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Surface modification of poly(l-lactide) and polycaprolactone biodegradable polymers using RF plasma discharge with sputter deposition of a hydroxyapatite target

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Surface modification of poly(l-lactide) (PLA) and polycaprolactone (PCL) biodegradable polymers by radio-frequency thermal glow discharge plasma is reported. Improved biocompatibility of PLA and PCL materials was obtained by employing hydroxyapatite target sputtering in Ar+ plasma as evidenced by the change of PLA and PCL properties from highly hydrophobic to hydrophilic, with absolute wettability obtained for both materials, and enhanced cell line EA-hy926 attachment to modified surfaces. Surface roughness and surface free energy were found to increase significantly for both polymers under prolonged plasma exposure and, a longer chain aliphatic PCL were found to display a marginally better post plasma-treated biocompatibility compared to PLA.

1. Introduction

Applications of biodegradable materials such as poly(l-lactide) (C3H4O2)n (PLA) and polycaprolactone (C6H10O2)n (PCL) in an environment where attachment and proliferation of living cells is important necessitate specific surface properties, namely, the surface roughness, to be within tens of nm (normally in 20–80 nm range and tailored to a specific cell line) and to be highly hydrophilic [1]. Although surface topology and physio-chemical properties have a major influence on the attachment of cells determining cell polarisation, cell morphology, spatial orientation of cell’s cytoskeleton components, the degree of order of intracellular transport and many other important parameters [2]. Recent experimental findings indicate that plasma surface treatment of PLA and PCL polymers increases surface roughness and reduces surface free energy and improves attachment of various biological cells on the surface [3]. However, despite a range of plasma treatment methods available (i.e. thermal and non-thermal plasma corona discharge, dielectric barrier discharge, etc.) inherent hydrophobicity and high-surface-free-energy properties of PLA and PCL are still poorly mend able in a controlled manner and hinder many applications [4,5]. While non-thermal plasma surface treatments are preferred for simplicity, modification of PLA and PCL under thermal plasma conditions remains less popular owing to inherent difficulties associated with identifying appropriate plasma conditions and complimenting target material(s) for a specific (bio-)material surface treatment.

In this Letter we report a pilot study on the effects of treatment of PLA and PCL polymers by a radio-frequency (RF) glow discharge plasma operated under Ar+ target sputtering mode to reduce hydrophobicity and surface-free energy towards improving living cell affinity. Hydroxyapatite (HA) Ca5(PO4)3(OH) was selected as a sputter target owing to HA natural (bio)-compatibility and ability to facilitate biological functions of various implant materials by plasma sputtering processes [6–8]. A dielectric HA serves as an ideal target material for an RF excited, plasma-assisted, sputtering of PLA and PCL to form a (bio-)ceramic/(bio-)resorbable polymer interface in support of live cell attachment studies.

2. Materials and methods

PURASORB® PL65 (Corbion Purac) with inherent viscosity (IV) at midpoint IVPLA=6.5 dl/g, and Polycaprolactone PURASORB®

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3. Results and discussions

Figs. 1 and 2 show the results of surface topology studies for stock L-PLA and PCL samples (see Fig. 1a for L-PLA, Fig. 2a for PCL), and for plasma-modified samples (see Fig. 1b–d for L-PLA, Fig. 2b–d for PCL); AFM topographic images, 2D fluorescent map of labelled EA-hy 926 cells; wettability images L-PLA and PCL with H2O and glycerol droplets. A summary of topological features for stock and plasma-treated L-PLA and PCL is given in Table 1, where changes for Rms, contact angles of water, θH2O, and glycerol, θgly, surface free energy, γ, including apolar, γsw, component accounting for Lifshitz/van der Waals (LW) type interactions, polar, γbl, component accounting for acid-base (AB) or donor-acceptor type interactions and total cell population are displayed as a function of the exposure time, t.

