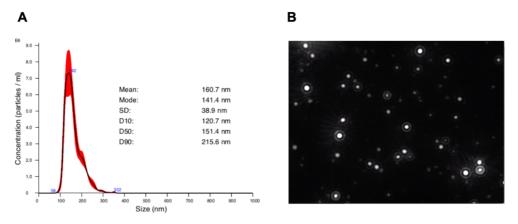
**Supporting information** 

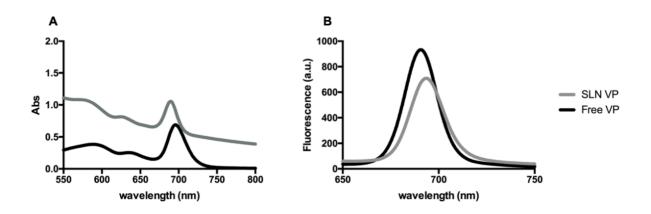
## Lipid nanoparticles as efficient verteporfin nanocarriers for photodynamic therapy of cancer

Tomás Mendes<sup>1</sup>, Andreia Granja<sup>1,\*</sup> and Salette Reis<sup>1</sup>.

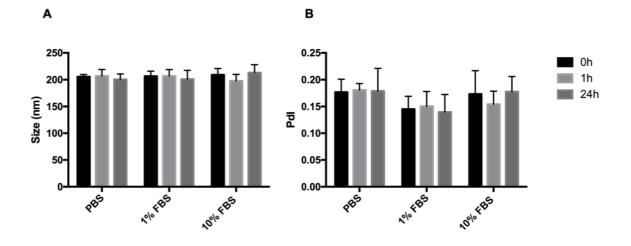
<sup>1</sup> LAQV, REQUIMTE, Departamento de Ciências Químicas, Faculdade de Farmácia, Universidade do Porto, R. Jorge de Viterbo Ferreira 228, 4050-313, Porto, Portugal.
\*corresponding author: aagranja@ff.up.pt



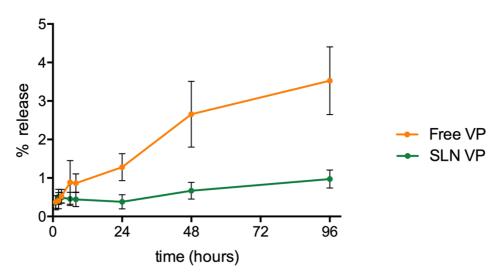
**Figure S1.** Particle size determinations obtained by NTA. Particle size distribution graph (**A**) and corresponding frame from the NTA captured video (**B**).



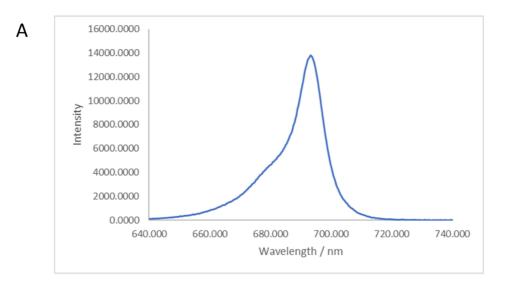
**Figure S2.** Absorption spectra (**A**) and fluorescence emission spectra (excitation at 581 nm) (**B**) of free VP and SLN VP (16  $\mu$ M of VP)



**Figure S3.** Nanoparticle colloidal stability into different biological relevant media. Nanoparticles were incubated with PBS, PBS with 1% FBS and PBS with 10% FBS for 1 h and 24 h at 37 °C and particle size and PdI were analyzed using a ZetaPALS Analyzer (Brookhaven Instruments Corporation; Software: Particle Sizing v.5 Brookhaven Instruments; Holtsville, NY, USA).



**Figure S4.** Cumulative release of VP (free and SLN-loaded) in PBS pH 7.4 (5% v/v of Tween 80). Data are expressed as mean ± SD (n=3)





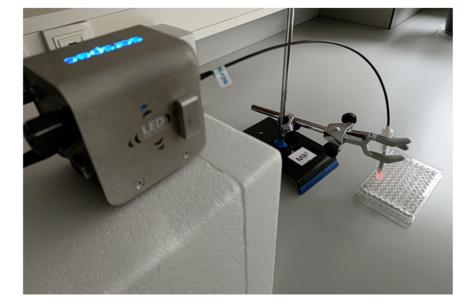


Figure S5. LED light source emission spectrum (A) and setup for the LED irradiation experiments (B)