MPC polymer regulates fibrous tissue formation by modulating cell adhesion to the biomaterial surface

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The aim of this study was to analyze the effects of 2-methacryloyloxyethyl phosphorylcholine (MPC) polymer on fibrous tissue formation and cell adhesion plaque (CAP)-forming reactions. Silastic elastomer (SE) plates coated (experimental group) and uncoated (control group) with MPC polymer were prepared for in vivo and in vitro experiments. For the in vivo animal experiments, SE plates were implanted subcutaneously in the rat dorsal region. At 4, 8, and 12 weeks, thicknesses of the fibrous tissue capsules in the experimental group were lower than in the control group. Likewise, the amount of collagen in the experimental group was lower than that of the control group. For the in vitro cell culture experiments, KMST-6 fibroblast cells in the experimental group demonstrated enhanced cell migration, accompanied with a weaker expression of vinculin and a larger amount of filopodia. Furthermore, weaker expressions of paxillin, talin, and ROCK1, but stronger expression of cofilin, were observed in the experimental group. Taken together, these results suggested that MPC polymer regulated fibrous tissue formation by modulating cell adhesion through changes in local CAPs and downstream signaling.

Keywords: 2-methacryloyloxyethyl phosphorylcholine polymer, Fibrous tissue, Cell adhesion plaque

INTRODUCTION

For the treatment of diseases such as cancer, trauma and myocardial infarction, numerous biomaterials and artificial devices have been used effectively as treatment methods. Despite the laudable treatment success achieved with these biomaterials, the downside is that biomaterials implanted into the body frequently induce inflammation and immune responses, often leading to fibrous scar formation. Although these reactions are a result of normal histological repair responses to foreign bodies, active scar formation may result in unfavorable outcomes, such as breast collapse after prosthetic implantation and stiff joints after artificial joint replacement. Against this backdrop of undesirable outcomes, an important consideration prior to and pertaining to the use of biomaterials and artificial devices is the inhibition of fibrous tissue formation.

When a biomaterial is implanted into a body, it induces a complex sequence of biological reactions in the local tissue, including cell contact, recognition, and signal transduction. The first event in vivo is cell migration to the biomaterial, and hence cell contact with the biomaterial. In light of this cellular phenomenon, the material surface is an important factor in biocompatibility estimation since it affects cell reactions and signal transduction. To the end of fabricating blood-compatible polymer surfaces, a novel coating material based on natural phospholipid cell membranes has been proposed.

2-methacryloyloxyethyl phosphorylcholine (MPC), which has a phospholipid polar group that mimics a biomembrane, was synthesized as a novel coating material by Ishihara et al. Numerous studies have shown that MPC is suitable for use as a biomaterial due to benefits such as suppressing protein adsorption, platelet adhesion, and denaturation of cells and proteins. However, MPC has several drawbacks, amongst which are poor mechanical properties caused by the introduction of water-soluble moieties, and in the case of coatings, weak bonding to the substrate leading to delamination. To improve its mechanical properties and reduce the propensity to delamination, MPC may be copolymerized with other methacrylate monomers, such as n-butyl methacrylate (BMA), n-hexyl methacrylate, and n-dodecyl methacrylate (DMA), through atom transfer radical polymerization (ATRP) or conventional free radical polymerization initiated by different methods.

Having circumvented the mechanical weaknesses through copolymerization, MPC polymers have since become attractive candidates for artificial organs and have shown potential in a wide range of medical device applications. For example, Kyomoto et al. reported its use in artificial hip joints, enhancing lubrication to reduce wear and prosthesis loosening. Huang et al. used it in soft contact lenses to prevent protein adsorption. The list goes on with MPC polymer being applied to myriad medical devices including...
biosensors, drug carriers, vascular stents, and urological devices.

