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Polymeric stent materials dysregulate macrophage and endothelial cell functions: implications for coronary artery stent

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Abstract

Background—Biodegradable polymers have been applied as bulk or coating materials for coronary artery stents. The degradation of polymers, however, could induce endothelial dysfunction and aggravate neointimal formation. Here we use polymeric microparticles to simulate and demonstrate the effects of degraded stent materials on phagocytic activity, cell death and dysfunction of macrophages and endothelial cells.

Methods—Microparticles made of low molecular weight polyesters were incubated with human macrophages and coronary artery endothelial cells (ECs). Microparticle-induced phagocytosis, cytotoxicity, apoptosis, cytokine release and surface marker expression were determined by immunostaining or ELISA. Elastase expression was analyzed by ELISA and the elastase-mediated polymer degradation was assessed by mass spectrometry.

Results—We demonstrated poly(*D,L*-lactic acid) (PLLA) and polycaprolactone (PCL) microparticles induced cytotoxicity in macrophages and ECs, partially through cell apoptosis. The particle treatment alleviated EC phagocytosis, as opposed to macrophages, but enhanced the expression of vascular cell adhesion molecule-1 (VCAM) along with decreased nitric oxide production, indicating ECs were activated and lost their capacity to maintain homeostasis. The activation of both cell types induced release of elastase or elastase-like protease, which further accelerated polymer degradation.

Conclusions—This study revealed that low molecule weight PLLA and PCL microparticles increased cytotoxicity and dysregulated endothelial cell function, which in turn enhanced elastase release and polymer degradation. These indicate polymer or polymer-coated stents impose a risk of endothelial dysfunction after deployment which can potentially lead to delayed endothelialization, neointimal hyperplasia and late thrombosis.

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Keywords

poly(*L*-lactic acid); PLLA; polycaprolactone; PCL; stent; coronary; polymer degradation; macrophage; endothelial cell; dysfunction; elastase

1. Introduction

Drug-eluting stents (DES) have remarkably reduced acute in-stent thrombosis and the occurrence of revascularization. However, late stent thrombosis associated with DES still remains unexpectedly high, which can be partially attributed to endothelial dysfunction and vasomotor disorders caused by stent, drug or polymer coatings.[1] Besides the stent geometry, the influence of incorporated drugs on inducing endothelial dysfunction has been studied in both animals and humans over the past decade.[2-4] In contrast, polymer coating-induced endothelial dysfunction has yet been reported despite of their proven hypersensitive effect. Biodegradable and resorbable polymers provide appealing substitutes to durable polymer coatings, since they are less likely inducing severe chronic inflammatory response and their degradation products can be cleared out of the human body. Coronary artery stents coated with or completely made of biodegradable polyesters, i.e. poly(lactic acid) (PLA), poly(lactic-*co*-glycolic acid) (PLGA) and to a lesser degree, poly(ϵ -caprolactone) (PCL), have been developed.[5-9] These polymers undergo hydrolysis and enzyme-mediated degradation (“biodegradation”),[10, 11] resulting in a release of low molecular weight degradation products. The released polymer debris is then removed by leukocytes and foreign body giant cells (“bioresorption”), as indicated by enhanced phagocytosis and cell activation. Activated leukocytes subsequently alter the phenotype of surrounding cells (i.e. endothelial cells) and make them actively involved in clearing and remodeling processes. Endothelial cells (ECs) are normally quiescent and maintain vascular homeostasis. The activation of endothelium promotes the adhesion of platelets and leukocytes while losing its cell-cell integrity, which gradually leads to vascular malfunctions such as atherosclerosis and stenosis.[12] For example, neointimal formation has been found with the degradation of polymer stents, [13, 14] as revealed by locally elevated inflammation and cell growth to intima. Although it can be alleviated by timed-release anti-proliferative drugs, locally delivered therapeutics often induce EC apoptosis/necrosis, impair re-endothelialization and lead to late thrombosis.[15, 16] It is unclear, however, whether there is a casual relationship between polymer debris and endothelial activation/dysfunction. Therefore, the objective of this study is to understand the interaction between polymer debris and ECs, which is particularly meaningful for polymer coronary stents. Here we use low molecular weight poly(*D,L*-lactic acid) (PLLA) and PCL microparticles as model molecules of degradation products to facilitate cellular uptake and to elucidate how macrophage and arterial endothelial cell functions can be regulated by degraded polymers. Result of this study will help interpret the occurring restenosis and late-thrombosis in polymer-coated stents and warrant more careful evaluation of endovascular polymer devices in general. It will enlighten both clinicians and biomedical engineers to improve the design of polymer-involved coronary stents from a perspective of minimizing the deteriorating effect of polymers on endothelium.

