Liposomes equipped with poly(N-isopropyl acryl amide)-containing coatings as potential drug carriers

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Table S1. Diameter and PDI of liposomes coated with PDA (L^D), PDA/pNiPAAm-NH₂ (L^{D/pNH_2}) or PDA/pNiPAAm-HB ($L^{D/pHB}$) for 30 min for L^D and 70 min for L^{D/pNH_2} and $L^{D/pHB}$ after dialysis.

Liposome	Diameter(STD) [nm]	PDI(STD)
LD	227(11)	0.29 (0.09)
LD/pNH2	297(37)	0.23(0.09)
L ^{D/pHB}	307(25)	0.18(0.03)

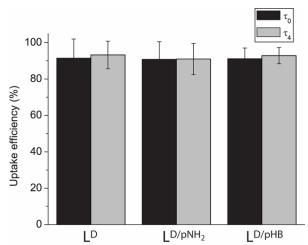


Figure S1. Interaction with macrophages considering shear stress: Uptake efficiency of L^D , L^{D/pNH_2} or $L^{D/pHB}$ by macrophages after 2.5 h at τ_0 or τ_4 .

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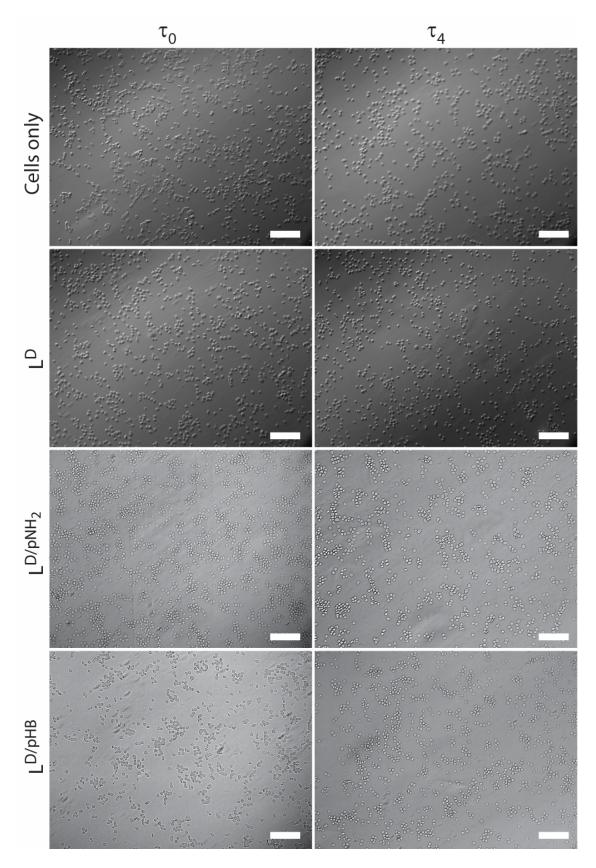


Figure S2. Overview bright field images of macrophages exposed to L^D , L^{D/pNH_2} or $L^{D/pHB}$ at τ_0 or τ_4 for 2.5 h. The scale bars are 200 μm .

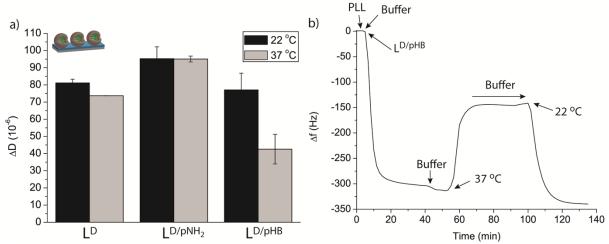


Figure S3. a) Adsorption of coated liposomes: Dissipation changes ΔD of PLL coated QCM-D crystals upon exposure to L^D , L^{D/pNH_2} or $L^{D/pHB}$ at 22 °C and 37 °C. b) QCM-D sample curve of crystals coated with PLL, $L^{D/pHB}$ at 22 °C followed by a temperature increase to 37 °C for 1 h constantly flowing buffer solution over the surface, and cooling down to 22 °C in the end. Since Δf before and after the temperature increase were found to be similar, the coated liposomes were stable adsorbed.

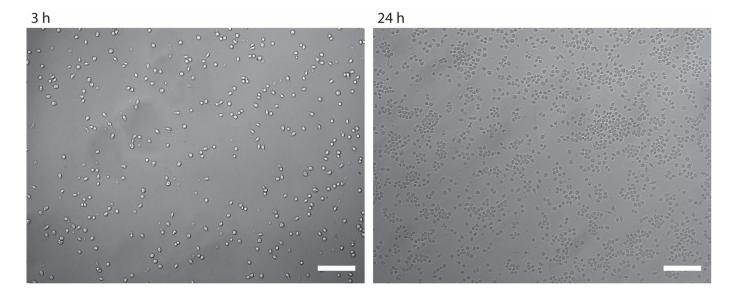


Figure S4. Overview bright field images of macrophages adhering to PDA coated surfaces for 3 h (top) and 24 h (bottom). The scale bars are $200 \mu m$.