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Reduced nucleotomy-induced intervertebral disc disruption through spontaneous spheroid formation by the Low Adhesive Scaffold Collagen (LASCol)

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A B S T R A C T

Back pain is a global health problem with a high morbidity and socioeconomic burden. Intervertebral disc herniation and degeneration are its primary cause, further associated with neurological radiculopathy, myelopathy, and paralysis. The current surgical treatment is principally discectomy, resulting in the loss of spinal movement and shock absorption. Therefore, the development of disc regenerative therapies is essential. Here we show reduced disc damage by a new collagen type I-based scaffold through actinidain hydrolysis—Low Adhesive Scaffold Collagen (LASCol)—with a high 3D spheroid-forming capability, water-solubility, and biodegradability and low antigenicity. In human disc nucleus pulposus and annulus fibrosus cells surgically obtained, time-dependent spheroid formation with increased expression of phenotypic markers and matrix components was observed on LASCol but not atelocollagen (AC). In a rat tail nucleotomy model, LASCol-injected and AC-injected discs presented relatively similar radiographic and MRI damage control; however, LASCol, distinct from AC, decelerated histological disc disruption, showing collagen type I-comprising LASCol degradation, aggrecan-positive and collagen type II-positive endogenous cell migration, and M1-polarized and also M2-polarized macrophage infiltration. Reduced nucleotomy-induced disc disruption through spontaneous spheroid formation by LASCol warrants further investigations of whether it may be an effective treatment without stem cells and/or growth factors for intervertebral disc disease.

1. Introduction

Back pain increases with age, affects 70–85% of people during their lives, and causes disability [1] with estimated healthcare costs up to $100 billion/year in the US [2]. The cause of back pain is multifactorial; however, intervertebral disc degeneration is one of the independent risk factors [3]. Intervertebral disc herniation, an injury with degenerative wear and tear, occurs even in the youth, limits daily and sports activities, and requires surgery in 10% [4].

The intervertebral disc has a complex structure with the nucleus pulposus (NP) encapsulated by the annulus fibrosus (AF) and endplates [5]. The collagenous, laminar AF surrounds the central, gelatinous NP, maintaining pressurization of the NP for support of compressive loading and facilitation of multidimensional spinal movement [5]. While the AF comes from the mesenchyme [6], the origin of the NP is the notochord [7]. Disc cells, positive for brachyury, CD24 (both disc NP markers) [8],...
tyrosine kinase with Ig and EGF homology domains 2 (Tie2) (disc NP progenitor marker) [9], collagen type V alpha 1 [10], and CD146 (both disc AF markers) [11], have a chondrocytic phenotype to produce matrix components comprising proteoglycans (principally aggrecan) and collagens (primarily types I in the AF and II in the NP) [12]. Furthermore, the disc is an immune-privileged, the largest avascular organ in the body [13]. Disc cells thus live under an extremely harsh environment—low glucose, oxygen, and pH and high osmolality and load fluctuation [14]. While age-related disc changes characterized by the loss of notochordal cells appear from early childhood as observed in people aged 11–16 years [15], ~40% of people aged under 30 years and 90% of those aged over 55 years present structural lumbar disc degeneration [16], which can cause back pain [17].

Herniated and degenerated discs can present not only back pain but also radicular pain, numbness, muscle weakness, and then paralysis in the worst scenario [18]. Despite successful conservative treatment outcomes with medication and physiotherapy in disc disease, non-responders reluctantly require surgery [4]. Surgical interventions predominantly include the excision of damaged discs, resulting in the function loss, immobilization, and potential additional complications due to the altered biomechanics [19]. Therefore, the development of new therapies for direct disc repair is an urgent issue.

Stem cells [20], growth factors [21], and tissue engineering [22] have been applied for disc regeneration. Each trial is effective; nevertheless, unclarified aspects still remain for the human body application. Mesenchymal stem cells are at risk of osteophyte [23] and also neoplasms [24] formation with an extremely low survival rate of intradiscally transplanted cells [20]. The bone morphogenetic protein-2 is associated with carcinogenesis [25]. Hence, interest has recently expanded to bioengineering approaches that exploit endogenous cell populations to restore the disc structure and function [22]. Injectable scaffolds using atelocollagen (AC), alginate, hyaluronan, and chitosan have been tested [26]. As a collagen-based scaffold, AC has often been used clinically, such as autologous chondrocyte-seeded AC for cartilage repair [27] and porous hydroxyapatite/AC composites for bone defect [28]. While several kinds of scaffold have shown the regenerative potential, no scaffolds are clinically available for disc treatment because of the undiscovered repair mechanisms as well as safety concerns [22].

In the tissue-engineering field, spheroïd formation has increased attention because of its 3D cell-culturing structure capable of mimicking the in vivo environment [29]. Morphological characteristics, metabolic activity, and function of cells are better maintained in 3D than 2D [29]. Spheroïd formation is effective in maintaining the phenotype of progenitor cells in the disc [9]. We thus developed a new collagen-based scaffold through the hydrolysis with actinidain protease, which was named as the Low Adhesive ScaFFold Collagen (LASCol) from the characteristic of its high 3D spheroïd-forming capability [30–32]. The LASCol is designed by removing major parts of the N-terminal and C-terminal telopeptides of collagen type I, while AC is similarly observed in all, 3 of which were dead in the control group, this test indicates a safe, relatively low antigenicity of LASCol. A low-immunogenic potential of LASCol makes it possible to apply directly without any immunosuppression even in xenogeneic transplantation.

In vitro, 7.0-mg/ml LASCol gel, consisting of Dulbecco’s modified Eagle’s culture solution, Reconstitution buffer, and 10.0-mg/ml LASCol solution from 5.0-mM hydrochloric acid and freeze-dried LASCol, was distributed at 1 ml in a 35-mm dish or each well of a 6-well plate and at 200 μm in each well of a 24-well plate at 37 °C overnight. Similarly, 2.1-mg/ml AC gel, consisting of 3.0-mg/ml AC solution, was distributed at the same amounts.