2. Methods

2.1 Preparation of stainless steel, PLLA and PCL microparticles

PLLA (RESOMER R202S, 10-18k Da, Sigma, St, Louis, MO) or PCL (1.25k Da, Sigma) was dissolved in dichloromethane (10% w/w) and the polymer solution was added to 1% poly(vinyl acid) (PVA) water solution drop-wise under continuous stirring at 3000 rpm (Silverson L4RT homogenizer, East Longmeadow, MA). The polymer solution was then slowly stirred at room temperature overnight to evaporate the organic solvent. The PVA was removed by repeated washing with DI water and the particles were collected by centrifugation and lyophilizing overnight. Particles less than 38 μm were obtained by sieving (H&C Sieving Systems, Columbia, MD). Stainless steel (SS) particles with around 10 μm in diameter were prepared by grinding and served as controls throughout this study. Particle size and morphology were visualized using a Hitachi S4200 high resolution scanning electron microscope (SEM, Schaumburg, IL) (Fig. 1).

2.2 Cell culture

Human blood-derived monocytes were obtained from Advanced Biotechnologies (Columbia, MD) and differentiated into human blood monocyte-derived macrophages.[17] Monocytes at a density of 2×10^6 cells/ mL were incubated with 10 mL DMEM (Invitrogen, Carlsbad, CA) with 20% fetal calf serum (Invitrogen), 10% human serum (Sigma), and 5 ng/ mL macrophage colony-stimulating factor (Sigma) for 9 days to differentiate into macrophages.[18] Human coronary artery ECs (HCAECs, Cell Applications, San Diego, CA) were cultured in endothelial cell growth medium (Cell Applications) until 80% confluent before seeding. Polymer or SS particles were UV sterilized for 1 hour prior to cell seeding. Macrophages (3×10^5 cells/ mL) were cultured with polymer particles (400 μg / mL of PLLA or PCL) or SS particles (8000 μg / mL) for 3 days before endpoint assays.

2.3 Phagocytosis and superoxide assays

After 3 days culture, macrophages or HCAECs were treated with green-fluorescent *Escherichia coli* (*E. coli*) particles for 2 hours according to the manufacturer's protocol (Vybrant® Phagocytosis assay kit, Life Technologies, Carlsbad, CA). Cells were stained with dihydroethidium (1 μg / mL DHE, Life Technologies) for 15 minutes. Cell nuclei were counterstained with Hoechst and imaged with an inverted Nikon Ti fluorescent microscope (Nikon Instruments, Melville, NY). Fluorescence intensities of *E. coli* particles phagocytized by activated cells and superoxide as indicated by DHE were measured using a plate reader (Tecan Group, Ltd Männendorf, Switzerland). Phagocytic activity and superoxide production were normalized to cell number quantified by Hoechst nuclear staining (n = 4 per polymer condition).

2.4 Cytotoxicity

Macrophage and endothelial cell cytotoxicity was measured using Cytotox One Homogeneous Membrane Assay (Promega, Madison, WI) by quantifying lactate dehydrogenase released from damaged cells according to supplier's protocol. Fluorescence

values were normalized to cell samples without material treatment (n=4 per polymer condition).

2.5 TNF- α and nitric oxide expression

After 3 day culture, media samples were collected and analyzed for secretion of tumor necrosis factor (TNF)- α , elastase, and nitric oxide (NO). TNF- α secretion was measured using Human TNF- α ELISA Max Standard Kit (Biolegend, San Deigo, CA) according to supplier's protocol. NO secretion was measured by incubating 100 μ L 1X modified Greiss reagent (Sigma) with 100 μ L media or nitrate standards for 15 minutes before reading the absorbance at 540 nm using a plate reader (Tecan). Nitrate standards (0-65 μ mol) were made using sodium nitrate (Sigma) (n=4 per polymer condition).

2.6 Immunostaining of VCAM-1 and Annexin V

Cells were fixed with 4% paraformaldehyde (PFA) for 15 minutes, blocked with 10% goat serum in Dulbecco's phosphate-buffered saline (DPBS) for 1 hour, and incubated with allophycocyanin-conjugated VCAM-1 antibodies (1:100 dilution, Abcam, Cambridge, MA) for 2 hours in DPBS. For Annexin-V staining, HCAECs were fixed with 4% PFA for 15 minutes, permeabilized with 0.25% TritonX-100 for 10 minutes, blocked with 10% goat serum for 1 hour, incubated with rabbit anti-human Annexin V antibodies (1:200 dilution, Abcam) overnight at 4°C, and then incubated with Alexa 488 goat anti-rabbit antibodies (1:100 dilution, Abcam) for 1 hour in DPBS. After washing twice with DPBS, cells were imaged on a LSM 710 META inverted confocal microscope (Zeiss, Thornwood, NY). The number of cells that were positive to each staining was counted and divided by the total number of cells per image field (n=6 images from three replicate experiments).

