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Engineering micropatterned surfaces to modulate the function of vascular stem cells

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ABSTRACT

Multipotent vascular stem cells have been implicated in vascular disease and in tissue remodeling post therapeutic intervention. Hyper-proliferation and calcified extracellular matrix deposition of VSC cause blood vessel narrowing and plaque hardening thereby increasing the risk of myocardial infarct. In this study, to optimize the surface design of vascular implants, we determined whether micropatterned polymer surfaces can modulate VSC differentiation and calcified matrix deposition. Undifferentiated rat VSC were cultured on microgrooved surfaces of varied groove widths, and on micropost surfaces. 10 μm microgrooved surfaces elongated VSC and decreased cell proliferation. However, microgrooved surfaces did not attenuate calcified extracellular matrix deposition by VSC cultured in osteogenic media conditions. In contrast, VSC cultured on micropost surfaces assumed a dendritic morphology, were significantly less proliferative, and deposited minimal calcified extracellular matrix. These results have significant implications for optimizing the design of cardiovascular implant surfaces.

1. Introduction

Cardiovascular disease is the leading cause of death in the United States. Coronary Artery Disease (CAD), a narrowing of the coronary arteries caused by plaque and inflammation, accounts for 16% of deaths and $108.9 billion in healthcare costs annually in the US. Plaque hardening, caused by osteoblastic deposition of calcified matrix, significantly increases the risk of plaque rupture, inflammation and, ultimately, myocardial infarct [1]. Furthermore, balloon angioplasty and stenting treatments to restore arterial patency cause vessel wall injury and can result in arterial narrowing from tissue restenosis. Even in the era of drug eluting stents, tissue restenosis remains a major complication of CAD treatment [2]. Second generation drug eluting stents, although significantly more effective than bare metal or first general drug eluting stents, still suffer from a restenosis rate of 12.2% [3].

The traditional model of vascular disease held that de-differentiated smooth muscle cells were responsible for intimal thickening in atherosclerosis, and for tissue restenosis following angioplasty and stenting. Furthermore, smooth muscle cells were implicated in calcified matrix formation in diseased arteries [4]. However, more recently, a population of multipotent vascular stem cells (VSCs) was found resident in the blood vessel wall [5]. In response to vascular injury, VSCs were shown to differentiate and contribute to vascular remodeling, to structural changes involving media thickening, and to reduced vessel lumen diameter. The VSC were also shown to be capable of osteogenic differentiation *in vitro*. Thus, it may be likely that VSC not only are responsible for vessel wall thickening but also for tissue calcification. Given the proliferative and differentiation potential of VSC, there is a critical clinical need to develop therapeutics that specifically target VSC function. Modulation of VSC behavior may lead to drastically improved clinical outcomes by preventing or limiting arterial narrowing and hardening.

Micro- and nano-topographic patterned biomaterial surfaces have been studied for biophysical regulation of cell functions such as orientation, proliferation, differentiation and tissue formation [6–13]. In particular, microgrooved surfaces have been shown to reduce vascular smooth muscle cell proliferation. Additionally, micropost substrates with varying micropost stiffness modulated human mesenchymal stem cell morphology, proliferation and differentiation [14], and directed endothelial cell migration [15]. In this study, we assessed the effect of microgrooves and microposts on rat vascular stem cell proliferation, osteogenic differentiation, and calcified matrix deposition.

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2. Materials and methods

2.1. VSC culture

Primary rat VSCs were obtained by using aorta tissue explant culture as previously described [5]. VSC cultures were maintained with Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin (PS) (complete medium) in an incubator at 37 °C. Cell cultures were maintained in a humidified 95% air–5% CO_2 incubator at 37 °C.

2.2. Micropatterned PDMS membranes

Polydimethylsiloxane (PDMS) membranes with micropatterned grooves and micropatterned posts were fabricated as previously described [8,13,15]. The PDMS was spun-coated onto photolithography patterned silicon wafers with appropriate microgrooves of various widths (10, 50, and 100 μm) and 3 μm depth, or with uniformly distributed cylindrical micropits of 7 μm depth, 3 μm diameter and 3 μm spacing. The PDMS was cured by baking on a hotplate, and the PDMS membrane was removed and cut to appropriate sizes. As an experimental control, unpatterned PDMS membranes were spun-coated and cured on a silicon wafer with a smooth surface. PDMS membranes were sterilized and coated with Cellstart (Life Technologies, Grand Island, NY) prior to rat VSC culture.

