

## Novel Top Coating Materials of Chemically Anchored Cytomimetic Thin films for Stent

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We have developed a new in-situ photopolymerization system which occurred at the interface of an acrylated phospholipid monolayer (1-palmitoyl-2-[12-(acryloyloxy)-dodecanoyl]-sn-glycero-3-phosphocholine) and an acrylated polymer substrate (acrylated poly (octadecyl acrylate-co-hydroxybutyl acrylate) using visible irradiation. The physicochemical properties of phospholipid thin films were confirmed using atomic force microscopy (AFM), scanning electron micrograph (SEM), water contact angles studies, and X-ray photoelectron spectroscopy (XPS). For the local drug delivery system for a potential antimetabolic agent, echinomycin release, in vitro, was substantially sustained from the polymerized phospholipid monolayer, due to the densely packed phospholipid molecules. Moreover, the polymerized phospholipid surfaces containing echinomycin substantially decreased smooth muscle cell proliferation, compared to the polymerized phospholipid surface without the drug. Finally, the blood compatibility of chemically anchored phospholipid thin films was evaluated using protein absorption and platelet adhesion, in vitro.