Lipid-coated silica nanoparticles for biomedical applications

COLL 33

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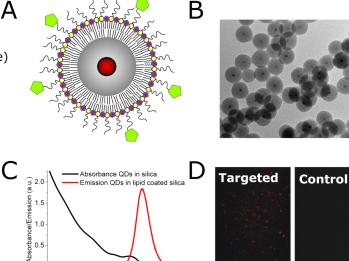
Silica particles as a nanoparticulate carrier material for contrast agents have received considerable attention the past few years, since the material holds great promise for biomedical applications. A key feature for successful application of this material in vivo is biocompatibility, which may be significantly improved by appropriate surface modification. In this study we report a novel strategy to coat silica particles with a dense monolayer of paramagnetic and PEGylated lipids. The silica nanoparticles carry a quantum dot in their centre and are made target-specific by the conjugation of multiple (alpha)v(beta)3-specific RGD-peptides. We demonstrate their specific uptake by endothelial cells in vitro using fluorescence microscopy, quantitative fluorescence imaging and magnetic resonance imaging. The lipid coated silica particles introduced here represent a new platform for nanoparticulate multimodality contrast agents.

(A) Schematic of the nanoparticle. Quantum dot (red) in silica (grey) incorporated in paramagnetic (red/blue) and pegylated (yellow) lipids. RGD peptides (green) were linked to PEG.

(B) TEM of quantum dots in 30 nm silica nanoparticles.

(C) Absorbance and Emission profile with emission peak at 626 nm.

(D) Endothelial cells incubated with $\alpha v\beta$ 3-targeted nanoparticles and control nanoparticles.



650 700

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400 450

500 550 600 Wavelength (nm)

0.0

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