Cell Shapes and Actin Alignment of Mesenchymal Stem Cells on Different Elastic Matrices with the Same Surface Topography

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Abstract

Mesenchymal stem cells (MSCs) can differentiate into multilineage cells. Thus, many efforts have been focused on finding effective ways to selectively differentiate the pluripotent cells into defined lineage-committed progenitor cells for their suitable use in regenerative medicine. Some studies showed that the lineage of MSCs was influenced by the surface topography where the cells were attached, while other studies showed that its lineage was decided by the rigidity of the matrix where the cells interfaced. Our study showed that both the surface topography and rigidity of the matrix simultaneously affected the shapes of MSCs through the alignment and organization of actin cytoskeletons when MSCs were grown on either soft (1 M Pa) or hard (2 – 5 GPa) polymers with groove patterns in various dimensions ranging from 500 nm to 50 μm. Cells were mostly aligned and elongated along all the patterns irrespective of the rigidity of the substrate. However, the degree of the elongation was affected by both the matrix rigidity and dimension of the patterns. This is further supported by the results showing that the orientation of alignment of actin cytoskeletons and their organization were affected by both the rigidity and the topography of the substrate.

Introduction

Mesenchymal stem cells (MSCs), which are capable of self-renewal and differentiation into various tissues, have attracted attention for use in regenerative medicine [1-4]. Many studies have demonstrated that MSCs can differentiate into osteoblasts, chondrocytes, adipocytes, muscle cells and even neuronal cells, and show respective functions in vivo after transplantation [1-6]. One of the approaches to utilize MSCs in regenerative medicine is transplantation of fully differentiated cells which are differentiated from MSCs in vitro [5,6]. The traditional method to induce the specific differentiation of stem cells in vitro is based on the treatment of chemicals such as several growth factor mixtures that lead stem cells to a specific lineage [5-8]. However, recent studies suggest that mechanical factors, such as rigidity or surface topography of the cell culture substrates, can also affect the stem cell fates [9-13]. For example, the surface topography in groove and ridge patterns, which have micron-scale widths and nanoscale depths, induced osteogenesis of the MSCs [12], whereas nanotopography in groove and ridge patterns (300 or 500 nm) induced neuronal differentiation of MSCs [13]. In addition, a recent report showed that MSCs can differentiate effectively on a substrate with disordered asymmetric post patterns, in contrast to the substrate with ordered post patterns, and demonstrated that symmetry or regularity of the surface topography is also important for the decision of the stem cell fates10. Another study demonstrated that soft matrices (0.1 – 1 kPa) induced neurogenic progenitor cells, while stiffer matrices (8 – 17 KPa) and rigid matrices (25 – 40 KPa) induced myogenic and osteogenic progenitor cells, respectively [11].

However, it has not been investigated whether cells can distinguish and respond to each mechanical cue when diverse combinations of various mechanical stimuli are given simultaneously. In addition, it has not been studied whether a specific mechanical cue can predominantly affect cellular behaviors among various mechanical cues. For example, even though it was known that the rigid substrate tends to direct osteogenesis of MSCs [11] and nanogroove topography induces neural differentiation of MSCs [13], there is no information about the differences between the cellular responses on the nanogroove patterns made of rigid materials and soft materials. However, this kind of research provides useful information to develop and optimize the cell culture substrates for specific stem cell differentiation.

Furthermore, it is also necessary to investigate the alteration of cytoskeletal systems while diverse combinations of various mechanical stimuli are given simultaneously to cells. Recent studies showed that cells respond to mechanical stimuli through the cytoskeletal system, which discerns the physical changes of microenvironments around the cells and passes the information on to downstream of the signal pathway to regulate cellular shapes and differentiation14-17. These results were also supported by several recent studies that showed that disrupting the cytoskeletal systems inhibited mechanical-cue-mediated morphology change and fate decision in stem cells11, 17. However, it is still elusive how cytoskeletal systems react to the diverse mechanical cues at the same time, and how they integrate the several mechanical signals.

Thus, we were motivated to investigate whether two different mechanical cues can simultaneously influence the cytoskeletons and cellular shapes. In this study, we examined the cellular shapes and the structures of actin cytoskeletons in MSCs on different elastic matrices with the same surface topography in various dimensions.

Materials & Methods

Cell Culture

Human MSCs were provided from PromoCell (Heidelberg, Germany) and maintained in low-glucose Dulbecco’s Modified Eagle’s Medium (Gibco, Germany) with 10% fetal bovine serum (FBS) in an incubator with a humidified atmosphere of 5% CO2 at 37°C. Cells were seeded on polymer surfaces in 24-well plates and grown to confluence. After reaching 80% confluence, cells were passaged and used for experiments.

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Medium (DMEM with 1 g/L glucose) containing 10% fetal bovine serum (FBS), 100 units/ml penicillin, and 100 µg/ml streptomycin [2,3]. The cells were incubated at 37°C and 5% fully humidified CO2. Cells in passage 6 were used for all experiments in this study.