In vivo, 42.0-mg/ml, 21.0-mg/ml, 14.0-mg/ml, and 7.0-mg/ml LASCol and 7.0-mg/ml AC gels were prepared from saline, Reconstitution buffer, and LASCol or AC solution. Based on the difference in solubility between LASCol and AC, 21.0-mg/ml LASCol and 7.0-mg/ml AC gels were compared. Effects of LASCol were also tested at other concentrations. Both gels had an injectable, low viscosity on ice and became stabilized at the body temperature of 37 °C within 1 min after the intradiscal injection, the storage modulus of which at 37 °C after 30 min was approximately ≥1.4 kPa, within a reported range in the viscosity of in-vivo disc NP tissues [33]. The LASCol gel becomes cloudy compared to the AC gel, which resulted from the difference in fibroïd (Fig. 1B).
A

N-terminal
telopeptide
Collagen type I
Low Adhesive
Scaffold Collagen
(LASColl)
C-terminal
telopeptide
Atelocollagen
(AC)

B

Injectable gels

0 min 1 min
LASColl
AC

C

Western blotting Human disc tissues

| Sample       | NP #1 | NP #2 | NP #3 | AF #4 |
|--------------|-------|-------|-------|-------|
| 49 kDa       |       |       |       | Brachyury |
| 37 kDa       |       |       |       | CD24   |
| 184 kDa      |       |       |       | Col5a1 |
| 118 kDa      |       |       |       | CD146  |
| 42 kDa       |       |       |       | Actin  |

D

Surgical procedures

Skin marking & incision Disc exposure AF incision & NP aspiration Gel injection Skin closure

E

Rat tail disc mid-sagittal sections

LASColl (+methylene blue) AC (+methylene blue) Control Intact

Vertebral body AF NP Disc

1 mm

(caption on next page)
Pfirrmann degeneration grade [34], median 2 ± 0.6 (2–4) (n = 12 for AF; age, 61.8 ± 11.6 [41–80] years; sex, 9 males and 3 females; Pfirrmann degeneration grade, median 3 ± 0.5 (2–4)). We carefully obtained disc AF tissues by box-cutting AF sharply with a scalpel and NP tissues from the first bite of tissues by a laminectomy rongeur without any violation of vertebral endplates, both of which were collected from discarded surgical waste. Immediately after surgery, disc NF and AF tissues were digested for cell isolation in Dulbecco’s modified Eagle’s medium (DMEM) with 10% fetal bovine serum (FBS), 1% penicillin/streptomycin, and 0.114% collagenase type 2 at 37 °C for 1 h in the NP and 12 h in the AF. Isolated cells were grown in 1% penicillin/streptomycin-supplemented DMEM with 10% FBS. To retain the phenotype, only first-passage cells were used for cell-counting (n = 6), immunofluorescent (n = 6), and real-time reverse transcription–polymerase chain reaction (RT–PCR) (n = 3) experiments following 72-h pre-culture. Cell samples collected abundantly were used across experiments. Experimental conditions were maintained throughout at 37 °C under 5% CO2, 2% O2 to stimulate the physiologically hypoxic disc environment [13,14].

The acquisition of human disc specimens specific for the NP and AF was validated by Western blotting using directly extracted proteins from residual tissues as described previously [35,36]. Briefly, tissues were homogenized using the MS-100R bead-beating disrupter for 30 s twice at 4 °C in the T-PER tissue protein extraction reagent with protease and phosphatase inhibitors. Soluble proteins were collected after centrifugation at 20,000 g for 15 min at 4 °C. Protein concentration was determined by the bichinchoninic acid assay. Equal 30-mg amounts of protein were mixed with the electrophoresis sample buffer and boiled for 5 min before loading onto a 7.5–15.0% polyacrylamide gel. Separated tissue proteins in the Tris–glycine–sodium dodecyl sulfate buffer system were transblotted electrically and probed with primary antibodies (1:200–1:1000 dilution) for brachyury [8] and CD24 [8] (disc AF markers), and collagen type I [12,30,32], for brachyury, collagen type II (LASCol constituent and disc AF matrix component) [12,30,32], and collagen type I (LASCol constituent) [12,30,32], and collagen type II (disc NP matrix component) [12] overnight at 4 °C for antigen-retrieval, permeabilized, and blocked. Multi-color immunofluorescence was performed at 1:400-diluted Alexa Fluor® 488, 568, and 647 secondary antibodies for 1 h at room temperature. Similarly, collagen type I alpha 1 [10], CD146 [11] (disc NF markers), and collagen type V alpha 1 [10] and CD24 [8] (disc NP markers), collagen type IV alpha 1 [110] and CD146 [11] (disc NF markers), and actin (loading control) for 12 h in 4 °C followed by secondary antibodies (1:2000 dilution). Signals were visualized by enhanced chemiluminescence. Images were obtained using the Chemilumino analyzer LAS-3000 mini. Consequently, an acceptable disc NP-specific and AF-specific marker expression was confirmed (Fig. 1C).

2.5 Animals

Twelve-week-old male Sprague–Dawley rats were used. Under general anesthesia, 1-cm longitudinal incision was made along the tail to expose the lateral portion of the caudal (C) disc. A #11 scalpel blade was inserted 1.5 mm into the disc. Nucleotomy by disc AF incision and NP aspiration with a 22-gauge catheter on a 5-ml syringe was conducted as described previously [37,38] (Fig. 1D).

Nucleotomy at C8–C9, C9–C10, and C10–C11 was performed to assess effects of materials in 44 rats (weight 427.1 ± 15.2 [405–450] g). Intrasidial 15-μl injection of 21-mg/ml LASCol gel, 7.0-mg/ml AC gel, and solvent control (saline and Reconstitution buffer) was conducted using a 25-gauge catheter, respectively. These three treatments were all applied in each rat tail to eliminate the individual difference and randomly at each disc segment. The injection volume was determined based on prior evidence [39]. Intradiscal gel injection without extrusion was preliminarily established by using a methylene blue dye (Fig. 1E). Radiographs were taken before and 7–56 d after surgery (n = 8). Then, magnetic resonance images (MRIs) were taken 28 d after surgery (n = 6). After imaging, vertebral body–disc–vertebral body functional spinal units were frozen with Super Cryoembedding medium quickly in liquid nitrogen. Mid-sagittal sections at 5-μm thickness were prepared for histology as described previously [40].

Nucleotomy at C8–C9, C9–C10, C10–C11, and C11–C12 was performed to assess dose-dependent effects of LASCol in 24 rats (weight 436.9 ± 17.8 [415–463] g). Intradiscal 15-μl injection of 42.0-mg/ml, 21.0-mg/ml, 14.0-mg/ml, and 7.0-mg/ml LASCol gels was conducted, respectively. These four concentrations were all applied in each rat tail to exclude the individual difference and randomly at each disc segment.

Radiographs were taken before and 7–56 d after surgery (n = 6).

