Culturing bone marrow cells with dexamethasone and ascorbic acid improves osteogenic cell sheet structure

M. Akahane, T. Shimizu, T. Kira, T. Onishi, Y. Uchihara, T. Imamura, Y. Tanaka

Nara Medical University Faculty of Medicine, Nara, Japan

Objectives
To assess the structure and extracellular matrix molecule expression of osteogenic cell sheets created via culture in medium with both dexamethasone (Dex) and ascorbic acid phosphate (AscP) compared either Dex or AscP alone.

Methods
Osteogenic cell sheets were prepared by culturing rat bone marrow stromal cells in a minimal essential medium (MEM), MEM with AscP, MEM with Dex, and MEM with Dex and AscP (Dex/AscP). The cell number and messenger (m)RNA expression were assessed in vitro, and the appearance of the cell sheets was observed after mechanical retrieval using a scraper. β-tricalcium phosphate (β-TCP) was then wrapped with the cell sheets from the four different groups and subcutaneously implanted into rats.

Results
After mechanical retrieval, the osteogenic cell sheets from the MEM, MEM with AscP, and MEM with Dex groups appeared to be fragmented or incomplete structures. The cell sheets cultured with Dex/AscP remained intact after mechanical retrieval, without any identifiable tears. Culture with Dex/AscP increased the mRNA and protein expression of extracellular matrix proteins and cell number compared with those of the other three groups. More bridging bone formation was observed after transplantation of the β-TCP scaffold wrapped with cell sheets cultured with Dex/AscP, than in the other groups.

Conclusions
These results suggest that culture with Dex/AscP improves the mechanical integrity of the osteogenic cell sheets, allowing retrieval of the confluent cells in a single cell sheet structure. This method may be beneficial when applied in cases of difficult tissue reconstruction, such as nonunion, bone defects, and osteonecrosis.

Cite this article: Bone Joint Res 2016;5:569–576.

Keywords: Osteogenesis; Cell sheet; Mechanical retrieval

Article focus
This study focused on the mechanical retrieval technique to create osteogenic cell sheets.

Key messages
Culture with dexamethasone/ascorbic acid phosphate improves the mechanical integrity of osteogenic cell sheets, allowing retrieval of confluent cells in a cell sheet structure.

Strengths and Limitations
Strength: this method may be beneficial for cases of difficult tissue reconstruction, such as nonunion and bone defects.

Limitations: Future studies should focus on human bone marrow stromal cells in order to determine if culture with Dex/AscP improves the integrity of osteogenic cell sheets.

Introduction
Mesenchymal stem cells (MSCs) are adult stem cells which can be isolated from tissues such as bone marrow,1,2 adipose,3 and skeletal muscle.4 Because MSCs are multipotent and can differentiate into osteoblasts,2,5 chondrocytes,6 and neurons,7,8 they have been used in regenerative medicine to treat
various diseases and tissue injuries, including tissue engineering using an in vitro cell culture. Bone marrow stromal cells (BMSCs) are commonly used for clinical treatment because they can be harvested from a patient’s pelvis by needle aspiration, a minimally invasive procedure compared with harvesting bone. Therefore, BMSC populations containing MSCs are a good source for creating tissue-engineered bone.

For clinical applications, the BMSCs harvested from patients are expanded in vitro, and then seeded onto scaffolding materials such as hydroxyapatite (HA) and β-tricalcium phosphate (β-TCP). When scaffolds seeded with BMSCs are transplanted into patients, more cells that are seeded on the scaffolds at implantation result in better bone formation. BMSCs are able to differentiate into osteoblastic lineage cells after subculture in osteogenic induction media containing dexamethasone (Dex), ascorbic acid phosphate (AscP), and β-glycerophosphate (β-GP). Combinations of BMSCs with a scaffolding material, such as HA and β-TCP, promote the formation of new bone tissue in vivo following subcutaneous transplantation, both when using freshly isolated bone marrow cells and culture-expanded BMSCs.

