PCR system (Applied Biosystems, USA). The primers were as follows for TaqMan® Gene Expression Assays (Applied Biosystems, USA): Col1A1 (Rn01637087_m1), Col2A1 (Rn01463848_m1), Sox9 (Rn01751070_m1), Acan (Rn00573424_m1), ITGa10 (Rn01533928_m1), Mmp13 (Rn01448194_m1), Lect1...
to expose the patellar tendon. The knee joint was shaved and disinfected. Under sterile conditions, a 1 cm incision was made in the skin via the medial parapatellar approach to expose the patellar tendon. The knee joint was flexed 90°, and MIA (0.2 mg dissolved in 50 μl of physiological saline; Sigma—Aldrich, St. Louis, MO, USA) was injected into the knee joint through a 27 G needle. We considered the rise of joint pressure due to intraarticular administration, so the same volume of saline was injected into the left knee joint using the same method. After injection, the skin was sutured.

2.3. Induction of OA by MIA

We used 20 male Wistar rats at 8 weeks of age. With the animal under anesthesia with isoflurane and oxygen inhalation, both knees were shaved and disinfected. Under sterile conditions, a 1 cm incision was made in the skin via the medial parapatellar approach to expose the patellar tendon. The femur and tibia were fixed in 20% formalin (Wako Pure Chemical) for 5–6 days. Each section was degreased and demineralized with acetone solution (Wako Pure Chemical) and ethylenediaminetetraacetic acid (EDTA) solution (Wako Pure Chemical). It was fixed again with 20% formalin, decalcified with EDTA solution, and embedded in paraffin wax. In the rat, the load distribution on the medial side of the femur is higher than on the lateral side [23,31,32]; therefore, the section of the medial femoral condyle was cut in the sagittal direction (3 μm sections). The sections were stained with Safranin O, collagen type I staining, and collagen type II staining in the same protocol. Microscopic images were captured under a BZ-9000 Bio-revo fluorescence microscope (Keyence, Japan).

Bone samples from Groups A, B, and C were analyzed histologically. We evaluated the distal part of the femoral articular cartilage and the intermediate region of the articular cartilage of the tibia based on the cartilage region determination method of Nomura et al. [33]. The Osteoarthritis Research Society International (OARSI) OA cartilage index system was used to score the histopathology, as established by Pritzker et al. [34]. The scoring system evaluates the grade according to the depth of cartilage damage and the stage according to the extent of damage over the joint surface. The score for grade × stage is calculated and can range from 0 to 24, with higher scores indicating more advanced cartilage degeneration. We show the OARSI score (Table 1). Scoring was performed by three single-blinded examiners.

2.6. Histological evaluation

Four weeks after MIA injection, six rats in Group A were sacrificed, and 4 weeks after transplantation of chondrocyte sheets (8 weeks after MIA injection), eight rats in Group B and six rats in Group C were sacrificed. The knee joint was exposed using the medial parapatellar approach and the collateral ligaments and anterior and posterior cruciate ligaments were severed. The femur and the tibia were then separated.

The femur and tibia were fixed in 20% formalin (Wako Pure Chemical) for 5–6 days. Each section was degreased and demineralized with acetone solution (Wako Pure Chemical) and ethylenediaminetetraacetic acid (EDTA) solution (Wako Pure Chemical). It was fixed again with 20% formalin, decalcified with EDTA solution, and embedded in paraffin wax. In the rat, the load distribution on the medial side of the femur is higher than on the lateral side [23,31,32]; therefore, the section of the medial femoral condyle was cut in the sagittal direction (3 μm sections). The sections were stained with Safranin O, collagen type I staining, and collagen type II staining in the same protocol. Microscopic images were captured under a BZ-9000 Bio-revo fluorescence microscope (Keyence, Japan).

Bone samples from Groups A, B, and C were analyzed histologically. We evaluated the distal part of the femoral articular cartilage and the intermediate region of the articular cartilage of the tibia based on the cartilage region determination method of Nomura et al. [33]. The Osteoarthritis Research Society International (OARSI) OA cartilage index system was used to score the histopathology, as established by Pritzker et al. [34]. The scoring system evaluates the grade according to the depth of cartilage damage and the stage according to the extent of damage over the joint surface. The score for grade × stage is calculated and can range from 0 to 24, with higher scores indicating more advanced cartilage degeneration. We show the OARSI score (Table 1). Scoring was performed by three single-blinded examiners.

2.7. Statistical analysis

The results are expressed as the mean ± standard deviation (SD). Gene expression levels in the cell sheets were analyzed using the Mann—Whitney U test. A p-value < 0.05 was considered to indicate a significant difference; however, changes in gene expression <2-fold are considered to be within the physiological limit and, therefore, such changes were considered to be unchanged despite being statistically significant. The damaged limb weight distribution ratio and histological assessment as measured by the OARSI score were evaluated using analysis of variance. The Tukey—Kramer test was used for post hoc testing. A p-value < 0.05 was deemed to be significant.