Owing to its ability to mimic the biomembrane, MPC polymer might possess the capacity to regulate unwanted fibrous tissue formation. To date, the evaluation of MPC polymer as a potential coating material for biomaterials has been limited to the analyses of its chemical properties and blood compatibility. This meant that the biocompatibility of MPC polymer, particularly its ability to inhibit fibrous tissue formation and cell adhesion, has not been conclusively demonstrated through in vivo and in vitro experiments. Nonetheless, it has been shown that MPC polymer influenced cell reactions. Konno et al. reported that cell mobility on MPC polymer surface was affected15 by its less adhesive surface, leading to accelerated cell aggregation and differentiation. Incidentally, cell adhesion plaque (CAP) plays an essential role in fibrous tissue formation and cell mobility. For this reason, we hypothesized that the MPC biomembrane could regulate the downstream signals of CAP to suppress protein adsorption and alter cell viability, thereby inhibiting fibrous tissue formation.

Silastic elastomers (SE) are very commonly used in maxillofacial prostheses due to their superior physical properties. However, they are not without shortcomings — amongst which are the tendency to induce fibrous tissue formation, absorb facial oils which can adversely impact prostheses longevity, and to provide a potential site for bacterial colonization. Nonetheless, with improved biocompatibility bestowed upon it by means of MPC polymer coating, silastic elastomers (SE) are well poised to be used in a wider range of medical applications. For this reason, silastic elastomer (SE) was selected in this study as the substrate material to be coated with MPC polymer. Our two-pronged experimental strategy comprised a subcutaneous implant in vivo model and in vitro cell culture to observe capsule scar formation and the regulation of downstream CAP signals, with a view to leveraging on these findings for new medical applications.

MATERIALS AND METHODS

Preparation of silastic elastomer coated with MPC polymer

This study was carried out in compliance with the guidelines of the International Organization of Standardization (ISO) 10996-3 (Biological evaluation of medical devices). MPC polymer (MPC-co-BMA) was prepared as described by Ishihara et al. (Ai Bio-Chips, Tokyo, Japan) (Fig. 1). Surface coating was carried out according to the method reported by Sibarani et al.21 Plates used for MPC polymer coating were fabricated from Silastic® MDX4-4210 BioMedical Grade Elastomer (Dow Corning Chemical Industries, Osaka, Japan). Curing agent and base elastomer (1:10) were thoroughly mixed and exposed to a vacuum of 710 mmHg for 30 minutes to remove trapped air. Then, in accordance with ISO-10996-3 recommendation

silastic elastomer (SE) was placed into a master mold (2×5×20 mm) and cured for 5 hours at 65°C. SE plates were cleaned by sonication in acetone (Wako Pure Chemical Industries Osaka, Japan) and alcohol. For MPC polymer coating, each SE plate was dipped in an EtOH solution containing 0.3 wt% MPC polymer. Solvent was slowly evaporated under an EtOH vapor atmosphere at room temperature for 24 hours, and SE plates were dried in vacuo overnight.

Surface analysis of SE plates coated with MPC polymer was done using X-ray photoelectron spectroscopy (XPS; AXIS His-165 Analytical, Shimadzu Corp., Kyoto, Japan). The take-off angle of the photoelectrons was 90°. The chemical composition of the SE plates was determined using an energy dispersive X-ray analyzer (EDX; JED-2300, JEOL, Japan) operated at an accelerating voltage of 15 kV and with a filament current of 60 µA.

Subcutaneous implantation in rats

All animal experiments were performed in accordance with the Guide for the Care and Use of Laboratory Animals (Tohoku University), 3rd edition. Twenty-four 10-week-old male SD rats were used in all the experiments. Rats were anesthetized by an intraperitoneal injection of sodium pentobarbital (30 mg/kg) (Wako Pure Chemical Industries, Osaka, Japan), and target regions were cleaned with 70% ethanol. Sagittal incisions of 2 cm were made 0.5 mm to the right and left of the dorsal centreline, and then subcutaneous pockets were created by blunt dissection in the thoracic region. SE plates coated and uncoated with MPC polymer (2×5×20 mm) were implanted subcutaneously in the left and right sides respectively. Skin wounds were sutured tightly using Polysorb suture No. 3 (Auto Suture, Norwalk, CT, USA).