Many recent studies have reported methods for creating cell sheets including thermo-responsive polymer-grafted culture dishes and mechanical retrieval. We have previously proposed solutions to undertake difficult tissue regeneration, including fracture nonunion and ligament reconstruction using scaffold-free cell sheet transplantation. We have also reported a technique for cell sheet injection we call ‘injectable bone’, that promotes osteogenesis in necrotic bone and implanted materials.

Preparation of the bone marrow stromal cells. BMSCs were prepared as previously reported. Briefly, BMSCs were obtained from the femur shafts of male seven-week-old Fischer rats. Male rats were chosen as they are larger than females and are therefore easier to handle. Both ends of each femur were removed from the epiphysis and the marrow was flushed out using 10 mL of standard culture medium, which consisted of MEM containing 15% FBS and antibiotics, expelled from a syringe through a 21-gauge needle. The obtained cells were collected in two T-75 flasks containing the standard culture medium. Once they were confluent, the cells were released from the culture substratum using trypsin/EDTA. In the present study, BMSCs were prepared from three different donor rats because the characteristics of BMSCs from primary culture may vary among each individual.

In vitro cell culture. The released BMSCs were seeded at a density of 1×10⁴ cells/cm² in order to assess cell sheet formation by macroscopic and microscopic observation. The cells were cultured with the standard medium alone (MEM group), the standard medium with 0.28 mM AscP (AscP group), the standard medium with 10 nM Dex (Dex group), or the standard medium with 10 nM Dex and 0.28 mM AscP (Dex/AscP group) for 14 days. The BMSCs were cultured in a humidified atmosphere of 95% air with 5% CO₂ at 37°C in 96-well plates for the cell proliferation assay, and 6-well plates for the assessment of ECM component expression and Western blotting. Macroscopic and microscopic observations were made of cells cultured in 100 mm dishes, by eye.

MTT assay. The cell proliferation of each group was measured using the MTT assay (3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulphophenyl)-2H-tetrazolium). Cells were seeded onto 96-well culture plates (1×10⁴ cells/cm²) and cultured with 100 µL of medium from each group for five days (n = 5). Then, the cell proliferation was determined using CellTiter 96 Aqueous One Solution Reagent (Promega, Madison, Wisconsin). Once the medium was removed and the cells were washed with PBS, 100 µL of medium and 20 µL of MTS solution were added per well and incubated for one hour at 37°C. Finally, formazan absorbance at 490 nM was measured in a microplate reader. The experiments in each group were performed in triplicate.

Materials and Methods

Experimental materials. Male seven-week-old Fischer rats were purchased from Japan SLIC Inc. (Shizuoka, Japan). Minimal essential medium (MEM), antibiotics (100 U/mL penicillin and 100 µg/mL streptomycin) and Trypsin/ethylene-diaminetetraacetic acid (EDTA) were purchased from Nacalai Tesque (Kyoto, Japan). Culture flasks and plates were from Corning (Corning, New York, New York), and fetal bovine serum (FBS) and phosphate-buffered saline (PBS) from Gibco (Life Technologies, Carlsbad, California). AscP was also purchased from Wako Pure Chemical Industries (Kyoto, Japan). Dex was purchased from Sigma (St. Louis, Missouri). Primer and probe sets and reagents for real-time quantitative polymerase chain reaction (PCR) were purchased from Applied Biosystems (Norwalk, Connecticut). β-TCP granules (OSferion 60, 60% porosity) were purchased from Olympus (Tokyo, Japan), and RNaseasy Mini Kits were purchased from QIAGEN Inc. (Venlo, The Netherlands).
RNA isolation and real-time quantitative PCR. The gene expression levels of collagen types I, laminin, fibronectin and osteocalcin were measured in each group. Total RNA was isolated from each group using RNeasy Mini RNA extraction kits. Cultured cells from each group in 6-well plates were rinsed twice with PBS and then treated with 350 μL of the isolation reagent and converted to complementary (c)DNA following the manufacturer’s protocol. To measure the messenger (m)RNA expression levels, real-time quantitative PCR (ABI StepOne Plus Real Time PCR System), using primers for rat cDNAs, as previously described. The target mRNA levels were compared after normalization to glyceraldehyde-3-phosphate dehydrogenase (GAPDH) mRNA levels as an internal standard that corrected for differences in the efficiency of reverse transcription between samples. The primers for the target mRNAs were collagen type I (Rn00801649 g1), laminin (Rn00564264 m1), fibronectin (Rn00569575 m1), osteocalcin (Rn01455285 g1) and GAPDH (Rn00014552 s1). The thermal cycling conditions were 20 seconds at 95°C for activation of the TaqMan Fast Universal PCR Master Mix, followed by 40 cycles of 1 second at 95°C for denaturing and 20 seconds at 60°C for annealing and extension. The experiments were performed in triplicate.