2.7 Elastase activity

In order to measure elastase activity, elastase standards (0-10 nmol) were made using human neutrophil elastase (Sigma). 200 μ L of media samples or standards were incubated with 200 μ L of 1mM granulocyte elastase substrate (Sigma) in DPBS for 1 hour at 37°C and glacial acetic acid was added to stop the reaction. Absorbance was measured at 415 nm by a plate reader (Tecan) (n=4 per polymer condition).[19]

2.8 Polymer degradation by elastase

PDLLA particles were suspended in DPBS with or without elastase (0.08U/mL, Sigma), and incubated at 37°C for 7 days. Insoluble polymer particles were filtered off using centrifugation units (Amicon-0.5ml 3K MWCO, Millipore, Billerica, MA) at 12,000g for 20 min. Degradation products in the filtered solution were confirmed by liquid chromatography coupled with mass spectrometry (LC-MS) (Variant 1000, Agilent, Santa Clara, CA) using a C18 column (Kinetex, Phenomenex, Torrance, CA) with an acetonitrile-H₂O mobile phase system. (n=3 per polymer condition).

2.9 Statistical analysis

In all experiments, analytical results are expressed as means \pm standard error of the mean (SEM). One-way ANOVA was used to determine if statistical differences exist between

groups. Comparisons of individual sample groups were performed using unpaired Student's t-test. For all experiments, $p < 0.05$ was considered statistically significant.

3. Results

3.1 Cell number and cytotoxicity of macrophages and ECs

The numbers of macrophages and ECs were determined by measuring the fluorescence intensity of Hoechst staining cell nuclei. The cell number significantly decreased in both cell types after being cultured with PLLA, PCL or SS microparticles for 3 days ($p < 0.05$, Fig. 2a) compared to no polymer control of the corresponding cell type. However, no significant difference in cell number was observed among the three test material types. The cytotoxic effect of these microparticles, presented by lactase dehydrogenase release from cells resulting from a damaged membrane, was shown in Fig. 2b. Apparently, the decrease of cell number correlates to the increase of cytotoxicity of materials. Compared to no polymer control, PLLA, PCL and SS microparticles showed a remarkable increase of cytotoxicity to macrophages ($p < 0.05$, Fig. 2b). Interestingly, there was no significant difference in cytotoxicity between PLLA and PCL microparticles, although PCL but not PLLA microparticles considerably increased cytotoxicity to ECs. As a negative control, SS microparticles has higher cytotoxicity to both cell types than PLLA and PCL microparticles ($p < 0.05$, Fig. 2b). These results demonstrate that the macrophages are more susceptible to cytotoxic effects of these microparticles than ECs as more severe cytotoxicity of the materials was seen in macrophages (Fig. 2b).

3.2 Phagocytic activity of macrophages and ECs

Phagocytic activity of cells was measured by quantitating the intensity of intracellular fluorescence emitted from green fluorescent-tagged *E. coli* particles with or without the treatment of microparticles. The addition of PCL and SS microparticles enhanced the phagocytic activity of macrophages compared to no polymer control ($p < 0.05$, Fig. 2c), whereas PLLA microparticles had no further effect on macrophage phagocytosis ($p < 0.05$). In contrast, the phagocytosis of *E. coli* particles by ECs was significantly decreased after adding PLLA, PCL and SS microparticles ($p < 0.05$, Fig. 2c), indicating the phagocytosis stimulated by these microparticles is cell type-dependent.

3.3 Apoptosis of ECs

Apoptosis of ECs induced by microparticles was measured by Annexin-V staining. Treatment of microparticles significantly promoted EC apoptosis compared to no particle control ($p < 0.05$, Fig. 2d-e). Apoptosis was even more dramatically increased with addition of PCL or SS microparticles, indicating a detrimental effect of these microparticles on EC survival.

3.4 VCAM-1 expression of ECs

The expression of VCAM-1 on ECs induced by microparticles was shown in Fig. 3a. Compared to ECs cultured with PLLA microparticles or no polymer control, ECs treated with PCL microparticles increasingly expressed VCAM-1 in both mono- and co-culture with macrophages. The quantitation of VCAM-1 positive cells showed that treatment of PCL and

SS microparticles resulted in a significantly higher number of VCAM-1 positive ECs compared to no polymer controls in both mono- and co-culture with macrophages ($p < 0.05$, Fig. 3b). In contrast, the addition of PLLA microparticles only slightly increased VCAM expression in both culture conditions compared to no polymer control.

