Factors secreted from dental pulp stem cells show multifaceted benefits for treating experimental temporomandibular joint osteoarthritis

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SUMMARY
Objective: Temporomandibular joint osteoarthritis (TMJOA) is a degenerative disease characterized by progressive cartilage degeneration, abnormal bone remodeling, and chronic pain. In this study, we aimed to investigate effective therapies to reverse or suppress TMJOA progression.
Design: To this end, we performed intravenous administration of serum free conditioned media from human exfoliated deciduous teeth stem cells (SHED-CM) into a mechanical-stress induced murine TMJOA model.
Results: SHED-CM administration markedly suppressed temporal muscle inflammation, and improved bone integrity and surface smoothness of the destroyed condylar cartilage. Moreover, SHED-CM treatment decreased the number of IL-1β, iNOS, and MMP-13 expressing chondrocytes, whereas it specifically increased PCNA-positive cells in the multipotent polymorphic cell layer. Notably, the numbers of TdT-mediated dUTP nick end labeling (TUNEL)-positive apoptotic chondrocytes in the SHED-CM treated condyles were significantly lower than in those treated with DMEM, whereas the proteoglycan positive area was restored to a level similar to that of the sham treated group, demonstrating that SHED-CM treatment regenerated the mechanical-stress injured condylar cartilage and subchondral bone. Secretome analysis revealed that SHED-CM contained multiple therapeutic factors that act in osteochondral regeneration.
Conclusions: Our data demonstrated that SHED-CM treatment promoted the regeneration and repair of mechanical-stress induced mouse TMJOA. Our observations suggest that SHED-CM has potential to be a potent tissue-regenerating therapeutic agent for patients with severe TMJOA.

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Introduction
The temporomandibular joint (TMJ) is a load-bearing articulation with a wide range of motion consisting of both rotation and translation. The articular cartilage in general synovial joints is a thin layer of hyaline cartilage that covers the entire articulating surface of each bone. The mandibular condylar cartilage however, uniquely consists of four layers: 1) a superficial fibrous layer covering the articular, 2) a proliferating polymorphic cell layer having multi-lineage differentiation potency toward osteoblasts, chondrocytes,
and adipocytes, 3) a mature hyaline-like cartilage layer producing cartilage matrix, and 4) a hypertrophic cell layer initiating endochondral ossification. This heterogeneous architecture plays an important role in maintaining condylar structure and function in the face of mechanical stress.

TMJ osteoarthritis (TMJOA) is a degenerative joint disease, characterized by progressive cartilage degeneration, abnormal bone remodeling, and chronic pain. The etiology of TMJOA is complex and multifactorial. However, it has been suggested that excessive malocclusion-induced mechanical stress affects TMJOA progression. It has also been shown that injury-activated chondrocytes play important roles in the pathophysiology of early stage TMJOA. Activated chondrocytes increase production of cartilage-degrading catabolic enzymes, proinflammatory cytokines, oxidative stress-inducers, and osteoclast-activators, such as matrix metalloproteinase-13 (MMP-13), interleukin-1β (IL-1β), inducible nitric oxide synthesis (iNOS), and receptor activator of nuclear factor (NF)-κB ligand (RANKL), respectively. During the progression phase, these tissue-degenerative TMJOA microenvironments promote degradation of cartilage matrix, chondrocyte apoptosis and necrosis, and abnormal resorption of subchondral bone, leading to irreversible articular damage and functional impairment. As the self-repair capability of the articular cartilage is limited, development of effective therapies to reverse or suppress TMJOA progression are currently under investigation.

Stem cell therapy holds great promise for the development and establishment of effective treatments for TMJOA and osteoarthritis (OA). It has been shown that transplantation of undifferentiated adult mesenchymal stem cells (MSCs) isolated from adipose tissue or bone marrow improved the OA-phenotype in clinical studies of animal TMJOA and Knee-OA models. However, most of these studies reported poor cell graft survival, suggesting that recovery of articular damage was primarily accomplished through paracrine trophic mechanisms. Stem cells secrete a broad repertoire of trophic and immunomodulatory factors, which can be collected as a serum-free conditioned medium (CM). Conditioned media from various stem cell types have been shown to exhibit considerable potential in treating a myriad of intractable diseases.