Western blotting. Secretion of ECM components was evaluated by Western blot analysis. Cell lysates were prepared using mammalian protein extraction reagent (M-PER; Pierce, Rockford, Illinois) containing protease inhibitor cocktails of 1 mM sodium orthovanadate and 25 mM NaF. The protein concentration was measured twice using a BCA Protein Assay Kit (Pierce) with bovine serum albumin as the standard. Cell lysate samples containing 10 μg of protein were separated by SDS-polyacrylamide gel electrophoresis and transferred to a polyvinylidene difluoride membrane (Invitrogen, Thermo Fisher Scientific, Waltham, Massachusetts). The membrane was incubated with antibodies against collagen type I (Rockland Immunochemicals Inc., Pottstown, Pennsylvania), fibronectin (Rockland Immunochemicals), laminin (Santa Cruz Biotechnology, Dallas, Texas) and actin (Merck MILLIPORE, Darmstadt, Germany) diluted in phosphate buffered saline (PBS-T) with 5% non-fat dry milk at 4°C overnight. The membrane was then washed in PBS-T and incubated with appropriate secondary antibodies at room temperature for one hour. The immune complexes were made visible with a chemiluminescent substrate (Western Lightning Plus-ECL; PerkinElmer, Waltham, Massachusetts). The signal intensities of the expressed protein bands were measured using densitometric analysis (ImageJ software; National Institute for Health, Bethesda, Massachusetts) and normalised to the ratio of the band densities in the MEM group, which was arbitrarily defined (by MA) as one unit.

Measurement of proteoglycan (PG) content. PG in the cell matrix in each group was measured. Briefly, cells of each group were cultured in 6-well plates (n = 3) and PG in the cell matrix at day 14 of each group was measured by ELISA (Proteoglycan Detection Kit, Astarte Biologics, Bothell, Washington). This experiment was performed in triplicate.

Subcutaneous transplantation of the composite of β-TCP granules and cultured cells. To assess the in vivo bone formation pattern, the cultured cells from all four groups were scraped off and transplanted together with β-TCP granules (n = 3). Briefly, the β-TCP scaffolds were combined with the cultured cells and then subcutaneously implanted into the backs of the eight rats. The β-TCP scaffolds were harvested after four weeks, fixed in 10% neutral-buffered formalin, decalcified with EDTA solution, and embedded in paraffin. The specimens were cut through the middle and stained with haematoxylin (H) and eosin (E). The experiments were performed in duplicate. Two of the authors (MA and TS) observed the H-E sections to assess whether there was bone tissue. MA knew the sample names, however TS was blinded. The authors got same result.

Osteogenesis at the bone defect site. To assess whether the complete cell sheet (cultured in Dex/AscP medium) could form bone tissue at the bone defect site, another animal study was performed. Briefly, femurs harvested from donor rats were irradiated (60 Gray) to obtain dead bone. Then, a bone defect was created at the shaft of the dead femur, and the defect was filled with β-TCP granules combined with cell sheets. Subsequently, the dead femur with β-TCP granules/cell sheet was subcutaneously implanted into the backs of rats. As a control, dead femur, in which the bone defect was filled with β-TCP granules alone, was transplanted subcutaneously. The femurs were harvested after four weeks, fixed in 10% neutral-buffered formalin, decalcified with EDTA solution, and embedded in paraffin. The specimens were cut through the longitudinal axis and stained with H and E. Two of the authors (MA and TS) observed the H-E sections to assess whether there was bone tissue. MA knew the sample names, however TS was blinded. The authors got same result.