The obtained results indicate that the measured values for surface roughness gradually increase to a maximum of ~30 nm for L-PLA and ~70 nm for PCL with plasma exposure due to thermo-chemical degradation of the polymeric material mostly caused by increased presence of free H– bonds and plasma and target species bombardment. The latter are reactive, positively charged CaO+, Ca2++, PHO+, PO2+, P+ and negative charged (PO4)3− species sputtered out from the HA target which are attracted by the surface of L-PLA and PCL. The result of these interactions is the formation of an amorphous calcium aliphatic compound with significantly altered populations of carbonyl (–C=O), hydroxyl (–OH) functional groups and, calcium-phosphate pre-nucleation layer on L-PLA and PCL materials. Earlier, Boyd et al. confirmed the formation of the amorphous phase developing at film growth stages for HA sputtering on metal substrates [12].

We postulate that these processes occur simultaneously resulting in the formation of an amorphous (bio-)ceramic/(bio-)resorbable polymer interface such as a-Ca–(L-PLA)–O3 on L-PLA, and a-Ca–(PCL)–O3 on PCL material. Plasma treatment of both L-PLA and PCL materials gradually change their surface properties from hydrophobic, with surface energy values of ~20 mJ/m² for L-PLA and ~30 mJ/m² for PCL to largely hydrophilic, with a maximum values of γ of ~73 mJ/m² reported for both materials indicating absolute wettability. It is established that surface free energy of hydrocarbons CnH2n+2 is strongly related to C- saturation, therefore the increase of hydrophilic properties in L-PLA and PCL materials is be attributed to strong de-hydrogenation process occurring during the plasma treatment.
Moreover, since amorphous solids traditionally exhibit lower surface energy values than crystalline counterparts, we postulate that l-PLA and PCL surface properties undergo marked transition from an amorphous to a crystalline state further supporting the probability of formation of a (bio-)ceramic/(bio-)resorbable polymer interface. The marked increase of the probability of formation of a (bio-)ceramic/(bio-)resorbable microstructure relative to L-PLA to inherent differences in their molecular structure is beneficial to overall cell viability. Notably, a longer chain aliphatic PCL polymer was found to display a marginally better post plasma-treated biocompatibility compared to shorter-chain l-PLA.

We attribute a major increase of surface roughness in PCL relative to l-PLA to inherent differences in their molecular structure: l-PLA while lacking a longer C-C backbone is less predisposed to severe thermal and physio-chemical modifications than PCL counterpart, and as PCL exhibits lower solubility, it is naturally more prone to molecular cross-linking during processing. PCL therefore displays more uneven surface with the most pronounced functional groups with additional O– ions supplied by the sacrificial HA target.

The latter is evidenced by minimal changes of $r^{\text{O\text{-}}}$ component. The reasons are twofold: one is the presence of local reactive molecular domains participating in radical formation, molecular cross-linking and/or C=C bond formations on polymer surface under plasma irradiation, the other are the interactions with the positive (and neutral) species from the HA target noted earlier. High intermolecular force interactions on plasma treated l-PLA and PCL contribute towards the increase of C-O and O-C=O functional groups with additional O– ions supplied by the sacrificial HA target.

Cell population

- Plasmainduced surfaces and retrospectively indicates that the functional groups with additional O– ions supplied by the sacrificial HA target.

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Living cell culture experimental the data are presented in Table 1. It shows an increased affinity of EA-hy 926 cell attachment to plasma modified surfaces and retrospectively indicates that the increase of surface roughness and modification of surface functional groups is beneficial to overall cell viability. Notably, a longer chain aliphatic PCL polymer was found to display a marginally better post plasma-treated biocompatibility compared to shorter-chain l-PLA.

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4. Conclusion

In summary, we have shown that surface properties of L-PLA and PCL bioresorbable polymers can be successfully modified employing thermal glow discharge plasma sputtering hydroxyapatite target. Surface roughness, surface free energy could be altered by plasma treatment to achieve maximum wettability of L-PLA and PCL materials enhancing permanent cell line EA-hy 926 attachment. We postulate the formation of a (bio-)ceramic/(bio-)resorbable polymer interface as the result of plasma-surface interaction and suggest that the reported complimentary (bio-)target sputtering approach is not exclusive to L-PLA and PCL polymers, but can aid in modification of other bioresorbable materials.

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