Human adult dental pulp stem cells (DPSCs) and stem cells from human exfoliated deciduous teeth (SHED) are self-renewing MSCs residing within the perivascular niche of the dental pulp. These cells are thought to originate from the cranial neural crest, which expresses early markers for both MSCs and neuroectodermal stem cells. Implantation of DPSCs, pre-differentiated toward chondrocytes, resulted in significant cartilage regeneration in a rabbit model of knee cartilage damage. The study demonstrated the therapeutic potential of local delivery of ex vivo culture-expanded DPSCs for treating OA. However, the benefits of these DPSC-secreted factors in the treatment of TMJOA, have not been examined. Notably, systemic administration of SHED serum free conditioned medium (SHED-CM) promoted functional recovery in various acute and chronic intractable disease models, such as ischemic brain injury, Alzheimer's disease, spinal cord injury, fulminant liver failure, liver fibrosis, bleomycin-induced fibrotic lung injury, autoimmune encephalomyelitis, type 1 diabetes, rheumatoid arthritis, and cardiac injury. In the present study, we investigated the therapeutic effects of SHED-CM administration in a mouse mechanical stress-induced TMJOA experimental model.

Methods

Preparation of SHED-CM

Human SHEDs were isolated and cultured as previously described. In brief, deciduous teeth from six- to twelve-year-old individuals were collected at Nagoya University and Tokushima University Hospital. This study was approved by the Institutional Ethical Committee of Nagoya University and Tokushima University Hospital and performed according to the principles of the Declaration of Helsinki (Permit No H-73 and No: 3,268 for Nagoya and Tokushima University, respectively). The pulp was gently removed and digested for 1 h at 37 °C in a solution containing collagenase type I (3 mg/ml) and dispase (4 mg/ml). Single-cell suspensions were plated on culture dishes in Dulbecco's Modified Eagle Medium (DMEM) supplemented with 10 % fetal bovine serum (FBS), then incubated at 37 °C in 5 % CO2. The human skin fibroblast line, derived from a 36-year-old individual, was obtained at passage 12 from the Health Science Research Resources Bank (Osaka, Japan). SHED (passage 9) and fibroblasts (passage 13) for CM preparation were seeded at 1 × 105 cells per dish. Cells at 70–80 % confluence were washed with phosphate-buffered saline (PBS) and serum-free DMEM twice, followed by culture media replacement with serum-free DMEM. Media were incubated for 48 h at 37 °C in a humidified atmosphere of 5 % CO2, then collected and centrifuged for 3 min at 440 g and 4 °C. Supernatants were then used as SHED-CM in TMJOA treatment protocols. We adjusted the protein concentration of each CM to 3 μg/ml with serum-free DMEM.

Mouse mechanical stress-induced TMJOA model and treatment with SHED-CM

Institute of Cancer Research (ICR) male mice, aged 11 weeks, were purchased from Japan SLC Inc (Shizuoka, Japan), kept individually in plastic cages, and maintained at ambient temperature (22–24 °C) in a 12 h light/dark cycle. Mice were fed a standard solid diet with water ad libitum throughout the experiment and were randomly divided into one control and three experimental groups (n = five for each group). An overview of the experimental design and workflow is presented in Supplemental Fig. 1.

In the experimental group, mechanical stress was applied to both TMJs by forced mouth opening for 3 h/day for five or 10 consecutive days by using a custom-made spring. This device kept the mandible in a maximal mouth opening position of 14 mm, and delivered a force of 2 N on each TMJ. As the mouth opening experiments, mice were anesthetized with a mixture of medetomidine (Domitor, Orion Corporation, Espoo, Finland), midazolam (Midazolam, Meiji Seika Pharma, Tokyo, Japan) and butorphanol (Vetorphale, Meiji Seika Pharma, Tokyo, Japan) at doses of 0.75, 4.0 and 5.0 mg/kg, respectively.

In one experimental group, both TMJs received only mechanical stress for 5 days only (pre-treatment group), while those in the other experimental groups received mechanical stress for 10 consecutive days and were injected daily with either 0.5 ml SHED-CM (SHED-CM group) or 0.5 ml DMEM (DMEM group) into the tail vein, from day six to day 10 [Fig. 1(B)]. For immunohistological analysis (except Proliferating Cell Nuclear Antigen, PCNA), mechanical stress followed by SHED-CM or DMEM treatment was carried out for three consecutive days [Fig. 1(C)]. For PCNA evaluation, mechanical stress was carried out for 6 days and mice received SHED-CM or DMEM on the last day [Fig. 1(D)]. When the animals were sacrificed, blood plasma was isolated and sent to SRL, Inc (Tokyo, Japan) where the level of C-reactive protein (CRP) was determined. Although no mechanical stress was applied to the TMJs of mice in the control group, the same anesthesia schedule was maintained. The weight of each mouse was measured immediately after the experiment (day 0), at the initial day of injection (day 6), and at the end of the experiment (day 10). All procedures performed in this study were approved by the Tokushima University Animal Care and Use Committee (Permit No: T30-95) and carried
out according to the Animal Research: Reporting of In Vivo Experiments (ARRIVE) guidelines.