2.6 Immunofluorescence

In vitro, in a 35-mm dish coated with 1-ml LASCol or AC gel, 3.0 × 105 human disc NP and AF cells/well were cultured in 2-ml 10% FBS-supplemented DMEM for 120 h. Fresh media of 500 μl were gently added at 72 h without exchange to prevent cell detachment. Cells were fixed in 4% paraformaldehyde for 30 min. To assess the phenotype, disc NP cells were stained with 1:100-diluted primary antibodies for brachyury (disc NP marker) [8], Tie2 (disc NP progenitor marker) [9], and aggrecan (disc matrix component) [12] or for brachyury, collagen type I (LASCol constituent) [12,30,32], and collagen type II (disc NP matrix component) [12] overnight at 4 °C, followed by 1:400-diluted Alexa Fluor® 488, 568, and 647 secondary antibodies for 1 h at room temperature. Similarly, collagen type I alpha 1 [10], CD146 [11] (both disc AF markers), and collagen type I (LASCol constituent and disc AF matrix component) or aggrecan primary antibodies and Alexa Fluor® 488, 568, and 647 secondary antibodies were used for disc AF cell staining. The 4’,6-diamidino-2-phenylindole (DAPI) was used for counterstaining. Images were photographed by the BX-Z700 microscope. The number of positive cells was counted in five random low-power fields (× 100) using the ImageJ software (https://imagej.nih.gov/ij/). Briefly, we created binary images at a fixed intensity level and counted positive cells included in the field. The positive cell percentage for brachyury, Tie2, aggrecan, and collagen types I and II in the NP and for collagen type V alpha 1, CD146, collagen type I, and aggrecan in the AF was calculated as relative to the total number of DAPI-positive cells.

In vivo, rat caudal disc sections were fixed with 4% paraformaldehyde for 30 min, decalcified in 10% ethylenediaminetetraacetic acid for 30 min, antigen-retrieval, permeabilized, and blocked. Multi-color immunofluorescence was performed with 1:200-diluted primary antibodies overnight at 4 °C for collagen types I and II to assess the composition of matrix [12] and scaffolds [30,32], for brachyury [8], Tie2...
[9], and aggrecan [12] to assess the phenotype of cells, or for ionized calcium binding adaptor molecule 1 (Iba1, pan-macrophage marker) [41], CD86 (M1-polarized macrophage marker, characterized by producing pro-inflammatory mediators) [42], and CD163 (M2-polarized macrophage marker, characterized by responses to anti-inflammatory and tissue remodeling) [43] to assess the infiltration of macrophages, followed by 1:400-diluted Alexa Fluor® 488, 568, and 647 secondary antibodies for 1 h at room temperature. The DAPI was used for counterstaining. Similarly, imaging and positive cell-counting were performed by creating binary images at a fixed level with ImageJ, and the percentage of positive cells for brachyury, Tie2, aggrecan, Iba1, CD86, and CD163 in the NP was calculated.

2.7. Time-lapse cell-counting and spheroid-counting

In vitro, in a 24-well plate coated with 200-μl LASCol or AC gel, 2.0 × 10^4 human disc NP and AF cells/well were cultured in 500-μl 10% FBS-supplemented DMEM. Fresh media of 100 μl were gently added every 72 h without exchange. Time-lapse photographing was performed every 6 h for 192 h. The number of cells and spheroids was counted in five random low-power fields (×100) using ImageJ. In this study, spheroid was defined as the aggregation of ≥ 3 cells.

2.8. RNA isolation and real-time RT-PCR

In vitro, in a 6-well plate coated with 1-ml LASCol or AC gel compared to the non-coated condition, 2.0 × 10^4 human disc NP cells/well were cultured in 2-ml 10% FBS-supplemented DMEM. Fresh media of 500 μl were gently added every 72 h without exchange. At 72 and 168 h, total RNA was extracted using the RNeasy mini kit, and 0.1-μg RNA was reverse-transcribed with random primers. Messenger RNA (mRNA) expression levels of chondrogenic SOX9 encoding SRY-box 9 (chondrogenesis regulator) [44], COMP encoding cartilage oligomeric matrix protein (cartilage turnover marker) [45], and TGFB1 encoding transforming growth factor beta 1 (chondrogenesis inducer) [46], relative to ATP synthase, H+ transporting, mitochondrial F0 complex subunit 1 (ATP5F1) [47] as an endogenous control were assessed in duplicate by real-time RT-PCR using a SYBR Green fluorescent dye. These target genes were chosen to analyze the trend of cartilage differentiation. To identify appropriate housekeeping genes for references, mRNA expression levels of 15 genes were evaluated using a human housekeeping gene primer set, and ATP5F1 was selected as a reference based on the highest stability in mRNA expression regardless of complementary DNA concentration. The commercialized, validated primer sequences were used as follows: SOX9, forward 5′-GTGAGATGAAATCTGTCTGGGAAATG-3′, reverse 5′-TGTGAGATGAAATCTGTCTGGGAAATG-3′; COMP, forward 5′-AGGTTACCACTCAAGCCAGAAG-3′, reverse 5′-GTTGACCACTCGTGAGGCAAG-3′; TGFB1, forward 5′-TCTCAGGGCATACCTAGCAG-3′, reverse 5′-GCTACGAGGAAGGCTCCTACA-3′; ATP5F1, forward 5′-GAAGAGGCGCTTCCATCAACCA-3′, reverse 5′-GTCTGGCGATTAAGGTTACCACTCAAGCCAGAAG-3′. Melting curve analysis was performed using the Dissociation Curve software to ensure that only a single product was amplified. Relative mRNA expression was analyzed using the 2−ΔΔCt method [48]. The value of non-coated control sample was set as 1.

2.9. Radiography

In vitro, lateral radiographs were taken using VPX-30E system and IXFR film (exposure time 40 s; distance 40 cm; current 3 mA; voltage 35 kV). Radiographs were measured twice at one-week intervals by each of two investigators blinded to the study purpose. Disc height was measured using ImageJ, normalized to adjacent vertebral body heights as the disc height index (DHI), shown as the percent of preoperative DHI (%DHI = [postoperative DHI/preoperative DHI] × 100) as described previously [49], and further normalized to the intact disc as the normalized %DHI (normalized %DHI = [experimental %DHI/intact %DHI] × 100) [50] (Fig. 4A).

2.10. MRI

In vivo, MRIs were taken using 4.7-T Varian Unity Inova 200 MHz MRI unit. Sagittal T2-weighted imaging and T2 mapping (repetition time/echo time 2000/30 and 60 ms [2 echoes], field of view 100 × 50 mm, slice thickness 2 mm, matrix size 512 × 512 mm, number of experiments 4, acquisition time 35 min) were performed. Region of interest was positioned in the disc center using the MATLAB software 9.1.0.441655 (R2016b), and mean T2 values were computed within the region as described previously [51].

2.11. Safranin-O staining

In vivo, safranin-O, fast green, and hematoxylin staining was performed to demonstrate proteoglycan distribution. Disc NP-cell number and safranin-O-positive area were measured using ImageJ. Briefly, we created binary images at a fixed intensity level and measured the area between vertebral endplates. We counted cells included in the area.