3. Results

3.1. Sheet thickness, cell count, and tissue evaluation of chondrocyte sheets

The macroscopic and microscopic findings for cell sheets are shown in Fig. 2A and B. Chondrocyte sheets contained cells at a high density, and these cells were easily harvested and manipulated without tearing after 14 days of culture. Chondrocyte sheets contained an average 1.32 ± 0.28 × 10⁶ cells and had a survival rate of

\[
\frac{\text{damaged limb load (g)}}{\text{(undamaged limb load (g) + damaged limb load (g))}} \times 100
\]

After MIA injection, the weight distribution was measured 14 times on days 1, 7, 14, 21, 28, 29, 31, 35, 38, 42, 45, 49, 52, and 55. Fig. 1 shows the schedule for MIA administration, transplant surgery, and load measurement.

2.5. Pain evaluation

An incapactance meter (BrainScience Idea Co., Ltd, Japan) was used to detect changes in the distribution ratio of the damaged limb to the undamaged limb, and the ratio served as the gauge for evaluating pain. This device is used widely to investigate pain and pain-alleviating effects [27,29,30]. The measurements were made when the rat’s hind legs were both positioned over the platforms and the rat was stationary. The weight distribution of both hind legs was measured 10 times, and the following formula was used to calculate the limb weight distribution ratio (%).

\[
\text{Limb weight distribution ratio } (\%) = \frac{\text{damaged limb load (g)}}{\text{(undamaged limb load (g) + damaged limb load (g))}} \times 100
\]
81.3% ± 2.4% and an average thickness of 29.53 ± 2.23 μm. The sheets formed a thick structure with integrated layers, which were examined histologically and by staining the cartilage matrix with Safranin O and toluidine blue.

3.2. qPCR of chondrocyte sheets

Gene expression in three chondrocyte sheets was evaluated and compared with that in passage 1 chondrocytes (Fig. 2C). The relative gene expression of chondrocytes was set at 1.0; the relative values were 1.5 times for Col1A1, 4.73 times for Col2A1, 1.6 times for Sox9, 3.10 times for Acan, 1.23 times for ITGa10, 1.83 times for Mmp13, 58.63 times for Lect1, and 9.57 times for Comp. This analysis confirmed an increase in the expression of all genes analyzed.

3.3. Pain evaluation

The normal limb weight distribution ratio is 50%. Fig. 3A shows the changes in the weight distribution ratio with time after MIA injection in each group. Up to 4 weeks after MIA injection, Group A was sacrificed, and Groups B and C were subjected to cell sheet transplantation surgery. At 8 weeks after MIA administration, Group B and C were sacrificed.

Table 1

<table>
<thead>
<tr>
<th>Grade (key feature)</th>
<th>Stage 1 (&lt;10%)</th>
<th>Stage 2 (10–25%)</th>
<th>Stage 3 (25–50%)</th>
<th>Stage 4 (&gt;50%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grade 1 (surface intact)</td>
<td>1</td>
<td>2</td>
<td>3</td>
<td>4</td>
</tr>
<tr>
<td>Grade 2 (surface discontinuity)</td>
<td>2</td>
<td>4</td>
<td>6</td>
<td>8</td>
</tr>
<tr>
<td>Grade 3 (vertical fissures, clefts)</td>
<td>3</td>
<td>6</td>
<td>9</td>
<td>12</td>
</tr>
<tr>
<td>Grade 4 (erosion)</td>
<td>4</td>
<td>8</td>
<td>12</td>
<td>16</td>
</tr>
<tr>
<td>Grade 5 (denudation)</td>
<td>5</td>
<td>10</td>
<td>15</td>
<td>20</td>
</tr>
<tr>
<td>Grade 6 (deformation)</td>
<td>6</td>
<td>12</td>
<td>18</td>
<td>24</td>
</tr>
</tbody>
</table>

Score = grade × stage

81.3% ± 2.4% and an average thickness of 29.53 ± 2.23 μm. The sheets formed a thick structure with integrated layers, which were examined histologically and by staining the cartilage matrix with Safranin O and toluidine blue.

3.2. qPCR of chondrocyte sheets

Gene expression in three chondrocyte sheets was evaluated and compared with that in passage 1 chondrocytes (Fig. 2C). The relative gene expression of chondrocytes was set at 1.0; the relative values were 1.5 times for Col1A1, 4.73 times for Col2A1, 1.6 times for Sox9, 3.10 times for Acan, 1.23 times for ITGa10, 1.83 times for Mmp13, 58.63 times for Lect1, and 9.57 times for Comp. This analysis confirmed an increase in the expression of all genes analyzed.