3.5 Expression of TNF- α , superoxide, and NO in macrophages and/or ECs

The TNF- α concentration in cell culture supernatants was determined by ELISA. Generally, the TNF- α expression increased in both macrophages and ECs by adding PLLA and PCL microparticles, although there was no significant difference compared to no polymer controls or between PLLA and PCL groups (Fig. 4a). The production of ROS was determined by staining cells with dihydroethidium that relocates from cytosol to nuclei after being oxidized to oxyethidium and emits red fluorescence. Results showed no significant differences among the test groups in each cell type (Fig. 4b).

NO production in EC mono-culture was quantified. Results showed the addition of PLLA, PCL and SS microparticles significantly reduced NO production in ECs, with the least NO production in ECs treated with SS microparticles (Fig. 5). PCL induced lower NO production than PLLA microparticles. However, no statistically difference was observed among these test materials.

3.6 Elastase activity and polymer degradation

The activity of elastase in cell culture supernatant was determined. The three types of microparticles promoted elastase secretion from both macrophages and ECs ($p < 0.05$, Fig. 6), compared to no polymer control. However, there was no significant difference between PLLA and PCL groups, although SS and PCL microparticles induced the highest elastase expression in ECs and macrophages, respectively. ECs also generally had a higher elastase activity than macrophages (Fig. 6).

The elastase-mediated degradation of PLLA microparticles was demonstrated by LC-MS (Fig. 7). Slow degradation of PLLA microparticles was observed in DPBS without elastase as identified by various lengths of PLLA polymer chain (repeating unit $m/z = 72$) at m/z 185, 257, 329, 401, 473, 545 and 617 [20, 21] (Fig. 7a). When elastases were added (Fig. 7b), the degradation of PLLA microparticles was greatly enhanced with an increase in the relative abundance of molecules at the various m/z values mentioned above.

4. Discussion

As an important group of implantable biomaterials, polyesters have been widely used for biomedical and clinical applications. Here we report the treatment of representative polyesters (PLLA and PCL) can lead to dysfunctional activation of ECs and macrophages, which in turn promotes further elastase-mediated polymer degradation and contribute to this degradation-phagocytosis-dysfunction loop. Low molecular weight microparticles were used in this study as model molecules of degradation products that could be quickly accessed by cells.

PLLA is generally considered non-toxic and non-inflammatory. For example, no cytotoxicity to macrophages and Schwann cells was observed during the interaction with PLLA films and micro-/nanofibers.[22, 23] In other studies, microparticles with a diameter of $\sim 1.6 \mu\text{m}$ were made from PLLA (100kDa) and PLGA (100kDa) blends which did not decrease cell viability of human lung cancer cells, showing PLLA/PLGA microparticles at these specific molecular weights and particle sizes are not cytotoxic to cancer cells.[24] However, PLLA microparticles of small molecular weight (10kDa) with $\sim 5 \mu\text{m}$ in diameter decreased cell viability of both human primary and murine RAW macrophages in a dose-dependent manner.[25] Moreover, pre-degraded PLLA microparticles ($<40 \mu\text{m}$) with a range of molecular weight from 99 to 53kDa induced phagocytosis and cell death of macrophages in mice.[26] These results indicate that the cytotoxic effect of PLLA on macrophages is closely related to phagocytosis, geometry, molecular weight and dose. Similarly, apoptosis or necrosis of macrophages mediated by the uptake of PCL nanoparticles (42kDa) was dose- and size-dependent, evidenced by the fact that a higher dose and a smaller particle size resulted in lower cell viability.[27] When high molecular weight PCL underwent hydrolytic degradation, macrophages phagocytized smaller-molecular-weight degradation products of PCL.[28] Especially, the fragments less than 3kDa could be completely degraded in phagosomes of macrophages.[29] In this study, we showed the low molecular weight PLLA and PCL microparticles (10k and 1.25kDa) increased phagocytic activity of macrophages and exhibited higher cytotoxicity than no treatment groups (Fig. 2b-c). As PCL degrades slowly, lower levels of macrophage activation and accumulation were expected compared to fast degrading PLLA and PLGA, thus causing less phagocytosis. However, both phagocytic activity and cytotoxicity of macrophages were enhanced in PCL groups (Fig. 2b-c), indicating the hydrolytic degradation of PCL in this study is possibly less important than the elevated particle uptake. The PCL microparticles being used in this study had a molecular weight of 1.25kDa which could be readily phagocytized by macrophages.[29] Moreover, the surface properties of PCL microparticles might contribute to the higher intracellular uptake. It was reported the cellular uptake of particles could be enhanced by surface-coated bovine serum albumin.[30] The adsorption of serum proteins on PCL is theoretically higher than on PLLA due to the higher hydrophobicity of PCL, which may enhance the uptake of PCL particles compared to PLLA particles. In summary, the increased phagocytic activities associated with low molecular weight polyester microparticles, PCL in particular, played a key role in the loss of macrophage viability and cell number in this study (Fig. 2a-c).