2.12. Statistical analysis

Data are expressed as the mean ± standard deviation in the text and box plots in the graphs. The Student t-test was used to assess effects of LASCol and AC treatments in in-vitro immunofluorescence. One-way analysis of variance (ANOVA) with the Tukey–Kramer post-hoc test was used to assess effects of multiple treatments in in-vivo MRI. Two-way repeated measures ANOVA with the Tukey–Kramer post-hoc test was used to assess effects of treatment and time in in-vitro cell-counting and spheroid-counting and in-vivo radiography, histomorphology, and immunofluorescence (experiments using replicates from the same donors) as well as in in-vitro real-time RT–PCR (target gene expression analysis shown as relative values of the control). In addition, intra-class correlation coefficient was calculated to determine intra-observer and inter-observer reliabilities for the measurement of radiographic parameters. The P-values of < 0.05 were regarded as statistically significant using IBM SPSS Statistics 23.0 (IBM, Armonk, NY).

3. Results

3.1. Spontaneous spheroid formation of human disc NP and AF cells on LASCol

First, we observed spheroid formation of human disc NP and AF cells on the LASCol gel in vitro. Both cell types proliferated in monolayer on the AC gel but demonstrated spheroid formation on the LASCol gel (Fig. 2A). Time-lapse photographing presented cell-aggregating spheroid formation on the LASCol gel only, the pattern of which was similar between disc NP and AF cells (Fig. 2B). Cell-counting analysis found no apparent increases in disc NP and AF cells on the LASCol but not AC gel (P < 0.001) (Fig. 2C). However, the number of spheroids, defined as ≥ 3 cell aggregation, progressively increased in both cells on the LASCol gel only (P < 0.001) (Fig. 2D). Spontaneous spheroid formation of disc NP and AF cells on the LASCol gel indicates a high compatibility of LASCol with both cell types.

3.2. Enhanced disc-cell and chondrocyte phenotype of human disc NP and AF cells on LASCol

Next, we assessed effectiveness of spheroid formation in human disc NP and AF cells on the LASCol gel in vitro. In disc NP cells, multi-color immunofluorescence showed elevated expression of brachyury (disc NP marker) [8], Tie2 (disc NP progenitor marker) [9], and aggrecan (disc...
Fig. 2. Spontaneous spheroid formation of human disc NP and AF cells on LASCol. (A) Human disc NP and AF cells cultured on the LASCol or AC gel in 10% FBS-supplemented DMEM at 120 h. Black triangles indicate spheroids. (B) Time-lapse human disc AF cells cultured on the LASCol gel in 10% FBS-supplemented DMEM at 72–90 h. (C) Changes in the number of human disc NP and AF cells cultured on the LASCol or AC gel. (D) Changes in the number of human disc NP and AF spheroids cultured on the LASCol or AC gel. The spheroid was defined as the aggregation of ≥3 cells. In (C) and (D), the number of cells and spheroids was counted in five random low-power fields (×100). Data are presented with box plots (n = 6). Two-way repeated measures ANOVA with the Tukey–Kramer post-hoc test was used.
In spheroids on the LASCol gel compared to on the AC gel (brachyury, \( P < 0.001 \); Tie2, \( P < 0.001 \); aggrecan, \( P = 0.001 \)). In addition, expression of collagen type II (disc NP matrix component) [12] in spheroids on the LASCol gel compared to on the AC gel, while collagen type I (LASCol constituent) [12,30,32] was comparable between on the LASCol and AC gels (collagen type II, \( P = 0.004 \); collagen type I, \( P = 0.39 \) (Fig. 3AC)). In disc AF cells, collagen type VI alpha 1 [10] and CD146 (both disc AF markers) [11] were more abundant in spheroids on the LASCol gel than on the AC gel (collagen type VI alpha 1, \( P = 0.02 \); CD146, \( P < 0.001 \)). Immunoactivity for aggrecan appeared to be higher on LASCol than on AC, showing the tendency toward increase on the LASCol gel although it did not reach statistical significance (\( P = 0.06 \) (Fig. 3BC)). Meanwhile, the percentage of immunopositive cells for collagen type I (disc AF matrix component) [12] was similarly high between on LASCol and AC, despite marked collagen type I expression in spheroids on the LASCol gel than on the AC gel (\( P = 0.20 \) (Fig. 3BC)). A high level of background signals detected in all measurements for collagen type I in disc NP and AC cells on LASCol and AC gels would indicate the presence of collagen type I-comprising LASCol and AC gels. Spontaneous spheroid formation on the LASCol gel is efficient in increasing disc NP-cell and AF-cell phenotypes compared to thin spreading of cells on the AC gel.

Then, we evaluated gene expression in human disc NP cells on the LASCol gel in vitro. Real-time RT–PCR displayed mRNA up-regulation of TGFB1 (chondrogenesis inducer) [46] on the LASCol gel relative to on the AC gel and non-coated control at 72 h (LASCol versus AC, \( P = 0.03 \); versus control, \( P = 0.009 \)). In addition, SOX9 (chondrogenesis regulator) [44], COMP (cartilage turnover marker) [45], and TGFB1 mRNA expression on the LASCol gel was up-regulated relative to on the AC gel and non-coated control at 168 d (SOX9: LASCol versus AC, \( P = 0.04 \); versus control, \( P = 0.02 \)) (COMP: LASCol versus AC, \( P = 0.02 \); versus control, \( P = 0.01 \); TGFB1: LASCol versus AC, \( P = 0.01 \); versus control, \( P = 0.02 \)). These gene expression on the LASCol gel all showed timedependent up-regulation between 72 and 168 h (SOX9, \( P = 0.001 \); COMP, \( P = 0.003 \); TGFB1, \( P = 0.002 \) (Fig. 3D). Spheroid-forming cultures on the LASCol gel stimulate chondrogenic growth of disc NP cells.

3.3. Delayed loss of radiographic disc height and MRI disc intensity by LASCol in a rat tail nucleotomy model

Based on a distinct spheroid-forming and phenotype-preserving characteristics of human disc NP and AF cells on LASCol in vitro, an in vivo study was designed using a rat tail nucleotomy model of disc damage which is well established because of the easy accessibility and high reproducibility [37,38]. First, radiographic analysis for the intradiscal 15-μl injection of 21.0-mg/ml LASCol gel, 7.0-mg/ml AC gel, and solvent control after nucleotomy was performed to assess effects of materials. For radiographic disc height measurements using DHI [49] (Fig. 4A), the intra-observer reliability was 0.913–0.947 by intraclass correlation coefficient while the inter-observer reliability was 0.934, all values of which indicated an acceptable reproducibility. Time-course radiographs demonstrated progressive disc space narrowing in solvent control discs (normalized %DHI at 7 d, 65.8 ± 8.0%; 56 d, 50.2 ± 11.9%) but relatively maintained disc height in LASCol-injected discs (normalized %DHI at 7 d, 78.6 ± 6.7%; 56 d, 77.4 ± 11.4%) and AC-injected discs (normalized %DHI at 7 d, 73.9 ± 9.8%; 56 d, 71.8 ± 11.5%) (Fig. 4BC). These normalized %DHI values were higher in LASCol-injected than control discs at 7–56 d (7 d, \( P = 0.005 \); 14 d, \( P = 0.003 \); 28 d, \( P = 0.002 \); 56 d, \( P < 0.001 \)). Meanwhile, statistical difference in normalized %DHI between AC-injected and control discs was detected only at 56 d (\( P = 0.001 \) (Fig. 4C)). Intradiscal injection of LASCol or AC expedites the maintenance of rat tail disc height.