Statistical analysis. This was performed in SPSS (Version 22, IBM, Armonk, New York). The statistical significance of differences between groups was determined using one-way analysis of variance with Tukey post hoc multiple comparisons tests. A p-value of < 0.05 was considered statistically significant.

Results

The appearance of the retrieved cell sheets in each group after scraping are shown in Figure 1. No obvious tears in the sheet structure of the Dex/AscP group were seen by macroscopic observation (Fig. 1a), and those cell sheets were easily removed from the wells. In contrast, the cell sheets in the MEM, AscP, and Dex groups appeared as fragments and incomplete structures. Tears and fragmentation of the sheet structure was found by microscopic observation (Fig. 1b) in the MEM, AscP, and Dex groups. In contrast, the Dex/AscP
The results of the present study clearly indicate that culture in medium with both Dex and AscP created better osteogenic matrix cell sheets than culture with either Dex or AscP alone. These culture conditions allowed for the mechanical retrieval of confluent cells in a single sheet structure.

The real-time PCR assessments of the mRNA expression levels of the ECM components are shown in Figure 3. The expressions of collagen types I and laminin were significantly higher in the Dex/AscP group than in the MEM group (p = 0.001 and 2.7×10⁻⁵, respectively). The expression of osteocalcin was significantly higher in the Dex and Dex/AscP group than in the MEM group (p = 0.003 and 0.001, respectively).

The Western blots for collagen type I, laminin and fibronectin are shown in Figure 4. More collagen type I was found in the Dex/AscP group than in the MEM group, whereas the bands of laminin and fibronectin were not increased.

Discussion

The results of the present study clearly indicate that culture in medium with both Dex and AscP created better osteogenic matrix cell sheets than culture with either Dex or AscP alone. These culture conditions allowed for the mechanical retrieval of the confluent cell layer as a single cell sheet structure that induced bridging bone formation in vivo. The results indicate that culture in Dex/AscP increases cell number and the expression of ECM components such as collagen type I and PG. The biosynthesis and deposition of this ECM by the cultured cells appears to bind the confluent cells into a stable sheet structure. Furthermore, osteoblastic differentiation is induced by the addition of Dex/AscP, which may be advantageous for early bone formation after transplantation.
Culturing bone marrow cells with dexamethasone and ascorbic acid improves osteogenic cell sheet structure

Collagen has recently received considerable attention as a favourable matrix for seeding cells. Collagen is important for many tissue engineering methods, and is widely used for bone repair and new bone formation. Collagen has recently received considerable attention as a favourable matrix for seeding cells. Collagen is important for many tissue engineering methods, and is widely used for bone repair and new bone formation. Collagen has recently received considerable attention as a favourable matrix for seeding cells. Collagen is important for many tissue engineering methods, and is widely used for bone repair and new bone formation.

Real-time polymerase chain reaction messenger RNA expression of (A) collagen type I, (B) laminin, (C) fibronectin and (D) osteocalcin.

Western blots showing that the expression of collagen type I was increased in the Dex/AscP group. The expression ratio of minimal essential medium was defined as 1 unit in the bar figure (upper panel) (Col1, collagen type I; LN, laminin; FN; fibronectin).

Proteoglycan measurement of the bone marrow stromal cells in the groups (*p = 0.002) (MEM, minimal essential medium; AscP, ascorbic acid phosphate; Dex, dexamethasone).
because it is biocompatible, improving MSC survival.  