Histological evaluation of fibrous tissue formation

Rats were anesthetized and sacrificed for histological evaluation at 2, 4, 8, and 12 weeks after surgery (six rats at each time point). SE plates with surrounding tissues were resected. The SE plates were examined using EDX, while the surrounding tissues were fixed.

![Fig. 1 Chemical formula of poly(MPC-co-BMA) (PMB, \textit{m}: 0.3, \textit{n}: 0.7).](image-url)
with 10% neutral buffered formaldehyde, followed by embedding in paraffin. Sliced sections (3 µm) were routinely stained with hematoxylin and eosin (HE) to evaluate tissue reactions such as inflammation and scar formation around the plates. Thickness of fibrous tissue capsule was measured with a microscope (BX51, Olympus, Tokyo, Japan), as reported previously.\textsuperscript{22}

Collagen staining and measurement
K61 Kit (Wako, Osaka, Japan) was used to stain the collagen in samples of surrounding tissues.\textsuperscript{21} Histological sections (6 µm) were deparaffinized and stained with a saturated solution of picric acid in distilled water containing 0.01% Fast green FCF. Sections were kept out of the light and incubated at room temperature for 15 minutes. After which, they were stained with a saturated solution of picric acid in distilled water containing 0.04% Fast green and 0.1% Sirius red F3B. After incubation in the dark at room temperature for 30 minutes, the stained histological sections were observed under a polarizing microscope (DM500, Leica, Wetzlar, Germany).

To measure the collagen content, 0.1% NaOH in absolute methanol (1:1, v/v, 200 µL) was added to the glass slide. Added NaOH was retrieved and transferred to a test tube, and the eluted color was read in a spectrophotometer at 530 and 605 nm (GloMax-Multi Detection System, Promega, Madison, Wisconsin, USA). The amounts of collagen and non-collagenous protein were calculated using the following formula:

\[
\text{Collagen (mg)} = \frac{(OD_{530} - 0.254 \times OD_{605})}{40.8}
\]
\[
\text{Non-collagenous protein (mg)} = \frac{OD_{605}}{2.04}
\]

Cell cultures on plate surfaces
SE plates coated and uncoated with MPC polymer were placed in 6-well tissue culture plates (Thermo Fisher Scientific, Rochester, NY, USA). Human fibroblast cells (KMST-6, obtained from the Cell Bank of the Old and Aging Medical Institute, Tohoku University, Sendai, Japan) were added to each well at a concentration of \(2 \times 10^5\) cells/well in a cell culture medium (Dulbecco Eagle’s MEM; Nissui Pharmaceutical, Tokyo, Japan) containing 10% fetal bovine serum (FBS; GIBCO BRL, Invitrogen Corp., Grand island, NY, USA). KMST-6 cells were maintained in a culture medium (α-MEM, GIBCO BRL) containing 10% fetal bovine serum at 37°C in a humidified atmosphere of air containing 5% CO\(_2\). Cells were observed daily by phase-contrast microscopy (DM500, Leica, Wetzlar).

**MTT assay**
Cell viability was evaluated using MTT assay.\textsuperscript{23} Briefly, coated and uncoated samples were set at the flat-bottom 96-well plates, and KMST-6 cells were seeded on it (10^4/well) with 100 µL of medium. After 24-hour incubation for attachment, the medium was replaced with a fresh serum-free medium. Then, each day from day 1 to day 6, six wells were added with 10 µL of MTT (5 mg/mL) (Wako Pure Chemical Industries, Osaka, Japan) and incubated for 4 hours. 100 µL of DMSO was also added to each well and mixed. Absorbance at 492 nm was measured using the spectrophotometer (GloMax-Multi Detection System, Promega, Madison, Wisconsin, USA).