Next, radiographic analysis for the intradiscal 15-μl injection of LASCol gels at varying 7.0 mg/ml concentrations after nucleotomy was performed to assess dose-dependent effects of LASCol. Time-course radiographs identified dose-dependent maintenance of height in LASCol-injected discs (normalized %DHI at 56 d in 42.0 mg/ml, 74.4 ± 8.2%; 21.0 mg/ml, 71.9 ± 7.8%; 14.0 mg/ml, 66.0 ± 5.2%; 7.0 mg/ml, 55.7 ± 11.2%) (Fig. 4DE). In fact, normalized %DHI values were higher in 42.0-mg/ml than in 7.0-mg/ml LASCol-injected discs at 7–56 d (7 d, \( P = 0.02 \); 14 d, \( P < 0.001 \); 28 d, \( P = 0.004 \); 56 d, \( P = 0.008 \)). Similarly, statistical difference in normalized %DHI between 21.0-mg/ml and 7.0-mg/ml LASCol-injected discs was observed at 14–56 d (14 d, \( P = 0.01 \); 28 d, \( P = 0.003 \); 56 d, \( P = 0.02 \) (Fig. 4E). Based on these findings, 21.0 mg/ml is proposed as a minimum requirement of intradiscal LASCol concentration.

To find clinically relevant differences between intrastradal LASCol and AC injection, MRIs were taken at 28 d. On T2-weighted images, LASCol-injected discs had higher intensity compared to solvent control discs, indicating preserved disc hydration (Fig. 4F). On T2-mapping images, LASCol-injected discs contained a larger yellow and green area than control discs, indicating a higher water content (Fig. 4F). Then, T2-mapping quantification found statistical T2-value difference between LASCol-injected and control discs (\( P = 0.03 \), although T2 values in all experimental discs were lower than intact discs (intact [33.7 ± 3.6 ms] versus LASCol [25.7 ± 4.0 ms], \( P = 0.001 \); versus AC [21.5 ± 1.9 ms], \( P < 0.001 \); versus control [20.5 ± 1.6 ms], \( P < 0.001 \) (Fig. 4G)). No osteophyte, neoplasia, or abnormal granulation was found in LASCol-injected discs. Radiographic and MRI findings indicate a reduced nucleotomy-induced disc disruption by LASCol, which is equal or better than AC.

3.4. Delayed loss of cell number and safranin-O-positive matrix by LASCol in the disc NP of a rat tail nucleotomy model

We then assessed rat tail LASCol-injected, AC-injected, and solvent control disc histomorphology by safranin-O, fast green, and hematoxylin staining to demonstrate proteoglycan distribution. In LASCol-injected discs, the presence of the LASCol gel was implicated as a safranin-O-negative NP region at 0 and 3 d. In particular, cell migration into the LASCol gel at 3 d was noteworthy (Fig. 5AB). At 7–56 d, safranin-O-positive matrix with cell migration was observed in LASCol-injected discs (Fig. 5AB). In AC-injected discs, the presence of the AC gel was similarly speculated at 0–7 d. However, only a few cells were found on the AC-gel surface at 3 and 7 d (Fig. 5AB). At 14–56 d, AC-injected discs also presented safranin-O-positive matrix; however, the area and cell number were substantially smaller than in LASCol-injected discs (Fig. 5AB). Meanwhile, control discs showed a gross loss, increased clefts, and progressive collapse of NP matrix throughout the study period (Fig. 5AB). Semi-quantification analysis found a reduced loss of cell number in the NP region of LASCol-injected discs compared to AC-injected and control discs at 7–56 d (cell number at 56 d in LASCol, 70.0 ± 15.5; AC, 26.2 ± 9.4; control, 3.2 ± 2.6) (all \( P < 0.001 \) (Fig. 5C). Delayed loss of safranin-O-positive disc NP area was also observed in LASCol-injected discs relative to AC-injected and control discs at 7–56 d (disc NP area at 56 d in LASCol, 251.6 ± 67.7 × 103 \( \mu \)m2; AC, 72.2 ± 41.5 × 103 \( \mu \)m2; \( P = 0.03 \), although T2 values in all experimental discs were lower than intact discs (intact [33.7 ± 3.6 ms] versus LASCol [25.7 ± 4.0 ms], \( P = 0.001 \); versus AC [21.5 ± 1.9 ms], \( P < 0.001 \); versus control [20.5 ± 1.6 ms], \( P < 0.001 \) (Fig. 4G)). No osteophyte, neoplasia, or abnormal granulation was found in LASCol-injected discs.

3.5. Decreased collagen type I-based LASCol and increased collagen type II-positive matrix in the disc NP of a rat tail nucleotomy model