Although PLLA is considered non-toxic to ECs, its unfavorable surface properties can decrease EC adhesion and dysregulate cell functions. For example, PLLA (800kDa) membrane reduced the viability of human umbilical vein ECs (HUVECs)[31] and PLLA (126kDa) nanofibers slightly reduced the number of adherent bovine aortic ECs after 3 days of culture[32]. In this study, the addition of PLLA and PCL microparticles decreased both cell number and viability of human coronary artery ECs (Fig. 2a-b). Polyesters have been reported to induce EC and macrophage apoptosis.[33, 34] Here the loss of cell number and viability seems partially due to cell apoptosis induced by polymer particles, as evidenced by Annexin V positive ECs after treatment with PCL particles (Fig. 2d,e). Again, polymer molecular weight, particle size, and surface modification may be responsible for the increased apoptosis observed with PLLA and PCL particles.

ECs phagocytize dead cells and cell debris, e.g. apoptosis bodies[35], senescent neutrophils[36] and red blood cells[37]. The phagocytosis of necrotic cells induces activation of ECs by releasing TGF- β 1 and IL-6 [38, 39] and increases ICAM-1 expression to facilitate macrophage adhesion.[40] In this scenario, phagocytosis dysregulates EC function and eventually leads to cell death.[37] ECs have also been shown to phagocytize polymer particles. [41-43] In contrast to enhanced macrophage phagocytosis, the phagocytic activity of ECs decreased upon microparticle treatment (Fig. 2c). However, as a marker of EC activation[44, 45], VCAM-1 expression increased significantly in ECs upon particle treatment (Fig. 3). Thus the phagocytic activity of ECs inversely correlates to their activation (Fig. 2c and 3). In order to understand the regulatory mechanism, we further quantified the production of TNF- α and superoxide since they are critical to endothelial activation. For example, autocrine TNF- α induces the expression of VCAM-1 in ECs and activates ECs to release other cytokines such as IL-6, IL-8, GM-CSF.[46, 47] TNF- α also contributes to EC activation via ROS production.[48, 49] In this study, we demonstrate the addition of PLLA and PCL microparticles did not significantly alter the production of TNF- α and ROS in either macrophages or ECs (Fig. 4), indicating a non-TNF- α nor ROS-dependent mechanism to regulate VCAM-1 expression. NO keeps the endothelium quiescent, inhibits the adhesion and activation of platelets and leukocyte and maintain the normal endothelium function. Therefore, we further revealed that the production of NO from ECs was significantly reduced by adding microparticles ($p < 0.05$, Fig. 5). Along with the increase of VCAM-1 expression, the loss of ability to produce NO leads to a pro-inflammatory vascular phenotype and indicates EC dysfunction after the microparticle treatment.[50] Therefore the reduced phagocytosis of microparticles in ECs (Fig. 2c) can be further considered as a consequence of NO-dependent EC activation (Fig. 2d and 3).

In addition to cytokines, extracellularly released proteases could also induce EC dysfunction. Elastase is one of such proteases that actively participate in vascular wall remodeling after injury by degrading extracellular matrices such as elastin.[51] Furthermore, it was shown that the enzymatic activity of leucocyte elastase led to endothelial cell damage. [52-54] Similar to leucocytes such as neutrophils and macrophages, ECs secrete an elastase-like enzyme which breaks down the same substrate that leucocyte elastase can degrade.[55] In this study, ECs exhibited a higher elastase activity than macrophages, and the treatment of cells with PLLA or PCL microparticles significantly enhanced elastase expression in both cell types (Fig. 6), which potentially contributed to EC dysfunction as discussed above. On the other hand, elastase expressed by macrophages and ECs could degrade polyesters and further enhance the cellular uptake of degradation products and activate cells. The degradation of PDLLA by neutrophil elastase as a model was demonstrated by mass spectrometry (Fig. 7), showing the active involvement of elastase in cell-material interactions and degradation product-induced EC activation and dysfunction.