**Adhesion plaque staining by immunofluorescence histochemistry**
KMST-6 cells were collected at day 3 after culturing on SE plates coated and uncoated with MPC polymer. SE plates were washed in phosphate buffered saline (PBS) and fixed with 4% paraformaldehyde. SE plates were then treated with 0.2% Triton X-100. After washing with Tris-buffer (TBS, pH 7.6), SE plates were incubated in 100 µL of blocking buffer. Anti-vinculin antibody (Sigma-Aldrich, St. Louis, MO, USA) was used at a 1:50 dilution. Plates were incubated with primary antibody for 60 minutes at room temperature, followed by incubation with rhodamine-conjugated secondary antibody (Chemicon International, Inc., Temecula, CA, USA). After sufficient rinsing with TBS, SE plates were incubated with Alexa Fluor 488 conjugated to phalloidin (5 IU; Cambrex, Walkersville, MD, USA) for 20 minutes at room temperature. After washing, SE plates were mounted on PLA-coated glass plates with a mounting medium. Cell adhesion and the cytoskeleton were observed with a confocal laser scanning microscope (LSM 510 META, Zeiss, Jena, Germany).

**mRNA expression levels of fibronectin-1 and focal adhesion kinase (FAK)**
KMST-6 cells cultured on SE plates coated and uncoated with MPC polymer were harvested using lysis buffer. Total RNA was isolated using the RNeasy Mini Kit (Qiagen, Hilden, Germany). RNA concentrations were spectrophotometrically measured at 260 nm, and RNA purity was confirmed from the ratio of absorbance at 260 nm to that at 280 nm. First-strand cDNA synthesis was performed using 1 µg of total RNA. RNA samples were reverse transcribed to cDNA using the SuperScript First-Strand Synthesis System (GIBCO BRL, Invitrogen Corp., NY, USA).

Real-time PCR was performed on a TP800BK Thermal Cycler Dice Real Time System (Takara Biochemicals, Kyoto, Japan) using a SYBR\textsuperscript{®} Premix Ex Taq\textsuperscript{™} kit (Takara Biochemicals, Kyoto, Japan). Thermal cycling conditions were as follows: initial hold for 2 minutes at 50°C and 10 minutes at 95°C, followed by 40 cycles at 95°C for 15 seconds and 60°C for 60 seconds. Primer pairs used for fibronectin-1 and FAK, as well as for the reference gene (housekeeping gene) β-actin, were designed using DINAsis software (Toshiba, Tokyo, Japan) and were produced by Nippon Gene (Sendai, Japan). The primers were fibronectin-1 (NM_212482), 5'-GAGCCATGTGTCTTTACCACT-T-3' (forward), 5'-AGTATTTCTGGTCCTGCTCA-T-3' (reverse); FAK (L13616), 5'-ACTTGAGCGATGTATGGAG-3' (forward), 5'-GCTGTCGAAGTACAGTTT-3' (reverse); and β-actin (NM_001101.3), 5'-
CTAAGCTAGTCCGTAGAAGCA-3' (forward), 5'-TGGCACCAGCACATGAA-3' (reverse). Relativity quantities normalized against β-actin were calculated, and all reactions were performed in triplicate. Quantification of mRNA expression levels was achieved using a standard curve, as reported previously.

Protein expression levels of ROCK1, coflin, paxillin, and talin
Protein expression was assessed by Western blotting analysis. Cells cultured on SE plates coated and uncoated with MPC polymer were lysed in RIPA buffer (1×TBS, 1% Nonidet P-40, 0.5% sodium deoxycholate, 0.1% SDS, 0.004% sodium azide, 1% sodium orthovanadate, 1% phenylmethylsulfonyl fluoride (PMSF), and 1% protease inhibitor cocktail). Lysates were centrifuged at 13,000 g for 5 minutes at 4°C to collect the supernatant. Protein concentration was measured using Bradford protein assay (Bio-Rad Laboratories, Hercules, CA, USA). Protein samples were separated by 12% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) with the Mini-PROTEAN Tetra cell (Bio-Rad Laboratories, CA, USA).