On the basis of histomorphological findings, we performed multicolor immunofluorescence for collagen types I and II to identify injected LASCol and AC gels, which were both developed from collagen type I. In LASCol-injected discs at 0 d, there was the collagen type I-positive, type II-negative structure, indicating the LASCol gel. The gel structure
Fig. 3. Enhanced disc-cell and chondrocyte phenotype of human disc NP and AF cells on LASCol. (A) Immunofluorescence of human disc NP cells cultured on the LASCol or AC gel in 10% FBS-supplemented DMEM at 120 h for brachyury (green), Tie2 (red), aggrecan (purple), DAPI (blue), and merged signals and for brachyury (green), collagen type I (purple), collagen type II (red), DAPI, and merged signals. (B) Immunofluorescence of human disc AF cells cultured on the LASCol or AC gel in 10% FBS-supplemented DMEM at 120 h for collagen type V alpha 1 (green), CD146 (red), collagen type I (purple), DAPI, and merged signals and for aggrecan (purple), DAPI, and merged signals. (C) Changes in the percentage of human disc NP cells positive for brachyury, Tie2, aggrecan, and collagen types I and II and AF cells positive for collagen type V alpha 1, CD146, collagen type I, and aggrecan on the LASCol or AC gel. Immunopositivity was counted in five random low-power fields (× 100) and calculated as relative to the total number of DAPI-positive cells. Data are presented with box plots (n = 6). The Student t-test was used. (D) Real-time RT-PCR for SOX9, COMP, and TGFβ1 in total RNA extracts from human disc NP cells cultured on the LASCol or AC gel in 10% FBS-supplemented DMEM at 72 and 168 h. The ATP5F1 was used as an internal control. Changes in SOX9/ATP5F1, COMP/ATP5F1, and TGFβ1/ATP5F1 mRNA expression relative to the non-coated control are shown. Data are presented with box plots (n = 3). Two-way repeated measures ANOVA with the Tukey-Kramer post-hoc test was used. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)
underwent progressive DAPI-positive cell migration at 3 d. Then, accelerated disappearance of the collagen type I-positive region and complementary appearance of collagen type II-positive disc NP matrix were observed at 7 d (Fig. 6). In AC-injected discs at 0 d, the collagen type I-positive, type II-negative structure relatively remained unchanged during the initial 7 d, the disappearance of which was decelerated compared to in LASCol-injected discs. There were also no obvious increases in collagen type II expression. Cell migration was found only on the gel-structure surface (Fig. 6). Meanwhile, in control discs, collagen types I and II expression was not clearly detected throughout possibly because of extensive nucleotomy (Fig. 6). Biochemical analysis further clarifies the future potential of collagen type I-positive LASCol for biological tissue repair with collagen type II-positive NP-like tissues.
Fig. 4. Delayed loss of radiographic disc height and MRI disc intensity by LASCol in a rat tail nucleotomy model. (A) The disc height index (DHI) was calculated by measuring, averaging, and normalizing the disc heights to the adjacent vertebral body heights at the anterior, middle, and posterior portions. (B) Lateral radiographs of rat tail 15-μL LASCol-injected (21.0 mg/ml), AC-injected (7.0 mg/ml), and solvent control discs taken before (pre-operation [pre-op]) and at 7, 14, 28, or 56 d after nucleotomy. Dark-gray, light-gray, white, and black triangles indicate LASCol-injected, AC-injected, solvent control, and untreated intact discs, respectively. (C) Changes in %DHI of LASCol-injected, AC-injected, and control discs. (D) Lateral radiographs of rat tail 15-μL LASCol-injected discs at varying concentrations (21.0, 21.0, 14.0, and 7.0 mg/ml) taken before (pre-operation [pre-op]) and at 7, 14, 28, or 56 d after nucleotomy. Dark-gray, intermediate-gray, light-gray, white, and black triangles indicate 42.0-mg/ml, 21.0-mg/ml, 14.0-mg/ml, and 7.0-mg/ml LASCol-injected, and untreated intact discs, respectively. (E) Changes in %DHI of LASCol-injected discs at varying concentrations. (F) Sagittal T2-weighted and T2-mapping MRIs of rat tail 15-μL LASCol-injected (21.0 mg/ml), AC-injected (7.0 mg/ml), and solvent control discs taken at 28 d after nucleotomy. Dark-gray, intermediate-gray, light-gray, and white triangles indicate LASCol-injected, AC-injected, solvent control, and untreated intact discs, respectively. Pink circles indicate the respective discs in T2-mapping images. (G) Changes in T2-mapping value of LASCol-injected, AC-injected, control, and intact discs. In (C), (E), and (G), data are presented with box plots (c, n = 8; e, n = 6; g, n = 6). Two-way repeated measures ANOVA with the Tukey–Kramer post-hoc test was used for (C) and (E). One-way ANOVA with the Tukey–Kramer post-hoc test was used for (G). (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

Fig. 5. Delayed loss of cell number and Safranin-O-positive matrix by LASCol in the disc NP of a rat tail nucleotomy model. (A) Safranin-O staining of rat tail 15-μL LASCol-injected (21.0 mg/ml), AC-injected (7.0 mg/ml), and solvent control discs sectioned at 0, 3, 7, 14, 28, or 56 d after nucleotomy in low-power fields (× 100). Black rectangles indicate the disc NP area shown as high-power field images in Fig. 5B. (B) Safranin-O staining of rat tail 15-μL LASCol-injected, AC-injected, and solvent control discs sectioned at 3 and 28 d after nucleotomy in high-power fields (× 400). (C) Changes in the number of cells in LASCol-injected, AC-injected, and control disc NP spaces. (D) Changes in safranin-O-positive area of LASCol-injected, AC-injected, and control disc NP spaces. In (C) and (D), data are presented with box plots (n = 6). Two-way repeated measures ANOVA with the Tukey–Kramer post-hoc test was used.
3.6. Migration of cells with the endogenous phenotype into LASCol in the disc NP of a rat tail nucleotomy model

Next, we performed multi-color immunofluorescence for brachyury, Tie2, and aggrecan to assess the phenotype of migrated cells into the LASCol gel. First, we thought these cells as remnant disc NP cells. In LASCol-injected discs, a substantial percentage of cells were brachyury-positive (3 d, 36.4 ± 9.9%; 7 d, 37.1 ± 4.8%; 14 d, 34.0 ± 8.3%; 28 d, 32.0 ± 4.5%; 56 d, 27.9 ± 5.2%), Tie2-positive (3 d, 39.7 ± 5.5%; 7 d, 22.7 ± 7.0%; 14 d, 20.3 ± 3.2%; 28 d, 22.2 ± 4.3%; 56 d, 17.7 ± 3.2%), and aggrecan-positive (3 d, 48.2 ± 6.2%; 7 d, 42.0 ± 5.4%; 14 d, 41.7 ± 8.6%; 28 d, 39.5 ± 4.6%; 56 d, 36.8 ± 3.2%) (Fig. 7AB). However, in AC-injected discs, immunopositivity for brachyury (3 d, 18.9 ± 3.7%; 7 d, 15.9 ± 3.1%; 14 d, 13.7 ± 3.7%; 28 d, 13.1 ± 2.3%; 56 d, 11.6 ± 1.6%), Tie2 (3 d, 15.0 ± 5.6%; 7 d, 10.1 ± 3.8%; 14 d, 11.9 ± 2.5%; 28 d, 7.5 ± 1.6%; 56 d, 6.1 ± 3.1%), and aggrecan (3 d, 29.9 ± 8.6%; 7 d, 31.3 ± 6.0%; 14 d, 29.6 ± 3.9%; 28 d, 28.3 ± 4.8%; 56 d, 26.0 ± 5.6%) was statistically lower than in LASCol-injected discs (all P < 0.01) (Fig. 7AB). Similarly, in control discs, immunopositivity for brachyury (3 d, 8.9 ± 2.4%; 7 d, 13.8 ± 4.8%; 14 d, 11.4 ± 3.2%; 28 d, 9.9 ± 2.0%; 56 d, 8.5 ± 2.0%), Tie2 (3 d, 8.4 ± 3.7%; 7 d, 11.7 ± 3.5%; 14 d, 8.5 ± 1.2%; 28 d, 5.5 ± 2.5%; 56 d, 7.0 ± 2.7%), and aggrecan (3 d, 16.3 ± 3.0%; 7 d, 16.2 ± 5.3%; 14 d, 17.2 ± 5.6%; 28 d, 14.0 ± 3.5%; 56 d, 9.1 ± 4.1%) was further low compared to in LASCol-injected discs (all P < 0.001) (Fig. 7AB). The presence of cells with brachyury-positive and/or Tie2-positive disc NP phenotypes supports the accelerated migration of remnant cells through LASCol-mediated spontaneous spheroid formation, potentially contributing to the repair process of nucleotomy-induced disc damage by expressing aggrecan as well as collagen type II.