This study enriches our knowledge on polymer-mediated endothelial dysfunction, which is particular important for polymer-coated drug-eluting coronary stents since late-thrombosis has been persistently observed after the drug is exhausted, indicating a possible role of polymer in preventing endothelial healing. Characterized by elevated inflammation and cell ingrowth from tunica media and adventitia, neointimal formation has been found with the

degradation of polymer stents [13, 14]. Although the acute thrombosis and neointimal formation can be alleviated by applying an anti-proliferative drug, as a consequence, locally delivered therapeutics often induce EC apoptosis/necrosis, impair re-endothelialization and lead to late thrombosis.[15, 16] Endothelial dysfunction causes plaque formation and plays a key role in atherosclerosis.[56] In parallel, neoatherosclerosis has been found in drug-eluting stents which occurred earlier with higher frequency than in bare metal stents, again indicating a drug and/or polymer effect on incomplete endothelialization.[57, 58] Here the impairment of coronary artery endothelial function by biocompatible polymers such as PLLA and PCL was demonstrated.

4.1 Limitations

The major limitation of this study is lack of demonstration of endothelial dysfunction in an animal model using, for example, an acetylcholine assay. We realized this and chose low molecular weight polymer microparticles to mimic the effect of degraded polymer debris *in vitro*. The size of polymer microparticles ranged approximately from 1 to 38 μ m in this study, which might limit the accurate interpretation of size effect of particles on cellular response, although it was obvious in this study that microparticles within this range elicited cell dysfunction, making fine-tuning particle size unnecessary. The third limitation is we selected PLLA and PCL and drew the conclusions based on these polymers due to their clinical potential as a degradable stent material. However, possibilities cannot be excluded that other degradable biopolymers may be less cytotoxic with reduced dysregulating effects on macrophages and ECs. More careful examinations need to be performed to evaluate degradable polymers for coronary stent applications.

5. Conclusions

A delayed re-endothelialization has been observed with polymer-coated DES, for which the local delivery of anti-proliferative drugs is considered to be responsible. However in this study, we report for the first time that endothelial cell dysfunction can be also induced by biodegradable polymers such as PLLA and PCL, which are widely used as biocompatible materials. Moreover, microparticles made of these materials could lead to apoptosis and death of macrophages and endothelial cells. Upon exposure to polymer microparticles, cells increased the release of proteases such as elastase for polymer degradation, which in turn activated both cell types and resulted in cell dysfunction. The results of this study add important knowledge to cellular/tissue response to polymer-coated DES and warrant more careful evaluation of biodegradable polymers for coronary stent application.

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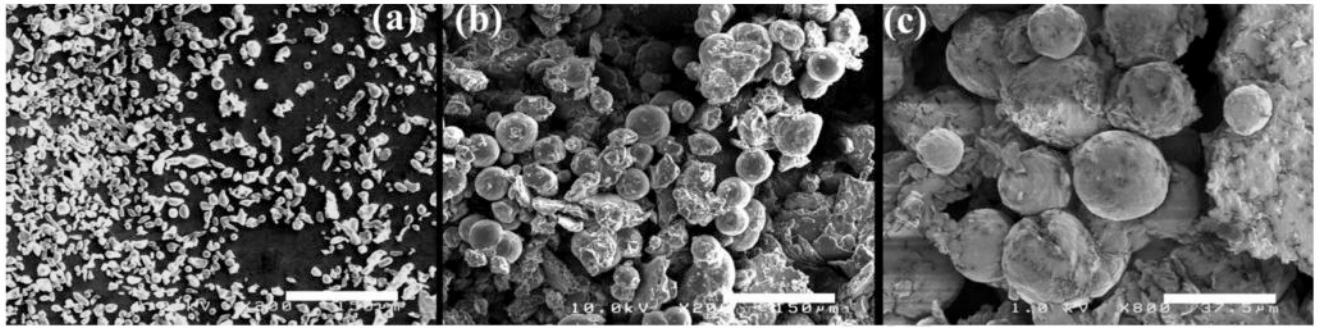


Figure 1.

SEM images of microparticles: (a) stainless steel, (b) PLLA and (c) PCL. Scale bar: 150 (a and b) or 37.5 (c) μm .