**Pain behavioral measurement**

Mechanical hyper-nociception was assessed using the von Frey microfilament procedure under blind conditions. A series of calibrated von Frey filaments (Aesthesio, Damnic global, CA, USA) were applied to the TMJ region. The head withdrawal threshold is defined as the lowest force applied by the filaments that produced a withdrawal response at least three times. Assessment of mechanical hyper-nociception were performed every day and the head withdrawal threshold was calculated as the mean value per joint of five mice per group.

**Micro-computed tomography (Micro-CT) analysis**

Murine mandibles from mice of all groups were resected, and the condyles were carefully separated from the soft surrounding tissues and fixed overnight in 70% ethanol. Condyles were then analyzed using high-resolution micro-CT (SkyScan 1,176 scanner and associated analysis software; Buruker, Billerica, MA, USA). Image acquisition was performed at 50 kV and 200 μA. During image acquisition, samples were tightly covered with plastic wrap to prevent movement and dehydration. To segment the bone image from background noise, thresholding was applied. The posterior region of the mandibular condyle in the midsagittal section was the region of interest (ROI). The resolution of the captured micro-CT images was 9 μm/pixel. For each ROI, microstructural parameters including the bone volume to trabecular volume ratio (BV/TV), trabecular thickness (Tb.Th), and trabecular separation (Tb.Sp) were analyzed.

**Histology and immunohistochemical analysis**

Following micro-CT examination, fixed TMJ specimens were decalcified in 30% ethylene diamine tetra acetic acid (EDTA) for 2 weeks prior to histology processing. Samples were dehydrated and embedded in paraffin for microtomy processing according to standard procedures. Tissue segments were isolated from three mice per experimental group, and subsequently, 5 μm sagittal sections were generated. Sections were deparaffinized and hydrated through immersion in a series of xylene and graded alcohol solutions. Sections were stained with hematoxylin and eosin (HE) for histological assessment and with toluidine blue (TB) for visualizing proteoglycans. Tartrate-resistant acid phosphatase (TRAP) staining identifying active bone-absorbing osteoclasts was carried out using a TRAP/AP staining kit (Wako, Osaka, Japan) according to the manufacturer’s instructions. Apoptotic cell death was analyzed using the TdT-mediated dUTP nick end labeling (TUNEL) assay (In situ Cell Death Detection Kit, Roche Life Science, Geneva, Switzerland) according to the manufacturer’s instruction.

For immunohistochemical analysis, sections were blocked with 5% (v/v) bovine serum albumin for 30 min, and incubated overnight with the following primary antibodies: rabbit anti-MMP13 IgG (1:100, ab75606, Abcam, Cambridge, UK), rabbit anti-iNOS IgG (1:500, ab3523, Abcam), rabbit anti-IL-1β IgG (1 μg/ml, ab9722, Abcam), and rabbit anti-PCNA IgG (1:250, ab92552, Abcam). As secondary antibody, the anti-rabbit IgG (VECTASTAIN Elite ABC, Vector Laboratories) was used and processed by diaminobenzidine (DAB) staining using the ABC reaction protocol (Vector Laboratories). After counterstaining with Mayer’s Haematoxylin (Sakura Finetek Japan Co, Tokyo, Japan), tissue images...
were captured using a universal fluorescence microscope (BZ9000; Keyence, Osaka, Japan). Time schedules of immunohisto staining for MMP13, iNOS, IL-1β and PCNA evaluation are presented in Fig. 1(B) and (C), respectively. Cells were counted in at least five images captured from three individual specimens in parallel experiments.

