Composite Dip Coating Improves Biocompatibility of Porous Metallic Scaffolds

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Abstract

Porous materials are becoming more common for bone implants, and it is increasingly important to find surface modification strategies that affect both the implant exterior and porous interior. In this study, selective laser melting (SLM) was used to create porous stainless steel implants 8 mm in diameter, which were subsequently dip coated with a composite polymethylmethacrylate-alumina (PMMA-Al₂O₃) film. Imaging with electron microscopy found evidence of the films at a depth of 2.2 mm into the porous implants, with dual-scale topography created by the native SLM stainless steel substrate and alumina nanoparticles. Energy dispersive X-ray spectroscopy confirmed the presence of the coating along the periphery of interior pores. In vitro tests with osteoblast-like cells showed greater cell metabolism on composite-coated samples compared to uncoated dense samples after seven days of culture.

Keywords

Dip coating, polymethylmethacrylate, additive manufacturing, porous materials, osseointegration, sol-gel preparation
1 Introduction

Porous metallic implant materials are becoming increasingly common for usage in bone and joint replacement due to their ability to mitigate stress-shielding effects[1]. The use of additive manufacturing enables effective scaffold design in bone implants by creating tunable mechanical performance[2] and channels for mass transport[3] by implementing porosity.

To encourage bone ingrowth, implants have been coated with composite materials, which can contain bioceramic particles such as calcium phosphate[4], rutile[5], and alumina[6]. One key limitation of conventional coating techniques, such as plasma spraying, is their line-of-sight processing[7], which prevents uniform coating deposition in the interior of porous structures, and their high-temperature processing. Changing the surface composition and topography in the interior of porous specimens is possible by electrochemical methods, such as micro-arc oxidation[8,9], where oxides of the base metal, calcium, or phosphorus are added to the surface. Composite materials have also been developed as a feedstock for additive manufacturing processes as a means of modifying the interior of porous structures compared to structures produced by traditional titanium feedstocks[10]. Recently, immersion techniques[11] and electrophoretic deposition[12] have been effective for depositing bioactive and organic coating materials on the interior of porous structures. Where alumina-based ceramic materials have been traditionally used for bulk implant components due to their excellent biocompatibility in bone applications[13], their integration in nanoparticle form on the interior of porous constructs has not been evaluated. Polymethylmethacrylate (PMMA), commonly known as a primary constituent in some bone cements, has recently been shown to have favourable osteogenic effects as an implant coating[14]. The potential for PMMA-Al₂O₃ organic composite material to coat the interior of porous metallic scaffolds remains unexplored.

The objective of this work was the development of a facile dip coating method to deposit a composite PMMA-alumina coating on the interior of an additively manufactured porous scaffold. Scanning electron microscopy (SEM) was used to confirm deposition, and the potential for improving osseointegration was explored with an *in vitro* cell metabolism assay.

2 Methods

2.1 Scaffold Production

Cylinders (*h* = 8 mm, *∅* = 8 mm) were designed in Autodesk Netfabb and hollowed to create body-centred cubic lattice struts rotated 45° about the X and Y axes. Fully dense implants as well as porous implants with strut diameters of 450 µm with a unit cell spacing of 1.2 mm, and therefore pores with an approximate throat diameter of 275 µm were formed. Scaffolds were built using 304L stainless-steel powder (< 45 µm, Carpenter Additive) using the selective laser melting (SLM) process (EOSINT M280, Germany). Laser power was set to 200 W, scan speed 800 mm/s, hatch spacing 80 µm, and layer thickness 40 µm during fabrication. Metal scaffolds were cleaned with ethanol and deionized water in an ultrasonic bath for 20 min. Scaffolds were either kept whole or sectioned longitudinally for subsequent coating.
2.2 Coating Deposition and Characterization

Polymethylmethacrylate (MW ~ 120,000, Millipore Sigma) and alumina (0.13µm, Al₂O₃, Baikowski) particles were obtained. Under continuous stirring, 10 g L⁻¹ PMMA was added to a mixed solvent containing 20% deionized water and 80% isopropanol (Millipore Sigma) and heated to 55°C, at which point the PMMA fully dissolved, only marginally increasing the viscosity. The PMMA solution was cooled back to room temperature, and Al₂O₃ particles were added to a concentration of 10 g L⁻¹. Metallic scaffolds were coated whole to produce specimens for cell viability assays or in halves for easier investigation of the midplane with electron microscopy. Scaffolds were attached to copper tape (see Figure S1), immersed into the PMMA-Al₂O₃ suspension under sonication for one minute, removed, and air dried at room temperature for 24 hours to evaporate solvent.

The mid-plane of the scaffold was sputter coated with carbon for conductivity, mounted to a SEM stub with silver paint and analyzed by SEM imaging (FEI Magellan 400) and energy dispersive X-ray spectroscopy (EDS) using an accelerating voltage of 10 kV.

2.3 Cell Viability

Saos-2 osteosarcoma cells (ATCC) were seeded on additively manufactured solid implants, porous implants, and PMMA-Al₂O₃ coated porous implants at a density of 10,000 cells/cm² in McCoy’s Modified 5A media. Cells were cultured at 37°C in an atmosphere of 5% CO₂ for seven days. After one, three, and seven days, a solution of 5% alamarBlue reagent in media (Life Technologies Inc.) was added to wells for 60 min. The dye was pipetted into a separate plate and fluorescence was measured at an excitation-emission wavelength of 540-580 nm. Statistical significance was determined using a two-way ANOVA in R with Tukey’s HSD test and p < 0.05.