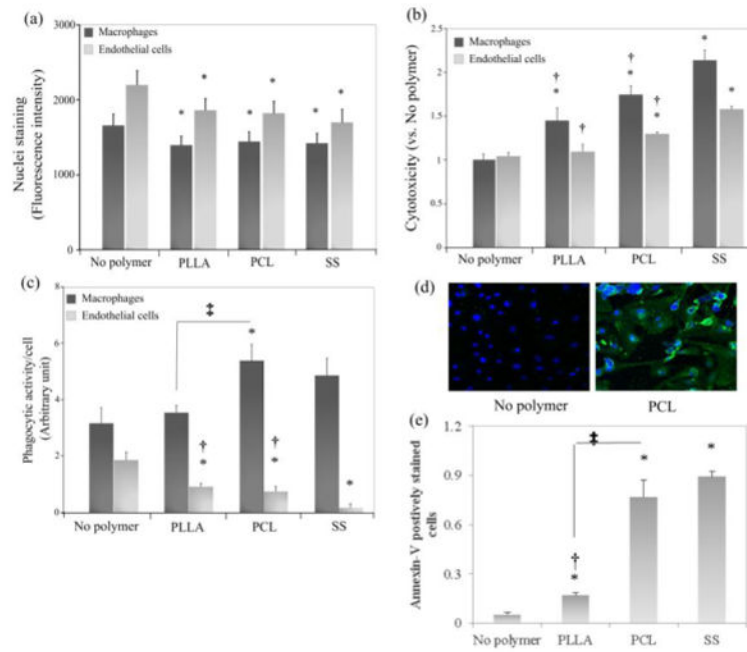


Figure 2.

Cytotoxicity, phagocytosis and apoptosis by interaction with microparticles: (a) Cell number of each group indicated by Hoechst fluorescence intensity; (b) cytotoxicity measured by lactate dehydrogenase released from damaged cells and normalized to no polymer control; (c) phagocytosis of fluorescent-tagged *E. coli* particles by activated macrophages and ECs. The fluorescence intensity emitted from *E. coli* particles was normalized to the corresponding cell number measured by Hoechst staining; (d) apoptosis in untreated (left) and PCL microparticle-treated ECs (right), blue: Hoechst, green: Annexin V; (e) quantification of apoptosis presented as the ratio of Annexin V-positive cells relative to the total cell number. * $p < 0.05$ versus no polymer control; † $p < 0.05$ versus SS control; ‡ $p < 0.05$ between PLLA and PCL groups connected by line.

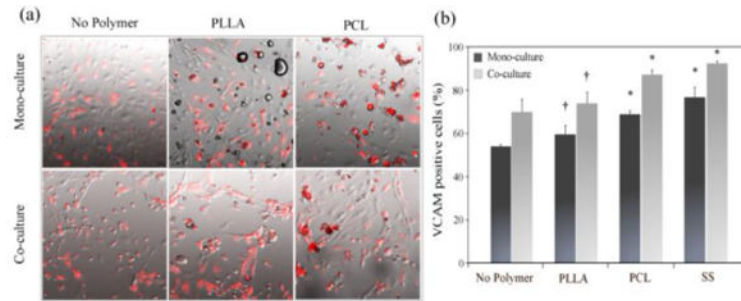


Figure 3. Microparticle-induced EC activation : (a) Merged images of APC-VCAM staining (red) and phase contrast; (b) VCAM-1 positive cells (%) relative to the total cell number in each sample. * $p < 0.05$ versus no polymer control; † $p < 0.05$ versus SS control.

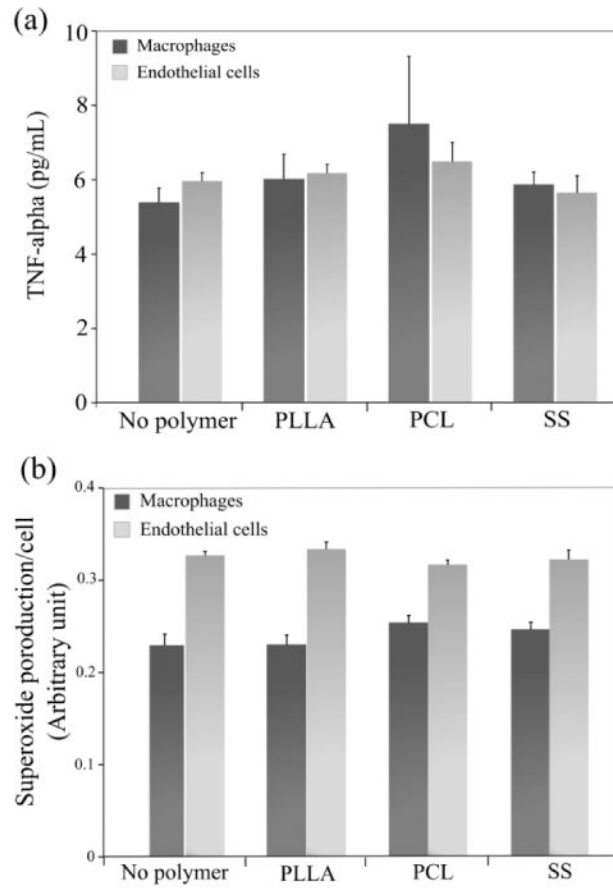


Figure 4. Microparticle-mediated macrophage activation: (a) TNF- α release into cell culture media which was determined by ELISA; (b) An intracellular superoxide level which was stained by DHE and normalized to the corresponding cell number measured by Hoechst staining.