Proteins were immobilized on a polyvinylidene fluoride (PVDF) transfer membrane by electro-transfer using a Mini Trans-Blot Cell (Bio-Rad Laboratories, CA, USA). Immobilized proteins were blocked overnight with 3% milk solution in 1×phosphate buffered saline (PBS) containing 0.1% Tween 20 (PBST), and then incubated for 30 minutes with each primary antibody (anti-ROCK1, 1:1000 dilution; anti-cofilin, 1:1000 dilution; anti-paxillin, 1:500 dilution; and anti-talin, 1:250 dilution). Membranes were then washed with PBST and incubated with HRP-conjugated donkey anti-rabbit IgG (1:2000 in PBST with 3% milk;...
Dako, Carpinteria, CA, USA) for 1 hour at room temperature. After washing with PBST, antibody reactivity was visualized using ECL (GE Healthcare, Buckinghamshire, UK) by exposure to X-ray film (Fujifilm Corp., Tokyo, Japan).

Statistical analysis
All data were presented as mean±standard deviation. Three independent cultures were used for each biochemical analysis. Statistical differences were analyzed by one-way ANOVA and Tukey’s test using JSTAT 12.5 software. Differences at \( p<0.05 \) were considered to be statistically significant.

RESULTS

XPS and EDX
As MPC was used to simulate the surface structure of a natural biomembrane, phosphate components were investigated. On the surface of the SE plate coated with MPC polymer, peaks for \( P_{2s} \) and \( P_{3p3} \) were observed in its XPS spectrum (Fig. 2). Similarly, the corresponding EDX spectrum also revealed the peak for phosphate (Fig. 3, upper right). Taken together, these results indicated that the SE plate was successfully coated with MPC polymer.

After 12 weeks of implantation, EDX analysis revealed that the MPC polymer coating remained on the SE plates removed from the animals (Fig. 3, lower), indicating that the MPC polymer coating remained stable and intact throughout the entire implantation period.

Histology
At 2 weeks after implantation, SE plates in the control and experimental groups were surrounded by fibrous connective tissue with slight inflammatory cell infiltration (Fig. 4). Moreover, at 2 weeks, fibrous tissue was thin and thickness did not differ between the experimental and control groups (\( n=6, p>0.05 \)).

In the control group, the thickness of fibrous connective tissue increased steadily at 4 and 8 weeks (Fig. 5). Besides, capillaries and some inflammatory cells were seen around the fibrous connective tissue. At 12 weeks, dense fibrous tissue was observed around the SE plate.

![Fig. 3 EDX imaging of SE plate surfaces in control (upper left) and experimental groups before implantation (upper right), and that of experimental group after 12 weeks of implantation (lower). ↑: Peaks for phosphate.](image-url)
In the experimental group, the thickness of fibrous tissue showed a gradual decrease at 8 and 12 weeks (Fig. 4). Moreover, at 4, 8, and 12 weeks, the surrounding fibrous tissue in the experimental group was thinner than that in the control group ($n=6, p<0.05$; Fig. 5). At 12 weeks, the fibrous tissue contained elastic fibers without inflammatory cell infiltration and capillary formation.

**Collagen staining**

Through a polarizing microscope, Sirius red and Fast green staining showed the collagen components of the fibrous tissue (Fig. 6). Figure 7 further shows the amounts of collagen fibers in fibrous capsules around the SE plates at 2, 4, 8, and 12 weeks for both experimental and control groups.