3 Results and Discussion

3.1 Coating Deposition

Dip coating with the PMMA-Al₂O₃ composite produced a coating on both the exterior and interior of the scaffolds. Where PMMA bonds to the substrate may be governed by bidentate ligands, the mechanical strength of the composite coating on the stainless steel substrate is possibly increased compared to other coatings with monodentate bonding coordination. Higher magnification images of a representative pore (Figure 1A) illustrate the presence of the composite coating at an interior site, where non-uniform thickness is observed around the periphery of the pore. There is also evidence of sintered stainless steel particles on the lattice struts within the pores, adding elements of microscale topography on both interior and exterior lattice sites. Nanoparticles of Al₂O₃ were observed to be uniformly distributed through the PMMA matrix without aggregation (Figure 1B) at interior sites of the scaffolds. These nanoparticles, in conjunction with inherent striations in the PMMA matrix, add elements of nanoscale topography to interior and exterior sites in the scaffold, which can be favourable for osseointegration[15]. Coating deposition on the interior appears consistent with the exterior (Figure 1C), where Al₂O₃ is evenly distributed through the PMMA.
Figure 1: (A) Image of a representative pore on the interior of the scaffold. Thickness of PMMA-Al₂O₃ around the pore periphery is not constant. (B) Surface topography of PMMA-Al₂O₃ composite coating at the scaffold interior. (C) Surface topography of the PMMA-Al₂O₃ composite coating at the scaffold exterior. The Al₂O₃ distribution is comparable at scaffold exterior and interior sites.

EDS maps at interior scaffold sites (Figure 2A) on the cross-section are shown in Figure 2B, 2C, and 2D. Each site has a different distance to the scaffold exterior. Elemental signals characteristic to the uncoated stainless-steel (Fe, Ni) were uniform on the cross-sectional surface, while signals characteristic to the composite coating (C, Al) displayed higher intensity directly around the pore periphery at depths of 0.4 mm, 1.2 mm, and 2.2 mm.

Figure 2: (A) Cross-sectional SEM image with bright regions representative of the coating. Sampling sites at various distances to exterior correspond to EDS maps at depths of 0.4 mm (Row B), 1.2 mm (Row C) and 2.2 mm (row D). Increased intensity of aluminum and carbon at the pore periphery indicates complete penetration of the coating into the interior pores.

3.2 Cell Viability

The results of cell viability assays on the solid implants, porous implants, and PMMA-Al₂O₃ coated porous implants are shown in Figure 3. After seven days of culture, the PMMA-Al₂O₃ coated scaffolds significantly outperformed dense stainless-steel samples, confirming what has been shown previously on two-dimensional substrates[14]. Statistically higher cell metabolism
was also observed from day one to day seven on both the porous implants without coating and PMMA-Al₂O₃ coated porous implants. These results suggest that the addition of an interconnected porosity network with a diameter of 275 µm allows for rapid cellular proliferation through the porous interior of the scaffold relative to the control. The addition of PMMA-Al₂O₃ composite coatings on porous scaffolds further improves osseointegration potential relative to uncoated implants. By coating the SLM implant surface, surface topography, chemistry, and wettability are modified, and it is probable that metallic particle release is impeded. All of these factors can change the dynamics of cell-surface interaction, contributing to the resulting increase in cell viability seen here.

![Relative Fluorescence](image)

**Figure 3**: Saos-2 cell viability on solid, porous, and coated porous AM parts. Porous implants coated with the PMMA-Al₂O₃ composite coating showed significantly higher cell proliferation after seven days compared to solid components, while all porous implants showed significantly more cell proliferation after seven days.

### 4 Conclusions

PMMA-Al₂O₃ composite films were successfully deposited on the porous interior of SLM stainless steel implants using a facile dip coating method. SEM and EDS maps confirmed that the coating penetrated at least 2.2 mm into the scaffold interior, indicating that this immersion technique is suitable for coating the complete interior of porous metallic implants with a biomedical composite material. The alumina nanoparticles also contributed to a nanoscale topography around the pore periphery. In vitro characterization of the coated scaffolds with Saos-2 cells showed statistically higher rates of cell metabolism when compared to fully dense structures with the same geometry. Therefore, the dip coating method is a promising approach for creating composite coatings on porous implants to improve the potential osteoconductivity of their interior pores. The successful demonstration of Al₂O₃ nanoparticles, PMMA, and stainless steel scaffolds as pilot materials should lead to investigation of other additives and substrates, such as titanium.
alloys, for polymer-composite coatings on porous SLM implants. Future work should also evaluate the limits of PMMA-Al2O3 integration in larger-sized implants.

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6 References


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\textbf{Figure S1:} Metallic scaffolds (grey) were either (A) Sectioned longitudinally and mounted along their mid-plane to copper tape (red), or (B) Mounted whole by their baseplate, prior to dip coating in the PMMA- $\text{Al}_2\text{O}_3$ suspension.