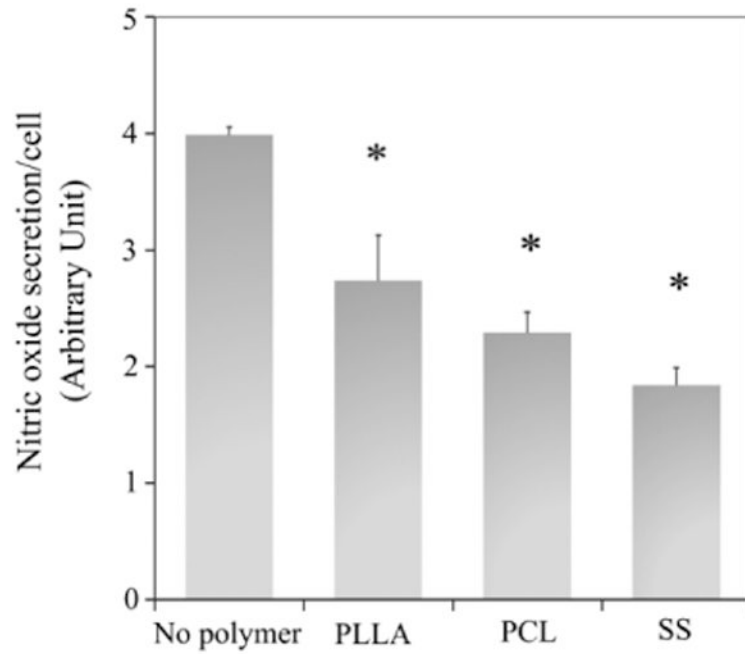


Figure 5. EC dysfunction induced by microparticles was determined by quantitating the amount of nitric oxide in supernatant (Greiss reagent). The absorbance was measured at 540nm and normalized to the cell number measured by Hoechst staining. * $p < 0.05$ versus no polymer control.

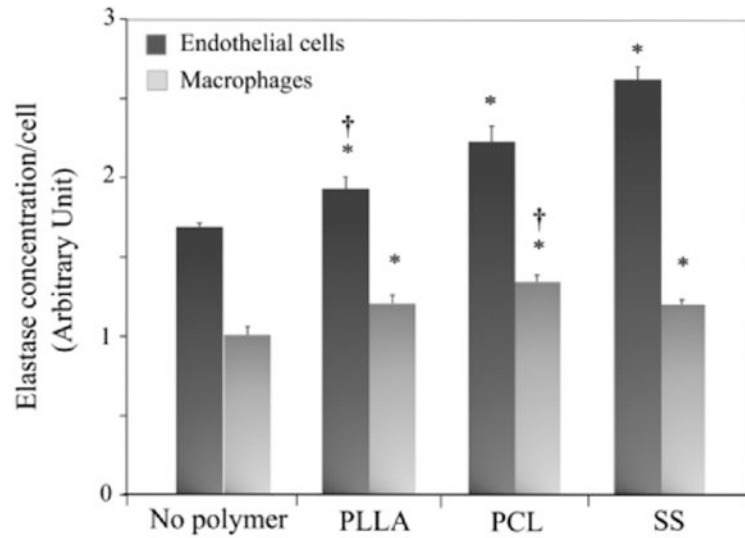


Figure 6.

Elastase production from cells after incubation with microparticles. Elastase production in cell culture media was determined by reacting with elastase substrate and comparing with the elastase standard at 415 nm. The absorbance value was normalized to the corresponding cell number measured by Hoechst staining. * $p < 0.05$ versus no polymer control; [†] $p < 0.05$ versus SS control.

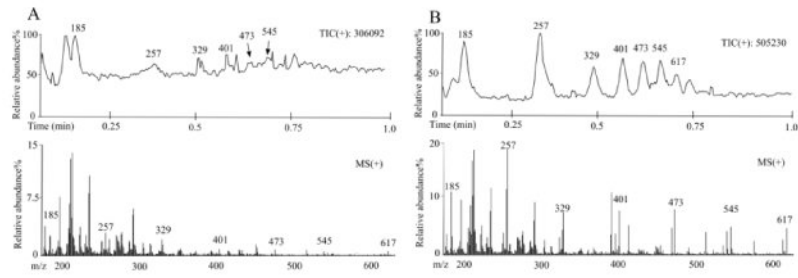


Figure 7.

Elastase-mediated polyester degradation. Mass spectrum of PLLA microparticle and degradation products. PLLA particles were suspended in DPBS without (a) or with (b) elastase and incubated at 37°C for 7 days. Degradation products in filtered solutions were confirmed by LC-MS with an acetonitrile-H₂O mobile phase system (positive ion mode).