3.7. Infiltration of macrophages into LASCol in the disc NP of a rat tail nucleotomy model

We further performed additional multi-color immunofluorescence for macrophage markers to identify residual cell migration other than remnant disc NP cells. In LASCol-injected discs, Iba1 (pan-macrophage marker) [41] was positive in a substantial percentage of cells into the LASCol gel (3 d, 59.6 ± 3.8%; 7 d, 67.9 ± 6.7%; 14 d, 58.7 ± 6.9%; 28 d, 48.0 ± 8.5%; 56 d, 32.2 ± 6.0%). This immunopositivity for Iba1 increased from 3 to 7 d (P = 0.04), indicating progressive macrophage infiltration to possibly accelerate LASCol degradation at earlier time points and gradual reduction of macrophages from 14 to 56 d after surgery. Moreover, cells in part were positive for CD86 (M1-polarized macrophage marker for pro-inflammation) [42] (3 d, 34.3 ± 6.0%; 7 d, 35.1 ± 5.3%; 14 d, 31.1 ± 4.5%; 28 d, 23.9 ± 3.3%; 56 d, 16.8 ± 4.2%) and/or CD163 (M2-polarized macrophage marker for anti-inflammation and tissue remodeling) [43] (3 d, 26.3 ± 4.5%; 7 d, 24.2 ± 5.5%; 14 d, 28.6 ± 5.0%; 28 d, 21.7 ± 4.6%; 56 d, 17.5 ± 4.0%) (Fig. 8AB). In AC-injected discs, cells observed on the gel surface were also Iba1-positive (3 d, 33.9 ± 7.0%; 7 d, 27.9 ± 10.8%; 14 d, 28.7 ± 6.3%; 28 d, 29.3 ± 5.7%; 56 d, 18.5 ± 4.2%), although the percentage was lower than in LASCol-injected discs (all P < 0.001). Immunopositivity for CD86 (3 d, 20.0 ± 3.1%; 7 d, 17.7 ± 7.9%; 14 d, 18.9 ± 3.6%; 28 d, 19.0 ± 4.9%; 56 d, 12.5 ± 3.5%) and CD163 (3 d, 14.9 ± 2.9%; 7 d, 9.9 ± 3.5%; 14 d, 9.7 ± 3.4%; 28 d, 14.0 ± 3.1%; 56 d, 8.0 ± 3.0%) was further low (CD86: 3–14 d, P < 0.001; 28 d, P = 0.046; 56 d, P = 0.08; CD163: all P < 0.001) (Fig. 8AB). There were fewer Iba1-positive cells in control discs (3 d, 20.1 ± 5.5%; 7 d, 24.1 ± 7.0%; 14 d, 12.8 ± 4.9%; 28 d, 15.6 ± 5.0%; 56 d, 12.1 ± 3.8%) compared to in LASCol-injected discs (all P < 0.001). Immunopositivity for CD86 (3 d, 10.5 ± 3.8%; 7 d, 11.8 ± 4.0%; 14 d, 7.7 ± 3.5%; 28 d, 9.7 ± 3.5%; 56 d, 8.8 ± 4.3%) and CD163 (3 d, 9.0 ± 3.6%; 7 d, 8.2 ± 1.9%; 14 d, 4.4 ± 3.0%; 28 d, 6.3 ± 3.3%; 56 d, 6.7 ± 2.4%) was also lower (all P < 0.001) (Fig. 8AB). In our proposed model, LASCol-mediated infiltration of M1-polarized and also M2-polarized macrophages would serve for the LASCol self-biodegradation. This could also be beneficial for the tissue repair process, in particular by M2-polarized macrophages. However, effectiveness of macrophage infiltration-mediated secondary inflammatory responses for the disc requires careful evaluations in the future.

4. Discussion

This is the first study to demonstrate the potential for intervertebral disc damage control by LASCol. In in-vitro experiments, compared to
AC, LASCol induced the formation of spheroids with an increased expression of human disc NP-cell and AF-cell phenotypes. In this study, the ratio of Tie2-positive cells on the LASCol gel was higher than that in other reports [9]. As Tie2 expresses at early phases of nucleus pulposus differentiation, LASCol-mediated spheroid formation appears to stimulate the progenitor potential, requiring further mechanistic investigations. In in-vivo experiments, LASCol induced a delayed progression of disc damage in a rat tail nucleotomy model. Radiographic and MRI findings were relatively similar between LASCol and AC, indicating the difficulty of complete structural disc repair by the applied version (concentration, dose, and volume) of both materials—a subject to be studied in the future. However, LASCol had distinct histological differences from AC in the cell number and matrix area of the disc NP. Furthermore, time-course histomorphology and immunofluorescence disclosed a marked migration of endogenous disc NP cells expressing aggrecan and collagen type II and infiltration of M1-polarized and also M2-polarized macrophages into LASCol with an accelerated degradation of collagen type I-positive LASCol. The observed reduction in structural disc damage and spontaneous invasion of endogenous cells and macrophages including the M2 by LASCol, suggesting the future potential for viable disc repair, is notable based on the spheroid-forming capability [22]. A stem cell-seeded tissue-engineered scaffold reports the maintenance of mechanical function and integration up to eight weeks [52]. Hyaluronan-based [53] and alginate-based [54] scaffolds present with enhanced NP-cell phenotypes and matrix production. These scaffolds with a disc regenerative potential are attractive; however, their repair mechanisms have not been fully clarified. At least, no scaffolds other than LASCol have shown spontaneous spheroid formation. While recent advances in the application of stem cells and/or growth factors have enabled radiographic disc height and MRI signal restoration, disc-specific phenotype maintenance, and increased matrix expression in animals [21] and discogenic pain relief and MRI signal improvement in humans [20], these treatments require the specialized institutions, equipment, scientists, technicians, strict laws and regulations, numerous medical expenses, and nevertheless unfavorable health complications. However, LASCol could stimulate disc-cell migration and disc-tissue repair by the scaffold only through a simple, temperature-sensitive gel injection. In addition, treatment of LASCol, developed from abundantly available porcine skin’s collagen type I, is much less expensive than cell transplantation and/or growth factor administration. Therefore, intradiscal LASCol injection is a new scaffold treatment option for non-critical but highly prevalent, intervertebral disc disease.