**LC-MS/MS analysis**

Secretome analysis of CM was performed according to a previously reported protocol. In brief, SHED-CM and fibroblast-CM (Fibro-CM) were concentrated using Amicon Ultra 3K filter (Millipore, Burlington, MA, US). For protein purification, methanol/chloroform precipitation was performed. Resulting pellets were...
resolved by MS buffer (8 M urea and 50 mM Tris–HCl pH 8.0), followed by reduction in 5 mM DTT (WAKO, Osaka, Japan) and alkylation in 27.5 mM iodoacetamide (WAKO, Osaka, Japan) for 30 min in the dark at room temperature. After being diluted 8 times in 50 mM Tris–HCl pH 8.0, proteins were digested with 50 ng Lys-C (WAKO, Osaka, Japan) and trypsin (Promega, Madison, WI, US) at 37 °C overnight. Peptides were purified by GL-Tips SDB (GL science, Tokyo, Japan) according to the manufacturer’s protocol. The concentration of peptides was measured using the Pierce™ Quantitative Colorimetric Peptide Assay Kit (Thermo, Waltham, MA, USA). Subsequently, peptides (268 ng each) were injected to an EASY-nLC™ connected to a Q-Exactive PLUS™ (Thermo) mass spectrometry system. Protein identification and label-free quantification were performed by Proteome Discoverer 2.2™ (Thermo).

Statistical analysis

The SPSS software package, version 22.0 (IBM, New York, USA) was used for statistical analysis. All data are expressed as the mean ± standard error of the mean (SEM). Differences between groups were compared using the Student’s t test or Mann–Whitney U test, according to the data type. To analyze three or more independent groups, we used repeated-measures analysis of variance (ANOVA) with Tukey’s post hoc test. A p value < 0.05 was considered to be statistically significant.

Results

SHED-CM restored bone integrity and surface properties of condyles in mice with TMJOA

Micro-CT images revealed a rougher condylar cartilage surface with severe subchondral trabecular bone loss in the pre-treatment and DMEM groups, whereas that of the SHED-CM group had a smoother surface and decreased subchondral trabecular bone resorption [Fig. 1 (E)]. We also measured BV/TV, Tb.Th, and Tb.Sp in several areas of the condylar subcondral bone. Notably, BV/TV was significantly (p < 0.01) higher in the SHED-CM group (0.42 ± 0.05 %) relative to the pre-treatment (0.25 ± 0.07 %) and DMEM groups (0.27 ± 0.06 %). The SHED-CM group also had a higher Tb.Th value (0.16 ± 0.08 μm) compared to the pre-treatment (0.09 ± 0.02 μm) and DMEM groups (0.11 ± 0.03 μm), with a significant difference (p < 0.05) noted between the SHED-CM and DMEM groups. Concurrently, Tb.Sp was significantly (p < 0.05) smaller in the SHED-CM group (0.21 ± 0.11 μm) relative to the pre-treatment (0.39 ± 0.25 μm) and DMEM groups (0.32 ± 0.17 μm) [Fig. 1 (F)].

SHED-CM suppressed masticatory muscle inflammation and pain and restored cellular alignment and matrix deposition of articular cartilage after TMJOA

Hematoxylin staining showed a massive infiltration of mononuclear immune cells into the damaged temporal muscle (TM) in the DMEM group, whereas the SHED-CM treated group exhibited markedly decreased infiltration. Of note, the inflammatory response in the synovial membrane (SM) in both the DMEM and SHED-CM groups was weak, suggesting the non-inflammatory nature of our mechanical stress-induced TMJOA model compared to the strong inflammatory nature of rheumatoid arthritis and synovitis [Fig. 2 (A)]. We further examined the effects of SHED-CM for TMJOA-induced mechanical hyper-nociception using the von Frey microfilament procedure. We found that SHED-CM treatment significantly suppressed TMJOA-induced hypersensitivity as indicated by reduced pain following SHED-CM treatment [Fig. 2 (B)].

Regarding the mandibular condyles, both pre-treatment and DMEM groups showed obvious OA-like lesions, such as a decrease in the condylar cartilage layer thickness and hyalinization of the cartilaginous matrix, and an increase in chondrocyte alignment irregularities. In contrast, the condylar cartilage of the SHED-CM group was similar to that of the sham control group, displaying obviously distinguished cell layers [Fig. 2 (C)]. Toluidine blue stained sections demonstrated that the proteoglycan positive area was significantly (p < 0.01) decreased in the pre-treatment and DMEM groups, accompanied with marked morphological changes. Meanwhile, condyles in the SHED-CM group were similar to those in the sham control [Fig. 2 (C) and (E)]. Modified Mankin morphological scores were significantly lower (p < 0.01) in the mandibular condylar cartilage samples from the SHED-CM group compared to those from the pre-treatment and DMEM groups [Fig. 2 (D)].