Different types of collagen are the main components of most fibrous connective tissues, such as tendon, ligament, and skin, as well as cartilage and bone. Collagen type I, which consists of two α1 and one α2 chains, is the most abundant protein in mammals, and a natural ECM that supports most tissues and cell structures. It is also present in bone and teeth and is the main organic component of bone. Laminin, which consists of α-, β-, and γ-chains, is the major protein component of the basement membrane, which forms a protein network foundation for cells, as well as organs. The laminin that is secreted and incorporated into the ECM belongs to the glycoprotein family and is an essential part of the structural scaffolding in most animal and human tissues. In the present study, culture with Dex/AscP increased cell number and the production of ECM components, including collagen type I and PG, resulting in the formation of an osteogenic matrix cell sheet. Unlike in methods that use trypsin/EDTA digestion, because the ECM in an osteogenic matrix cell sheet remains intact, they are suitable for skeletal reconstruction applications such as bone defects and fracture nonunions. Transplantation of an osteogenic matrix cell sheet includes both necessary features of cell transplantation and ECM transplantation, which together, can fill in gaps between bones and artificial bone, as well as to unite artificial bones. PGs are a major component of the animal extracellular matrix and are formed of glycosaminoglycans (GAGs) covalently attached to the core proteins. PG and GAG side-chains form supramolecular aggregates that interconnect the collagenous network in connective tissues, and play a significant role in regulating the mechanical behaviour of the extracellular matrix, particularly in soft tissues.

The standard procedure for osteogenic differentiation of stem cells is treatment of culture cells with Dex, AscP and β-glycerophosphate (β-GP). Langenbach and Handsche13 reported that Dex induces osteogenic differentiation of BMSCs by increasing the expression of mitogen-activated protein kinase phosphatase-1 (MKP-1), which dephosphorylates and activates runt-related transcription factor 2 (Runx2). AscP leads to the increased secretion of collagen type I, which facilitates osteogenic differentiation. Others reported that Dex prevents apoptosis of BMSCs and promotes MSC proliferation. In our method of cell sheet fabrication, we cultured BMSCs with Dex and AscP, therefore enhancing osteogenic potential and ECM (collagen type I and PG) production, and resulting in a
Culturing bone marrow cells with dexamethasone and ascorbic acid improves osteogenic cell sheet structure

Recently, many researchers have reported methods for fabricating cell sheets, e.g., using special culture plates or multifunctional copolymers. Specifically, Yang et al. reported that the use of culture dishes coated with thermosensitive polymers, which have been clinically applied for the treatment of corneal surface dysfunction. Other researchers reported on the mechanical retrieval method of cell sheet fabrication by a scraping technique, which formed bone tissue within a subcutaneous site, consistent with our previous results. However, these other groups created their cell sheet in MEM with Dex, AscP and β-GP. In our pilot study, it was difficult to scrape off the cell sheet when cells were cultured with Dex, AscP and β-GP as calcium deposition disturbed the process of smooth scraping (data not shown).

There are a few limitations to the present study. First, the study was performed on rat BMSCs. Because human BMSCs are used in the clinical application of cell sheets created using our technique, human BMSCs should be used to determine whether culture with Dex/AscP improves the integrity of osteogenic matrix cell sheets. Secondly, although we assessed the expression of the ECM components collagen types I, laminin, fibronectin and PG to show the potential mechanism of cell sheet formation, there are many other ECM components that were not assessed. Thirdly, we cultured the BMSCs in only a single concentration of Dex and AscP. To determine the optimal concentrations of Dex and AscP that create stable cell sheets, a range of treatment doses are necessary. However, no obvious tears were found in the sheet structures cultured with Dex and AscP, indicating that the cell sheets may be useful for tissue engineering methods of bone regeneration. Finally, we did not compare the characteristics of cell sheets created using our method with those of cell sheets created by other techniques, such as by using a thermosensitive polymer. Therefore, other techniques for making cell sheets from BMSCs should be similarly tested. Future studies will address these issues.

In conclusion, the data from the present study indicate that culture with Dex and AscP increased the cell number and expression of the ECM components collagen types I and PG, resulting in a stable cell sheet structure. These culture conditions allowed for the mechanical retrieval of the confluent cell layer as a single cell sheet structure that induced bridging bone formation in vivo. Because of its clear benefit to cell sheet structure, this method may be beneficial when applied to cases where tissue reconstruction is difficult.
References

1. Akahane M, Ueha T, Dohi Y, et al. Secretory osteocalcin as a nondestructive osteogenic marker of tissue-engineered bone. J Orthop Sci 2011;16:622-628.