In this study, in-vivo delayed reduction of safranin-O-positive matrix by LASCol is speculated to come from cells migrating into LASCol expressing aggrecan and collagen type II as well as brachyury and/or Tie2, similar to endogenous disc NP cells. The LASCol would promote the internal migration of remnant disc NP cells, accelerate the formation of cell-aggregating spheroids encouraging the original disc NP

**Fig. 7.** Migration of cells with the endogenous phenotype into LASCol in the disc NP of a rat tail nucleotomy model. (A) Immunofluorescence of rat tail 15-μl LASCol-injected (21.0 mg/ml), AC-injected (7.0 mg/ml), and solvent control discs sectioned before (pre-operation [pre-op]) and at 3, 7, 14, 28, or 56 d after nucleotomy for brachyury (green), Tie2 (red), aggrecan (purple), DAPI (blue), and merged signals. White rectangles indicate the LASCol or AC gel. (B) Changes in the percentage of cells in LASCol-injected, AC-injected, and control disc NP spaces positive for brachyury, Tie2, and aggrecan. Immunopositivity was counted in the whole disc NP low-power fields (× 100) and calculated as relative to the total number of DAPI-positive cells. Data are presented with box plots (n = 6). Two-way repeated measures ANOVA with the Tukey–Kramer post-hoc test was used. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)
phenotype maintenance, e.g. brachyury+, Tie2+, aggrecan+, and collagen type II+, and also up-regulating chondrogenic SOX9, COMP, and TGFB1 gene expression, and facilitate viable disc-tissue repair with the presence of matrix-producing cells. Consequently, this treatment is potentially suitable for discs with early to intermediate stages of degeneration that cells can still be alive [55]. Long-term effectiveness of LASCol to prevent the progression of age-dependent disc degeneration should be studied in the future.

When the efficacy of LASCol depends on endogenous disc NP-cell migration and aggrecan-positive and collagen type II-positive tissue repair, rapid self-biodegradation of collagen type I-based LASCol is advantageous to restore the original tissue environment as the difference in biochemical characteristics between collagen types I (two pro-alpha 1[I] chains and one pro-alpha 2[I] chain) and II (homotrimers of alpha 1[II] chains) is distinct [56]. In this study, infiltration of M2-polarized macrophages as well as M1-polarized macrophages into LASCol is a possible approach to accelerate this degradation. As no macrophages exist in healthy discs but marked macrophage infiltration is observed in aged discs, the M1/M2-polarization of macrophages corresponds to the degeneration severity [57]. Of course, macrophage infiltration into the disc induces mechanical hyperalgesia and pro-inflammatory cytokine production [58]. The originally avascular nature of the disc without macrophage infiltration [57,58] requests a careful interpretation on the involvement of secondary inflammatory responses. Nevertheless, the transition of macrophage polarization from a pro-inflammatory M1 to a pro-healing M2 can be a strategy to improve a poor healing potential of the intervertebral disc [57]. In addition, the design of scaffolds is also critical as macrophages acquire a more tissue-regenerative M2-polarized phenotype on scaffolds with larger fiber and pore dimensions [59]. Following further justifications of the 3D microstructure of scaffolds and roles of inflammation in the disc, the infiltration of M1-polarized and also M2-polarized macrophages could prove useful for not only the biodegradation of LASCol but also the repair process of disc tissues.

This study has several limitations. In vitro, human disc cells surgically obtained had variations in age, sex, and disc degeneration severity, although the observed findings were consistent regardless of these parameters. In vivo, rat tail discs are biologically and mechanically different from rat lumbar discs as well as humans. Rodents retain notochordal cells in the disc NP throughout their lives, resulting in fewer age-related disc pathologies [60]. Mechanical differences between cervical/lumbar and caudal segments and between bipeds and quadrupeds are not negligible [60]. In addition, surgical induction of nucleotomy-induced disc disruption in rat tails does not completely mimic clinical situation of degenerative and herniated discs in humans [61]. Therefore, preclinical studies of LASCol using larger-sized animal models in which disc NP notochordal cells disappear should be carried out. In this study, despite the observed in-vitro compatibility of LASCol with both disc NP and AF cells, in-vivo analysis was primarily focused on disc NP tissues, requiring future clarifications regarding LASCol.

| Immunofluorescence | Rat tail disc NP |
|--------------------|----------------|
| **Pre-op** | **3 d** | **7 d** | **14 d** | **28 d** | **56 d** |
| Iba1 | CD68 | LASCol | CD163 | DAPI |
| Merge | Merge | Merge | Merge | Merge |

Fig. 8. Infiltration of macrophages into LASCol in the disc NP of a rat tail nucleotomy model. (A) Immunofluorescence of rat tail 15-μl LASCol-injected (21.0 mg/ml), AC-injected (7.0 mg/ml), and solvent control discs sectioned before (pre-operation [pre-op]) and at 3, 7, 14, 28, or 56 d after nucleotomy for Iba1 (green), CD86 (purple), CD163 (red), DAPI (blue), and merged signals. White rectangles indicate the LASCol or AC gel. (B) Changes in the percentage of cells in LASCol-injected, AC-injected, and control disc NP spaces positive for Iba1, CD86, and CD163. Immunopositivity was counted in the whole disc NP low-power fields (×100) and calculated as relative to the total number of DAPI-positive cells. Data are presented with box plots (n = 6). Two-way repeated measures ANOVA with the Tukey-Kramer post-hoc test was used. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)
mediated disc repair. Further biomechanical and structural analyses of intradiscal LASCol also need to be conducted. As the material property, alternative LASCol based on collagen type II but not type I might be preferable to repair disc NP tissues since disc NP matrix comprises collagen type II. The development of collagen type II-based LASCol is technically capable; however, the quality control and mass production are difficult due to a limited acquisition amount of raw materials, e.g. pig knee articular cartilage. Collagen type I-based LASCol can be supplied stably and affordably for future clinical applications. It should also be determined in the future which is the best suited shape to the future clinical intradiscal application of LASCol, injectable gel or other shapes including sponge, fragment, flare, and powder.

5. Conclusion

The LASCol, a new collagen type I-based scaffold that we developed by actinidain hydrolysis, has distinct characteristics from conventional AC of an improved water-solubility, faster biodegradability, decreased by actinidain hydrolysis, has distinct characteristics from conventional shapes including sponge, fragment, and powder. Future clinical intradiscal application of LASCol, injectable gel or other pig knee articular cartilage. Collagen type I-based LASCol can be supplied stably and affordably for future clinical applications. It should also be determined in the future which is the best suited shape to the future clinical intradiscal application of LASCol, injectable gel or other shapes including sponge, fragment, flare, and powder.

Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.biomaterials.2020.119781.

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