To evaluate the adverse side-effects of CM-treatment, we examined plasma CRP levels. We found that there were no significant differences between the CRP levels of the sham operated, pre-treatment, DMEM-treated, and SHED-CM-treated groups [Supplemental Table 1].
SHED-CM suppressed osteoclastogenic activity in the subchondral bone layers of condyles

TRAP-positive osteoclasts were observed in the mineralized subchondral bone layers of mandibular condyles [Fig. 3 (B)]. The subchondral bone samples from the DMEM group (13.20 ± 4.36) exhibited significantly higher numbers of TRAP-positive cells (p < 0.01) relative to those from the SHED-CM (3.40 ± 2.06) and control groups (3.67 ± 1.70) [Fig. 3 (D)]. Taken together, this indicates that SHED-CM administration might downregulate osteoclastogenesis, leading to a decrease in bone resorption.

SHED-CM inhibited chondrocyte apoptosis in the subchondral bone of condyles

TUNEL assays were performed to determine whether apoptosis of abnormal chondrocytes was preferentially induced in degraded cartilage [Fig. 3 (A)]. An obvious increase in the number of TUNEL positive cells was observed in the DMEM group (76.97 ± 11.18 %) compared to the SHED-CM group (34.26 ± 6.97 %) [Fig. 3 (C)], SHED-CM injection led to a decrease in TUNEL positive cells, thus providing a protective effect against cartilage degradation.

SHED-CM suppressed the expression of cartilage degradation-associated factors and promoted the proliferation of multipotent polymorphic cells

Immunohistological analysis showed that expression of the pro-inflammatory factor IL-1ß (95.56 ± 2.02 %) and cartilage matrix degradation factors MMP-13 (44.43 ± 7.90 %) and iNOS (48.8 ± 3.96 %) were elevated in the DMEM group when compared to the sham treated control (IL-1ß; 79.96 ± 4.81 %, MMP-13; 26.69 ± 1.52 %, iNOS; 38.96 ± 1.63 %) [Fig. 4(A) and (B), (D), (E), (H), (I) and (N)]. The number of cells found positive for these factors increased in both multipotent polymorphic cells and the mature chondrocyte layer. In contrast, signal intensities of those exacerbating factors were reduced (IL-1ß; 80.60 ± 4.95 %, MMP-13; 32.15 ± 5.78 %, iNOS; 38.72 ± 2.52 %) in the SHED-CM group [Fig. 4 (C), (F), (J) and (N)]. Furthermore, the majority of multipotent polymorphic cells were negative for these factors. Importantly, we found that PCNA-positive proliferating polymorphic cells were markedly increased in the SHED-CM treated group (sham group; 11.92 ± 3.40 %, DMEM group; 5.95 ± 2.70 %, SHED-CM group; 45.13 ± 4.56 %) [Fig. 4(K)–(N)]. These findings suggest that SHED-CM administration can protect against cartilage destruction and enhance cartilage formation in patients with TMJOA.

Secretome analysis of conditioned medium

Secretome analysis of SHED-CM resulted in the identification of 1,426 proteins. According to Gene Ontology cellular component classification, the number of cell surface, extracellular, or membrane proteins was 894. SHED-CM contained 51 of these proteins at levels at least 10 times greater than detected in Fibro-CM [Fig. 5 and Supplemental Fig. 2]. These molecules are known to be involved in the processes of anti-fibrosis, anti-apoptosis, anti-inflammation, proliferation, differentiation, and migration of chondrocytes.

Discussion

We previously reported on the therapeutic effects of systemically administrated SHED-CM against anti-collagen type II antibody-induced arthritis (CAIA)18. However, no study to date has examined the effect of SHED-CM administration against TMJOA. To the best of our knowledge, this is the first study reporting on the promising potential of SHED-CM administration as an effective treatment of TMJOA. In this study, we used a mechanical-stress induced TMJOA model consisting of five consecutive days of TMJ damage followed by treatment with DMEM or CM for the next five days of TMJ damage. The initial TMJ damage induced pain and reduced condylar cartilage thickness and matrix deposition. Subsequent DMEM treatment worsened the pathophysiology of TMJOA. In contrast, SHED-CM treatment reduced pain and repaired osteoarthritic TMJ through the inhibition of the pro-inflammatory response, apoptosis, and matrix degradation while enhancing proliferation and matrix deposition of the cartilage layer. Our study revealed the remarkable articular cartilage regeneration activity of SHED-CM, which restored homeostasis in the mechanically destroyed TMJ.