At 2 weeks, fibrous connective tissue was composed of numerous collagen fibers and scattered non-collagenous proteins such as proteoglycans and elastin in the experimental group. At 4 weeks, non-collagenous protein fibers gradually increased in both groups. Nonetheless, the amount of collagen in the experimental group was lower than that in the control group ($n=6, p<0.01$). At 8 weeks, fibrous tissue in the experimental group comprised a large number of elastic fibers and other non-collagenous components—in sharp contrast to the control group. At 12 weeks, there was a gradual increase in non-collagenous proteins in fibrous tissue for both groups, with the experimental group showing a higher level ($n=6, p<0.01$).
Analysis of cytoskeleton and cell adhesion

Figure 8 shows the extent of cell growth in both the experimental and control groups. In the control group, numerous KMST-6 cells with a spindle-shaped morphology—which is a typical fibroblastic shape—were observed, as well as signs of colony formation. In the experimental group, there were scattered round-shaped cells over the surface but no signs of colony formation. Figure 9 shows the cell growth curves measured with MTT assay. It could be seen that cell proliferation on the MPC plate was slower than on the control plate.

In Fig. 10, phalloidin-positive cytoskeleton could be clearly seen. For the control group, KMST-6 cells had a fully stretched shape with close contact to the SE plate. In the experimental group, round-shaped cells were seen on the coated surface with a large number of filopodia. The cells contained large amounts of short and dense skeletal structure.

Confocal images of immunohistochemical staining showed that vinculin expression was significantly weaker in the experimental group than in the control group (Fig. 11).

mRNA expression levels of fibronectin-1 and FAK

The mRNA expressions of fibronectin-1 and FAK were quantified using real-time PCR. Figure 12 shows the relative expression levels of fibronectin-1 and FAK mRNAs in KMST-6 cells that adhered to the experimental and control surfaces. The relative expression levels of fibronectin-1 and FAK mRNAs in
the experimental group were clearly lower than those in the control group (n=3, p<0.01).

Protein expression levels of ROCK1, coflin, paxillin, and talin
Protein lysates of KMST-6 cells were analyzed using the Western blotting technique, and the results thereof are shown in Fig. 13. In the experimental group, the protein expressions of ROCK1, paxillin, and talin were significantly downregulated ($n=3$, $p<0.01$). However, the expression of coflin protein in the experimental group was significantly upregulated ($n=3$, $p<0.01$).

DISCUSSION
Fibrous tissue formation after biomaterial implantation can lead to treatment failures, serious clinical problems, or loss of implanted device function$^{26}$. To prevent fibrous tissue formation, intervention methods include administration of 5-fluorouracil, interferon-alpha 2b, and corticosteroids, UV irradiation or cryodesiccation therapy, and surgical treatment$^{27}$. However, these treatments are accompanied with various side effects, hence providing a strong impetus to improve the biocompatibility of implant materials to prevent unwanted fibrous tissue formation. Against this background, MPC polymer was developed as a material to inhibit harmful reactions at the interface between tissue and material$^{6}$. Thus far, MPC polymer has attracted considerable attention because of its ability to influence cell reactions$^{8}$. 

protein expressions of ROCK1, paxillin, and talin were significantly downregulated ($n=3$, $p<0.01$). However, the expression of coflin protein in the experimental group was significantly upregulated ($n=3$, $p<0.01$).
Inflammation and immune reactions around an implanted material will lead to development of fibrous tissue, and even increased fibrous tissue growth. Therefore, the thickness of fibrous tissue capsules and the collagen-synthetic response of fibroblasts are important parameters in the evaluation of a material’s biocompatibility\(^{27}\). In the present study, MPC polymer-coated materials exhibited inhibition of fibrous tissue formation and collagen synthesis. In the same vein, the fibrous tissue surrounding MPC polymer-coated materials showed higher synthesis rates of non-collagenous proteins, such as elastin and proteoglycans, without an active inflammatory response. Our results thus showed that MPC polymer might have the capacity to regulate the quantity and quality of fibrous tissue formation. This meant that MPC polymer could contribute to preventing the deformation of and assisting in the smooth movement of implantation areas.