Substrate preparation

Substrates for cell culture were fabricated on two different materials, polydimethylsiloxane (PDMS) and SU-8. For patterning structures on PDMS substrates, PDMS were cast on a master mold which was photo-lithographically fabricated in SU-8 on a silicon wafer, or master patterns in etched silicon surfaces after e-beam lithography. Micropatterned SU-8 substrates were fabricated on tissue culture plates (TCP) by soft lithography18 using a PDMS replica mold. In detail, PDMS replica molds were placed on SU-8-coated TCP surfaces and exposed to UV-rays for 10 min., and the PDMS molds were removed and cured for 20 min. For nanoscale patterns, since soft lithography using PDMS molds cannot guarantee accurate features, a hard polyurethane acrylated polymer mold coated with an anti-adhesion PDMS layer19 was used for the nanopattern transfer process. The rest of the process is the same as the microscale SU-8 molding described above. All the substrates were prepared for cell culture by coating with 10 µg/ml fibronectin for 2 h in a 5% CO2 cell culture incubator. PDMS substrates were treated with oxygen plasma before the coating.

Cell staining

After the cells were cultured on various substrates, they were fixed in 3.7 % formaldehyde for 1 h, rinsed in phosphate buffered saline (PBS), and permeabilized by treating with 0.1 % tritonX-100 for 15 min. Then, actin cytoskeletons were stained with rhodamine-conjugated phalloidin, and the nuclei were stained with 4’-6-diamidino-2-phenylindole (DAPI).

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for this phenomenon, further studies are needed for the identification of the threshold values of the stiffness ranges sensed by MSCs on different elastic matrices.

When MSCs were cultured on the patterned substrates with the dimensions of 0.5, 1, 20, and 50 µm (groove and spacing width, Figures 2(d-k)), the cells oriented along the direction of patterns such that the longest axes of the cells were aligned in parallel to the grooves and ridges. In addition, the cells on the PDMS groove and ridge patterns were elongated more than those on SU8 substrates with the same pattern of structures. On the PDMS substrate with 1-µm groove patterns, it was commonly observed that cells were extremely elongated more than 1 mm. In addition, some parts of the cells on that substrate showed morphological similarity to neuronal cells composed of a single cell body and highly branched neuritis, while this was not observed on the SU8 1-µm groove patterns. This result indicates that cells can sense the surface topography and matrix elasticity simultaneously. Furthermore, it is very interesting that the differences in the cell elongation and shape change on the PDMS and the SU8 substrates were observed only on the patterned substrates (especially on the 1-µm groove patterns), while no difference was observed on flat surfaces. Through this result, we propose that there are some interactions between sensing the matrix elasticity and sensing the surface topography in the cells.

To corroborate the results (Figure 2), high-magnification images for fluorescent microscopic imaging. Actin cytoskeletons were stained with rhodamine-conjugated phalloidin, and the nuclei were counterstained with DAPI. In the cells on the SU8 substrates with 0.5-µm groove patterns, actins were strongly expressed only at the edge of the cells (Figure 3a). On the SU8 substrates with the 1-µm groove patterns, actin fibers aligned in parallel to the patterns were clearly observed in both the middle and the edge of the cells (Figure 3b). In the case of the 20-µm groove made of SU-8, even though the cell bodies were aligned along the patterns, most of the actin fibers inside the cells were biased by about 60 degrees (Figure 3c). However, actins on the 50-µm patterns were not organized in a specific orientation; these actins were biased and formed meshwork. On the patterned PDMS substrates, strong actin fibers were not observed as much as on the patterned SU8 surfaces (Figures 3e-g). It was observed that the filament-like actins were aligned along the pattern, and the actins formed meshwork on all the patterned PDMS substrates.

Subsequently, different cell shape parameters, such as the degree of elongation, distinct form, and alignment of actin cytoskeleton, were observed more significantly on substrates with different stiffnesses compared to flat substrates. Unfortunately, sufficient data were not accumulated to explain these phenomena. Further investigation is necessary to understand the mechanism of the substrate stiffness effect on the dynamic rearrangement of cytoskeletons in MSCs.

We have demonstrated that the matrix elasticity and the surface topography simultaneously affect cell morphology and cytoskeleton rearrangement. Furthermore, our results reveal that the matrix elasticity (1 MPa and 2 GPa) has a greater effect on changes of MSC shapes and cytoskeletons with surface topography than without surface patterns. Therefore, we propose that there are some interconnections between sensing the matrix elasticity and sensing the surface topography in the cells. In conclusion, the combination of the matrix elasticity and the surface topography are considered to apply matrix mechanical cues for specific differentiation of stem cells in regenerative medicine.

**Conclusion**

The authors have declared that no competing interests exist.

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**Actin-Alignment of MSCs grown on various substrates**

To corroborate the results (Figure 2), high-magnification images (×800) of actin cytoskeletons were taken using a confocal laser scanning microscope (CLSM). Figure 3 shows some representative images of the cells cultured on the same substrates previously used for this study. The scale bars represent 20 µm and the direction of the groove patterns are represented by double arrow headed lines above the scale bars.

**Figure 3: Confocal laser microscopic images of actin cytoskeletons of MSCs on different substrates. Actin cytoskeleton of the cells on (a)–(d) SU8 substrates and (e)–(h) PDMS substrates. Dimensions of groove patterns in each substrate are following; (a) 500 nm, (b) and (c) 1 µm, (d) and (e) 20 µm, and (f) and (g) 50 µm. The scale bars represent 20 µm and the direction of the groove patterns are represented by double arrow headed lines beside of the cells.**

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Author Contributions

SH Kim performed the experiment and data analysis, G Lee and S Park did analysis and wrote the manuscript. The funding for the research was acquired by S Park.

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