Similar to previous studies, the TMJOA experimental model in the present study was established by applying mechanical stress to the mandibular condyle24–26. This model was characterized by OA-like degenerated lesions, chondrocyte alignment irregularities in the condylar cartilage layers, subchondral bone loss, and marked depletion of proteoglycans. Furthermore, these results were consistent with results previously reported for early TMJOA generated by surgical manipulation of the joint24, local application of chemicals25,26. IGF-1, a pleiotropic growth factor, has been involved in osteoarthritis, synovitis, TMJOA has a primarily non-inflammatory origin26,37. The pathological process in TMJOA is characterized by deterioration and abrasion of the articular cartilage and local thickening and remodeling of the underlying bone26. These changes are frequently accompanied by the superimposition of secondary inflammatory reactions. Surgical, genetic, or chemical methods cannot successfully mimic the real conditions encountered in patients with TMJOA, because of the high inflammatory arthritic conditions present in these induced animal models. Excessive mechanical stress is characterized as a key factor inducing mandibular condylar cartilage degradation in the TMJ. Therefore, our mechanical stress protocol of forced mouth opening is a useful tool for generating an experimental TMJOA model that can be used to evaluate the initiation and advancement of TMJOA.

Our data show that systemic intravenous administration of SHED-CM effectively repaired the lesions induced by TMJOA. Local injection allows for lower drug dosages and elicits fewer side effects, both systemic and local, when compared to conventional systemic intravenous injection. However, the needle pricks for local intra-articular injection can cause substantial tissue damages in TMJ. Our data show that the level of CRP in the SHED-CM treated group was similar to that of the sham-operated group, indicating there were little to no adverse side-effects after CM-treatment. Moreover, a small volume of systemic CM injection effectively restored injured TMJ. Taken together, systemic CM injection may avoid the iatrogenic disorders caused by needle pricks and provide considerable advantages in a clinical setting.

We characterized the soluble factors in SHED-CM by performing a LC-MS/MS analysis and found that SHED-CM contained 51 of the array proteins at levels more than 10-fold higher than those detected in Fibro-CM. A cluster analysis of these proteins identified 12 proteins known to provide beneficial effects in treating TMJOA and OA (Fig. 5). For instance, the identified insulin-like growth factor (IGF) and hepatocyte growth factor (HGF) are multifunctional proteins that protect cells from apoptosis and suppress inflammatory reactions. In cartilage repair, IGF promotes proliferation and maturation of chondrocytes, and the synthesis of their extracellular matrix. In bone remodeling, IGF enhances osteoblastogenesis, matrix deposition, and mineralization38,39. Although the circulating half-life of the free form of IGF is reported to be around 20 min, IGF binding protein 5 (IGFBP-5) may form a complex with IGF in SHED-CM and extend its half-life40. HGF enhances cellular motility,
proliferation, and proteoglycan synthesis of chondrocytes and activates osteoblast for bone regeneration. Transforming growth factor β (TGF-β) promotes the differentiation of chondrocytes and the synthesis of collagen and proteoglycans, thereby maintaining cartilage homeostasis. R-Spondin 2 (SPON2), an activator of Wnt/b-catenin signaling, promotes endochondral ossification through induction of chondrogenic differentiation. Secreted frizzled related protein 1 (SFRP1), a Wnt signaling antagonist, is required for the maintenance of immature proliferating chondrocyte precursors. Syndecan (SDC), a cell surface heparan sulfate proteoglycan, promotes chondrocyte proliferation and maintains cartilage matrix homeostasis. Platelet derived growth factor (PDGF), is a potent mitogenic and chemotactic factor for all mesenchymal originating cells, including chondrocytes and MSCs, while pleiotrophin (PTN) and midkine (MDK) are involved in the mitotic activation of chondrocytes. It has been reported...
that MDK, a heparin-binding growth factor, promotes the proliferation of articular chondrocytes in vivo when administrated subcutaneously in normal mice\textsuperscript{48}, suggesting that systemically administrated MDK may reach the joint. Despite the relatively low concentrations of chondrogenic factors (1–10 ng/ml) measured in SHED-CM, the combinatorial effects of these multiple factors may provide prominent therapeutic benefits in treating TMJ OA. To clarify the detail mechanisms of SHED-CM-mediated articular cartilage regeneration, in future it is necessary to examine the accessibility of therapeutic factor in SHED-CM toward injured TMJ.