2. Ohgushi H, Caplan AI. Stem cell technology and bioceramics: from cell to gene engineering. J Biomed Mater Res 1999;48:913-927.

3. Tsuji W, Rubin JP, Marra KG. Adipose-derived stem cells: implications in tissue regeneration. World J Stem Cells 2014;6:312-321.

4. Úsas A, Maciulaitis J, Maciulaitis R, et al. Skeletal muscle-derived stem cells: implications for cell-mediated therapies. Medicina (Kaunas) 2011;47:469-479.

5. Akahane M, Ohgushi H, Yoshikawa T, et al. Osteogenic phenotype expression of allogeneic rat marrow cells in porous hydroxyapatite ceramics. J Bone Miner Res 1999;14:561-568.

6. Watanabe H, Yodoi Y, Yamanaka S, et al. Human autologous cultured expanded bone marrow mesenchymal stem cells provide osteoblastic cell growth and noninvasive cell sheet harvesting. Tissue Eng Part C 2010;26:921-932.

7. Woodbury D, Schwarz EJ, Prockop DJ Black IB. Adult rat and human bone marrow stromal cells differentiate into neurons. J Neurosci Res 2000;61:364-370.

8. Brazelton TR, Rossi FMV, Neshet G. From marrow to brain: expression of neuronal phenotypes in adult mice. Science 2002;295:1775-1779.

9. Ohishi M, Schipani E. Bone marrow mesenchymal stem cells. J Cell Biochem 2010;109:277-282.

10. Salem HK, Thiemermann C. Stromal cell sheets: current understanding and clinical status. Stem Cells 2010;28:585-596.

11. Ueha T, Akahane M, Shimizu T, et al. Utility of tricalcium phosphate and osteogenic matrix cell sheet constructs for bone defect reconstruction. W J Stem Cells 2015;5:673-682.

12. Uchihara Y, Akahane M, Shimizu T, et al. Osteogenic Matrix Cell Sheets Facilitate Osteogenesis in Irradiated Rat Bone. BioMed Res Int 2015;629:168.

13. Matsushima A, Kotobuki N, Todokoro M, et al. In vivo osteogenic capability of human mesenchymal cells cultured on hydroxyapatite and on β-tricalcium phosphate. Avit Organs 2009;33:474-481.

14. Kawate K, Yajima H, Ohgushi H, et al. Tissue-engineered approach for the treatment of steroid-induced osteonecrosis of the femoral head: transplantation of autologous mesenchymal stem cells cultured with beta-tricalcium phosphate ceramics and free vascularized fibula. Avit Organs 2008;30:969-962.

15. Ter Brugge PJ, Jansen JA. In vitro osteogenic differentiation of rat bone marrow cells subcultured with and without dexamethasone. Tissue Eng 2002;9:821-331.

16. Nakamura A, Dohi Y, Akahane M, et al. Osteoclast secretion as an early marker of in vitro osteogenic differentiation of rat mesenchymal stem cells. Tissue Eng Part C Methods 2009;15:169-180.

17. Bianco P, Robey PG. Stem cells in tissue engineering. Nature 2001;414:118-121.

18. Petitie H, Viateau V, Bensaid W, et al. Tissue-engineered bone regeneration. Nat Biotechnol 2000;18:959-963.

19. Dong J, Kojima H, Uemura T, et al. In vivo evaluation of a novel porous hydroxyapatite to sustain osteogenesis of transplanted bone marrow-derived osteoblastic cells. J Biomed Mater Res 2001;57:208-216.

20. Shimizu T, Sekine H, Yamato M, et al. Cell sheet-based myocardial tissue engineering: new hope for damaged heart rescue. Curr Pharm Des 2009;15:2807-2814.

21. Kanai N, Yamato M, Okano T. Cell sheets engineering for esophageal regenerative medicine. Ann Transl Med 2014;2:28.

22. Akahane M, Nakamura A, Ohgushi H, et al. Osteogenic matrix sheet-cell transplantation using osteoblastic cell sheet resulted in bone formation without scaffold at an ectopic site. J Tissue Eng Regen Med 2008;2:196-201.