In general, biomaterials such as metals and bone graft materials emphasize cell adhesion. On the contrary, improvements to MPC have rendered it to be a biomaterial that mimics a biological cell membrane and which inhibits protein adsorption and cell adhesion\(^{29}\). The inhibition of cell adhesion has a negative impact on fibrous tissue formation\(^{30}\). Therefore, in this study, cultured cells on MPC polymer exhibited a round-shaped morphology and weaker expression of vinculin protein. Vinculin, acting through binding to F-actin, plays an important role in focal cell adhesion, in a manner similar to integrins\(^{30}\). At the same time, large amounts of filopodia immunofluorescently labeled with F-actin were observed in cultured cells on MPC polymer. These findings thus demonstrated the enhanced cell migration and weakened cell adhesion of the cultured cells on MPC polymer.

Fibronectin (FN) secreted from cells plays a major role in cell adhesion. In an in vitro study using bovine serum by Sawada et al.\(^{31}\), MPC polymer surface registered a decrease in FN deposition. Similarly, results of the present study showed a significant reduction in FN expression of fibroblasts in contact with the MPC polymer. This meant that MPC polymer might have the capacity to diminish FN secretion upon contact with cells, thereby resulting in inhibited cell adhesion.

Integrin binding to FN leads to CAP formation, and at the same time induces downstream signaling. In other words, CAP formation mediated by FN induces signal transduction. Recent research has demonstrated a close relationship between integrin, vinculin, talin, and paxillin molecules and cell adhesion\(^{30}\). These molecules combine with fibronectin in the local microenvironment to form CAP. After CAP formation, cells perceive mechanical changes, and in response regulate their growth, secretion, plasticity and other physiological functions\(^{30}\).

In the present study, the expression of the downstream signaling molecules of FN was investigated using the Western blotting technique and a decrease in talin expression was detected. Owing to inhibited FN expression, there was a decrease in CAP formation and hence a decrease in talin expression —because inhibited FN expression affected downstream signal transduction. Besides, a downregulated paxillin expression reflected a cell migration trend, and this finding was augmented by a weaker expression of focal adhesion kinase (FAK). FAK represents a 125-kDa protein which is localized to focal adhesion and which is activated by tyrosine phosphorylation in response to integrin clustering\(^{32}\). Taken together, these results showed that the MPC polymer inhibited CAP formation by affecting downstream molecular signaling.

Enhanced cell migration on MPC polymer has been reported\(^{30}\), whereby actin expression is an important parameter of cell migration capacity. ROCK1 and cofilin regulate actin metabolism through the Rho signaling pathway. ROCK1 maintains cell morphology by increasing the stability of actin-based stress fibers. ROCK1 also phosphorylates and activates LIM kinase, which phosphorylates cofilin. Cofilin stimulates actin depolymerization and changes in cell structure, and the phosphorylation of cofilin by LIM kinase suppresses its activity\(^{30}\). In this study, cultured cells on MPC polymer showed decreased ROCK1 and increased cofilin activity, leading to accelerated depolymerization of actin. Active actin metabolism with a large amount of filopodia formation represented enhanced cell
migration, affecting collagen metabolism by fibroblasts as a result. Cell adhesion has a close relationship with collagen. Okajima et al. found that with reduced secretion of Type I collagen—which is a major matrix component, fibrous tissue formation was prevented.

Results of the present study suggested that fibrous tissue formation was prevented on MPC polymer-coated material because of decreased adsorption of proteins on the MPC polymer-coated silicon surface. In a study by Abraham et al. which investigated the adsorption of laminin, collagen, and fibronectin on polymer membrane surfaces, it was shown that the adsorption of proteins on biomaterial surfaces modulated cellular viability and restricted proliferation. In the context of the present study, the modulated cellular interaction with the MPC polymer-coated surface resulted in decreased secretion by and amplification of fibroblasts. Together with the concomitant regulation of signaling pathways such as Rho and FAK, scar formation was evidently attenuated.

In conclusion, this study showed the usefulness of artificial devices coated with MPC polymer, which mimicked a natural cell membrane. Nonetheless, for MPC polymer-coated implant materials to be used in a wide range of clinical applications, further research must be done to understand the role of direct metabolism exchange between cell membrane lipids and artificial phospholipids.

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