3.3. Pain evaluation

The normal limb weight distribution ratio is 50%. Fig. 3A shows the changes in the weight distribution ratio with time after MIA injection in each group. Up to 4 weeks after MIA administration, Group A was sacrificed, and Groups B and C were subjected to cell sheet transplantation surgery. At 8 weeks after MIA administration, Group B and C were sacrificed.

Table 1

OARSI osteoarthritis cartilage histopathology scoring.

<table>
<thead>
<tr>
<th>Grade (key feature)</th>
<th>Stage 1 (&lt;10%)</th>
<th>Stage 2 (10–25%)</th>
<th>Stage 3 (25–50%)</th>
<th>Stage 4 (&gt;50%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grade 1 (surface intact)</td>
<td>1</td>
<td>2</td>
<td>3</td>
<td>4</td>
</tr>
<tr>
<td>Grade 2 (surface discontinuity)</td>
<td>2</td>
<td>4</td>
<td>6</td>
<td>8</td>
</tr>
<tr>
<td>Grade 3 (vertical fissures, clefts)</td>
<td>3</td>
<td>6</td>
<td>9</td>
<td>12</td>
</tr>
<tr>
<td>Grade 4 (erosion)</td>
<td>4</td>
<td>8</td>
<td>12</td>
<td>16</td>
</tr>
<tr>
<td>Grade 5 (denudation)</td>
<td>5</td>
<td>10</td>
<td>15</td>
<td>20</td>
</tr>
<tr>
<td>Grade 6 (deformation)</td>
<td>6</td>
<td>12</td>
<td>18</td>
<td>24</td>
</tr>
</tbody>
</table>

Score = grade × stage

Fig. 1. Experimental schedule. Four weeks after administration of MIA, Group A was sacrificed, and Groups B and C were subjected to cell sheet transplantation surgery. At 8 weeks after MIA administration, Group B and C were sacrificed.

Fig. 2. Macroscopic and histological findings, and gene expression in cell sheets. (A) Representative images of a cell sheet attached to a white polyvinylidene membrane. (B) Hematoxylin–eosin, Safranin O, toluidine blue, and collagen type II staining of the cartilage matrix. Collagen type I did not stain (>40, scale bar = 50 μm). (C) The expression of Col1A1 and ITGa10 increased, but there was no significant difference (p > 0.05). Expression of Sox9 and Mmp13 increased significantly but not more than 2-fold (*p < 0.05, >2-fold). Expression of Col2A1, Acan, Lect1, and Comp increased significantly by >2-fold ("p < 0.05, >2-fold).
injection, the ratio did not differ significantly between any groups. Fig. 3B shows the limb weight distribution ratio on the last day was 47.4% ± 1.0% in Group A, 49.4% ± 0.2% in Group B, and 48.4% ± 0.6% in Group C; the value was significantly higher in Group B than in the other two groups.

3.4. Macroscopic evaluation

No signs of infection such as swelling or redness of the knee joint were observed macroscopically throughout the entire study. Fig. 3C shows the macroscopic findings for femoral condylar and tibial articular cartilage in each group compared to normal. Normal femoral condylar articular cartilage did not stain with indigo carmine and its surface was smooth. In Group A, damage was observed in the medial condyle. Minor injury was observed in Group B, and severe injury was observed in the medial and lateral condyles in Group C. Tibial articular cartilage damage was observed in all groups as for the femur, but there was less evidence of erosion in Group B than in Group C.

3.5. Histological evaluation

Fig. 4A shows the histological findings for femoral condylar articular cartilage and tibial articular cartilage. Safranin O staining of the femur and tibia showed loss of chondrocytes and decreased proteoglycan staining in Group A, but the superficial cartilage layer was stable. Similar findings were observed in Group C, but thinning of the articular cartilage was also observed. By contrast, in Group B, the superficial cartilage layer and partial staining were retained in the femur, and only mild damage was found in the surface cartilage layer of the tibia. Immunostaining showed collagen type II staining in all intact cartilage, but collagen type I staining was observed only in the superficial regions of cartilage in normal, Group A, and Group B.

3.6. Histological scoring

Fig. 4B shows the OARSI scores for the femur and tibia in the three groups. The score for the femur was significantly lower (indicating less damage) in Group B (5.83 ± 3.02 points) than in Group A (8.28 ± 4.87 points) and Group C (11.5 ± 4.5 points). For the tibia sites, the scores did not differ significantly between groups: 9.89 ± 5.09 points in Group A, 11.33 ± 4.57 points in Group B, and 13.83 ± 2.26 in Group C.

4. Discussion

We have previously investigated defects created in articular cartilage of various animals such as rats [26], rabbits [27], and mini pigs [28], and we have transplanted chondrocyte sheets to confirm the efficiency of articular cartilage repair. The purpose of this study was to investigate the application of allogeneic chondrocyte sheets in a rat model of nontraumatic early arthritis.