2.3. Immunofluorescence staining of VSC mutlipotency markers

Rat VSC were characterized for multipotency markers as previously described [5]. Rat VSC were immunofluorescently stained for Sox10 (R&D Systems, Minneapolis, MN) and Neurofilament-M (Abcam, Cambridge, MA), and counter-stained with DAPI (Sigma–Aldrich, St. Louis, MO). The samples were imaged with a Zeiss fluorescent microscope (Zeiss).

2.4. Morphological assessment of VSC on micropatterned membranes

Rat VSC morphology on micropatterned membranes was analyzed with fluorescent staining of the actin cytoskeleton. Rat VSC were cultured on the micropatterned and control PDMS membranes for 24 h. After 24-h culture the cells were fixed, permeabilized and fluorescently stained with Alexa Fluor 546 phalloidin (Life Technologies, Grand Island, NY) and DAPI. The fluorescently stained samples were imaged with the Zeiss fluorescent microscope.

2.5. VSC proliferation on micropatterned membranes

The effect of micropatterned surfaces on rat VSC proliferation was measured using the Click-It 5-Ethyl-2′-deoxyuridine (EdU) proliferation assay (Life Technologies, Grand Island, NY). Rat VSC were cultured on micropatterned and control PDMS membranes for 48 h in complete medium.

After 48 h culture in complete medium, the rat VSC were pulsed with EdU for 1 h. The cells were then fixed, permeabilized, and fluorescently stained. For EdU detection cells were incubated with Alexa Fluor azide 488, and then counter-stained with DAPI. The stained samples were imaged with a Zeiss Fluorescence microscope. The images of EdU and DAPI staining were used to count proliferative cells and total number of cells using ImageJ software. Proliferation rate for each surface treatment was determined as the percentage of cells incorporated with EdU.

2.6. Osteogenic differentiation of VSC on micropatterned membranes

Rat VSC were cultured on micropatterned and control PDMS membranes in complete medium until 70% confluence. The cultures were then switched to osteogenic induction media and cultured for 4 weeks to induce osteogenic differentiation and calcified matrix deposition as described previously [5].

Osteogenic differentiation of and calcified matrix deposition by rat VSC on the micropatterned surfaces was assessed with alizarin red staining. Alizarin red complexes with calcium and produces a bright red/orange stain that can be imaged with light microscopy. Alizarin red solution was prepared by mixing 1 g of Alizarin Red S in 50 mL distilled water and adjusted to pH 4.3 using 10% ammonium hydroxide. The micropatterned and control samples were washed with PBS and then incubated with Alizarin Red for 20 min. After staining, the cells were washed with distilled water and then imaged with a Zeiss light microscope.

2.7. Statistical analysis

One-way ANOVA with post-hoc Tukey's multiple comparison (95% confidence interval) was used to determine significant differences in proliferation rates between surface treatments.

3. Results and discussion

3.1. Rat VSC expressed pluripotency markers

We first examined the effects of parallel microgrooves on rat VSC morphology, proliferation and osteogenic differentiation. All experiments were conducted with low passage, undifferentiated rat aortic VSC. Rat VSC multipotency was verified by immunofluorescent staining of Sox10 and NFM. As shown in Fig. 1, all cells expressed Sox10 and NFM. Diffuse staining of NFM was observed in the cell cytoplasm (Fig. 1B). Sox10, a transcriptional factor, was primarily localized in the nucleus (Fig. 1C and D). The Sox10 and NFM staining profile, consistent with previous results [5], suggested that VSCs in culture were in an undifferentiated state.

3.2. 10 μm microgrooved surfaces constricted and oriented VSC

Microgrooved PDMS membranes composed of parallel alternating ridges and grooves were produced with groove and ridge widths of 10, 50 or 100 μm (Fig. 2). To assess the effects of parallel microgrooves on VSC morphology, the cells were cultured on 10, 50 and 100 μm microgrooved PDMS membranes (Fig. 2). Actin cytoskeletal staining with fluorescent phalloidin showed that cell morphology was guided by the topography of the micropatterned surfaces. VSC cultured on control surfaces had flat, spread morphology with no particular cytoskeletal or nuclear orientation. On the microgrooves, VSCs generally exhibited an elongated morphology and cytoskeletal alignment parallel to microgroove orientation. On the 10 μm width microgrooves, the cellular actin cytoskeleton and membrane protrusions were aligned along the pattern direction (Fig. 2B). On the wider patterned surfaces (50 and 100 μm), the cellular actin cytoskeleton and membrane protrusions showed a partial alignment with the microgrooves, resulting in less cell elongation than that on narrower microgrooves (Fig. 2C and D). Overall cell size, however, was similar on all cell surfaces thereby indicating that the cells spanned across channels on the smaller widths.

The results of these morphological studies suggest that the smaller 10 μm microgrooves were more effective in orienting VSC actin cytoskeleton and membrane protrusion parallel to the