Recent reports have revealed that MSCs-derived extracellular vesicles (EVs) play important roles in the trophic effects exhibited in MSCs-transplantation therapy. Importantly, several studies have demonstrated a cartilage regenerating activity of MSCs-derived exosomes, a type of EVs released from cells through fusion of endosomal multivesicular bodies with the plasma membrane\textsuperscript{49}. Exosomes carry multiple heterogeneous proteins, micro-RNAs and lipids, which are involved in many diverse biochemical and cellular processes, such as communication, structure and mechanics, inflammation, exosome biogenesis, tissue repair and regeneration.

Fig. 5

LC-MS/MS analysis of SHED-CM and Fibro-CM. In SHED-CM, a total of 1,426 proteins were identified. From these, 51 proteins were more than 10 times higher in SHED-CM compared to the Fibro-CM.
and metabolism. Exosomes express endosome-related proteins, such as Alix, tumor susceptibility 101 (TSG101), integrin α chains, and tetraspanins (TSPANs, CD9, CD63, CD81), which are members of a protein superfamily characterized by the presence of four transmembrane and two extracellular domains. In the list obtained from our SHED-CM proteome analysis, we identified many exosome markers, such as TSPAN 4, CD9, and Integrin α 3 (ITGA3), suggesting that SHED cells may produce a larger number of exosomes than previously thought. In the future, it would be necessary to clarify the roles of exosomes and trophic factors in SHED-CM-mediated cartilage and bone regeneration of the mandibular condylar.

As previously described, TMJOA is characterized by mandibular condylar cartilage degradation due to mechanical stress. Mechanical stress of the mandibular condylar cartilage induces the expression of IL-1β, an inflammatory cytokine closely related to the progression of TMJOA. Since the fibrocartilage covering both the mandibular condyle and articular eminence is avascular, these fibrocartilage cells have limited ability for self-repair, similar to the hyaline cartilage in other synovial joints. Therefore, once the breakdown in the joint starts, TMJOA can be crippling, leading to a variety of morphological and functional deformities. This highlights the importance of suppressing cartilage degradation during the early stages of TMJOA. However, so far no treatment remedy has been developed for the management of severe TMJOA. Furthermore, despite the fact that many studies have demonstrated the potential of using embryonic stem (ES) cells for cartilage regeneration, many concerns raised by the use of ES cells, such as possible tumorigenesis, ethical issues regarding the use of embryos and potential allo-geneic immune rejection, make this approach problematic. The use of induced pluripotent stem (iPS) cells may overcome most of these issues. However, this technology is still in its infancy and includes many unknown parameters. In the present study, we used SHED-CM for the treatment of TMJOA through cartilage regeneration and our results suggested that SHED-CM administration can attenuate cartilage degeneration, promote cartilage regeneration, and protect against the progression of TMJOA induced by mechanical stress. Therefore, SHED-CM administration may represent an effective treatment strategy for patients with mechanical stress-induced TMJOA.

In conclusion, we showed that SHED-CM contained multiple therapeutic factors with the potential for treating TMJOA. Importantly, we did not observe any adverse side-effects during the experimental treatment period. Our findings suggest that SHED-CM not only inhibited the articular degradation cascade but also regenerated the injured articular by promoting proliferation of the multipotent polymorphic cell layer and production of cartilage matrix in mice with TMJOA. Thus, SHED-CM may provide a novel articular regenerative therapy for patients with severe TMJOA.

Studies involving humans or animals

This study was approved by the Institutional Ethical Committee of Nagoya University and Tokushima University Hospital and performed according to the principles of Helsinki Declaration (Permit No H-73 and No: 3,268 for Nagoya and Tokushima University, respectively). All procedures performed in this study involving animals were approved by the Tokushima University Animal Care and Use Committee (Permit No: T30-95).

Contributions
Study conception and design: NO, ET, AY. Acquisition of data: NO, FK, NH, HM, YL, LX, TS. Provision of study materials or patients: HH, TI. Analysis & interpretation of data: All authors. Writing of first manuscript draft: NO, ET, AY. Critical manuscript revision and approval of final manuscript: All authors. AY had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

Conflict of Interest
The authors declare that there is no conflict of interest.

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Supplementary data
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References