The prepared chondrocyte sheets had a survival rate of 80% or more, sufficient thickness, abundant extracellular matrix, and significant increase in the expression (>2-fold) of Col2A1, Acan, Lect1, etc.
and Comp. Collagen type II and aggrecan are the major components of cartilage matrix and are essential for the maintenance of normal chondrocytes [35]. Collagen type II is resistant to tensional force [36]. Aggrecan is also involved in providing resistance to joint loading by forming a complex through interactions with proteins such as hyaluronic acid [37]. Lect1 encodes a chondromodulin I protein, and Miura et al. [38] reported that it has an antiangiogenic effect in cartilage tissue and promotes cartilage formation. The STAT signaling pathway regulates transcription of multiple genes that regulate cell proliferation and differentiation, including osteoblast differentiation, and bone-related factors such as alkaline phosphatase and collagen type X [39,40]. Chondromodulin I inhibits the STAT signaling pathway [41]. Klinger et al. [42] reported that chondromodulin I strongly suppresses chondrocyte hypertrophy and endochondral ossification and helps to stabilize cartilage repair tissue.

Cartilage oligomeric matrix protein (COMP) is a noncollagenous protein that is important for multicellular networks and is known to stimulate chondrocyte proliferation and cartilage formation [43,44]. Some have reported interactions between COMP and collagen types I and II, and that these interactions play a role in maintaining the stability of the collagen network [45,46]. Although COMP is found in various tissues, it is most abundant in tissues that bear mechanical loading such as tendons and cartilage [47]. Our data suggest that chondrocyte sheets exhibit sufficient expression of genes important to maintaining a stable cartilage matrix.

Using the limb load distribution ratio as an index of pain, we observed a significant pain-relieving effect in the rats that received transplanted chondrocyte sheets. Our results are similar to those reported for transplantation of allogeneic chondrocyte sheets in a rabbit articular cartilage defect model at 4 weeks after transplantation reported by Ito et al. [27] and 12 weeks after transplantation reported by Tani et al. [30].

Macroscopic observations showed no inflammatory findings in the knee such as swelling or reddening. In addition, the transplanted group (Group B) showed evidence of suppression of cartilage degeneration compared with the nontransplanted groups (Groups A and C), which did not receive transplanted sheets.

Histologically, the adhesion of the margin and bottom of the cell sheet was good, and tissue was significantly repaired on the femur sites. On the tibia sites, there was no significant difference between groups that received or did not receive chondrocyte sheet transplantation. However, there was a tendency toward less progression of cartilage degeneration in Group B. In this study, the cell sheet was transplanted into the femoral condylar aspect, and significant cartilage repair was obtained on the femur side. These findings suggest that degeneration of the tibia sites may have been suppressed by a paracrine effect.

Udo et al. [24] also used an MIA arthritis model, but our macroscopic findings and histological scoring of Groups A and C were more favorable in our study. As noted earlier, the effects of MIA depend on the timing and dose. Although there are no reports of the residual time in joints, Udo et al. reported the progress of cartilage degeneration for 12 weeks after low-dose MIA injection. Future studies should extend the observation period and investigate the effectiveness of chondrocyte sheet transplantation. We did not measure inflammatory factors in the synovium or infrapatellar fat pad (IFP) in this study. Udo et al. observed decreased inflammation in the IFP and reversible changes 7 days after low-dose MIA injection. Synovitis and cartilage degeneration are typical features of knee joint OA, and it is necessary to confirm the effectiveness of chondrocyte sheet transplantation against synovitis.
5. Conclusion
In this study, we transplanted allogeneic chondrocyte sheets into rats with OA induced by low-dose MIA injection. Transplantation of cell sheets alleviated pain, as shown by a higher limb weight ratio in the group that received transplanted sheets. Histological findings showed less degeneration in rats that received transplanted sheets. These data suggest that, in this nontraumatic early arthritis model, transplantation of allogeneic chondrocyte sheets reduced the effects of OA, induced cartilage repair, and inhibited the progression of articular cartilage degeneration.

Declarations of interest
M.S. is one of the inventors on the patent (WO2006093151) submitted by CellSeed Inc. for the manufacturing process of chondrocyte sheets. M.S. receives research funds from CellSeed Inc.

Acknowledgements
This research was supported by the Project Focused on Developing Key Evaluation Technology: Manufacturing Technology for Industrialization in the Field of Regenerative Medicine (No. 14525207) and by the Project Focused on Developing Key Evaluation Technology: Evaluation for Industrialization in the Field of Regenerative Medicine (No. 15667006 to M.S.) from the Japan Agency for Medical Research and Development.

We are grateful to the Support Center for Medical Research and Education, Tokai University.

References