23. Ma D, Ren L, Liu Y, et al. Engineering scaffold-free bone tissue using bone marrow stromal cell sheets. J Orthop Res 2010;28:697-702.

24. Akahane M, Ueha T, Shimizu T, et al. Cell sheet injection as a technique of osteogenic supply. Int J Stem Cells 2010;3:138-143.

25. Nakamura A, Akahane M, Shigematsu H, et al. Cell sheet transplantation of cultured mesenchymal stem cells enhances bone formation in a rat nonunion model. Bone 2010;46:418-424.

26. Inagaki Y, Uematsu K, Akahane M, et al. Osteogenic matrix cell sheet transplantation enhances early tendon graft to bone tunnel healing in rabbits. Bio Med Res Int 2013;942:192.

27. Akahane M, Shigematsu H, Todokoro M, et al. Scaffold-free cell sheet injection results in bone formation. J Tissue Eng Regen Med 2010;4:404-411.

28. Shimizu T, Akahane M, Ueha T, et al. Osteogenesis of cryopreserved osteogenic matrix cell sheets. Cryobiology 2013;66:326-332.

29. Toma Y, Dohi Y, Ohgushi H, et al. Osteogenic activity of bone marrow-derived mesenchymal stem cells (BMSCs) seeded on irradiated allogeneic bone. J Tissue Eng Regen Med 2012;6:96-102.

30. Kim HJ, Kim U-J, Yunjak-Novakovic G, et al. Influence of macroporous protein scaffolds on bone tissue engineering from bone marrow stem cells. Biomaterials 2005;26:4442-4452.

31. Oh S-A, Lee H-Y, Lee JH, et al. Collagen three-dimensional hydrogel matrix carrying basic fibroblast growth factor for the cultivation of mesenchymal stem cells and osteogenic differentiation. Tissue Eng Part A 2012;18:1077-1100.

32. Bertassoni LE, Swain MV. The contribution of proteoglycans to the mechanical behavior of mineralized tissues. J Mech Behav Biomater 2014;36:91-104.

33. Langenbach F, Handschef J. Effects of dexamethasone, ascorbic acid and β-glycerophosphate on the osteogenic differentiation of stem cells in vitro. Stem Cell Res Ther 2013;4:117.

34. Song H, Caplan AI, Dennis JE. Dexamethasone inhibition of confluence-induced apoptosis in human mesenchymal stem cells. J Orthop Res 2009;27:216-221.

35. Wang H, Pang B, Li Y, et al. Dexamethasone has variable effects on mesenchymal stromal cells. Cytotherapy 2012;14:423-430.

36. Nishida K, Yamato M, Hayashida Y, et al. Corneal reconstruction with tissue-engineered cell sheets composed of autologous oral mucosal epithelium. N Engl J Med 2004;351:1187-1196.

37. Kim YS, Lim JY, Donahue HJ, et al. Thermoresponsive terpolymeric films applicable for osteoblastic cell growth and noninvasive cell sheet harvesting. Tissue Eng Regen Med 2005;11:30-40.

38. Yang J, Yamato M, Nishida K, et al. Corneal epithelial stem cell delivery using cell sheet engineering: not lost in transplantation. J Drug Target 2006;14:471-482.

Funding Statement

M. Akahane reports funding received from the Translational Research Network Program from the Japan Agency for Medical Research and Development (AMED) which is related to this article.

Author Contribution

M. Akahane: Designed the study, Prepared the manuscript.
T. Shimizu: Assisted in the study processes and preparations.
T. Onishi: Nara medical University Faculty of medicine
Y. Uchihara: Assisted in the study processes and preparations.
T. Imamura: Assisted in manuscript preparation, Critically reviewed the manuscript.
Y. Uchihara: Assisted in manuscript preparation, Critically reviewed the manuscript.

ICMJE conflict of interest

None declared.

© 2016 Akahane et al. This is an open-access article distributed under the terms of the Creative Commons Attribution license (CC-BY-NC), which permits unrestricted use, distribution, and reproduction in any medium, but not for commercial gain, provided the original author and source are credited.