

# Mapping the L-tryptophan Capped Copper Nanocluster Mediated Binding and Targeted pH-Responsive Release of Doxorubicin via Fluorescence Resonance Energy Transfer (FRET)

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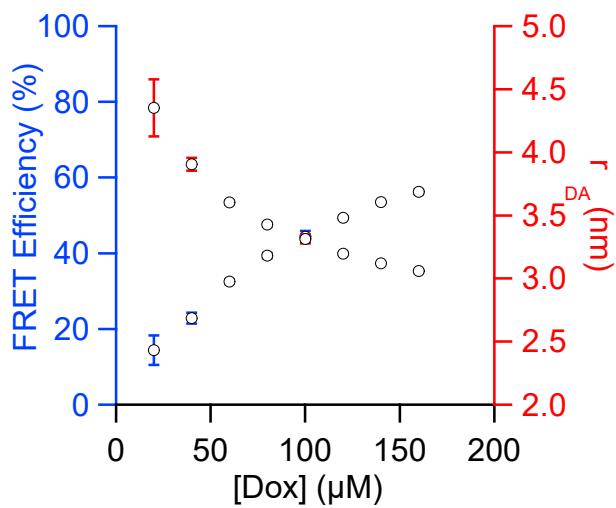
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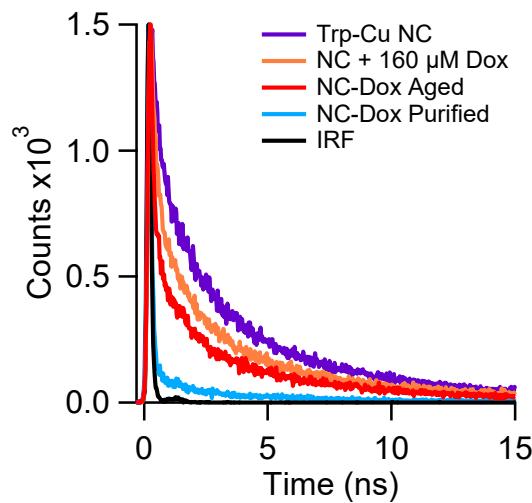
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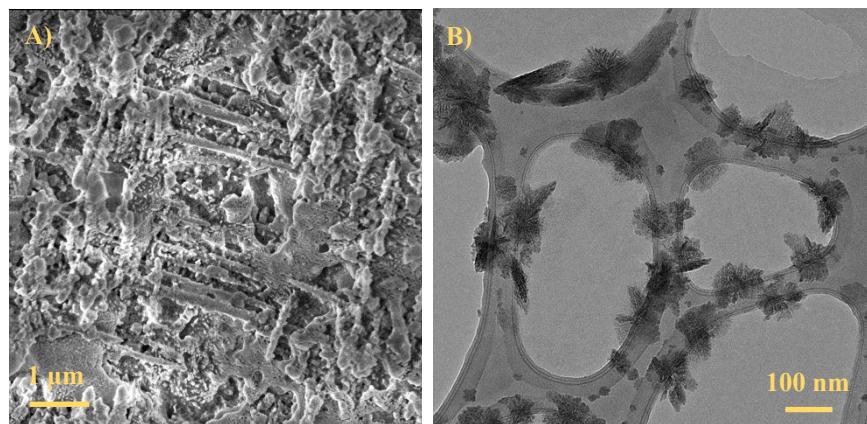
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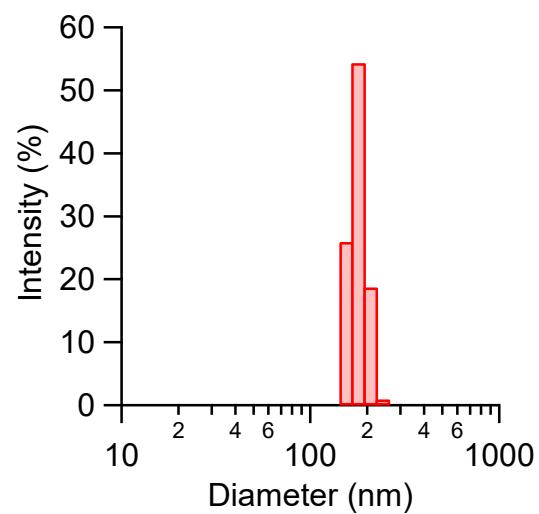
**Figure S1:** Changes in the FRET efficiency (blue) and donor-acceptor distance ( $r_{DA}$ ) (red) at increasing concentrations of Dox calculated from time-resolved FRET measurements.



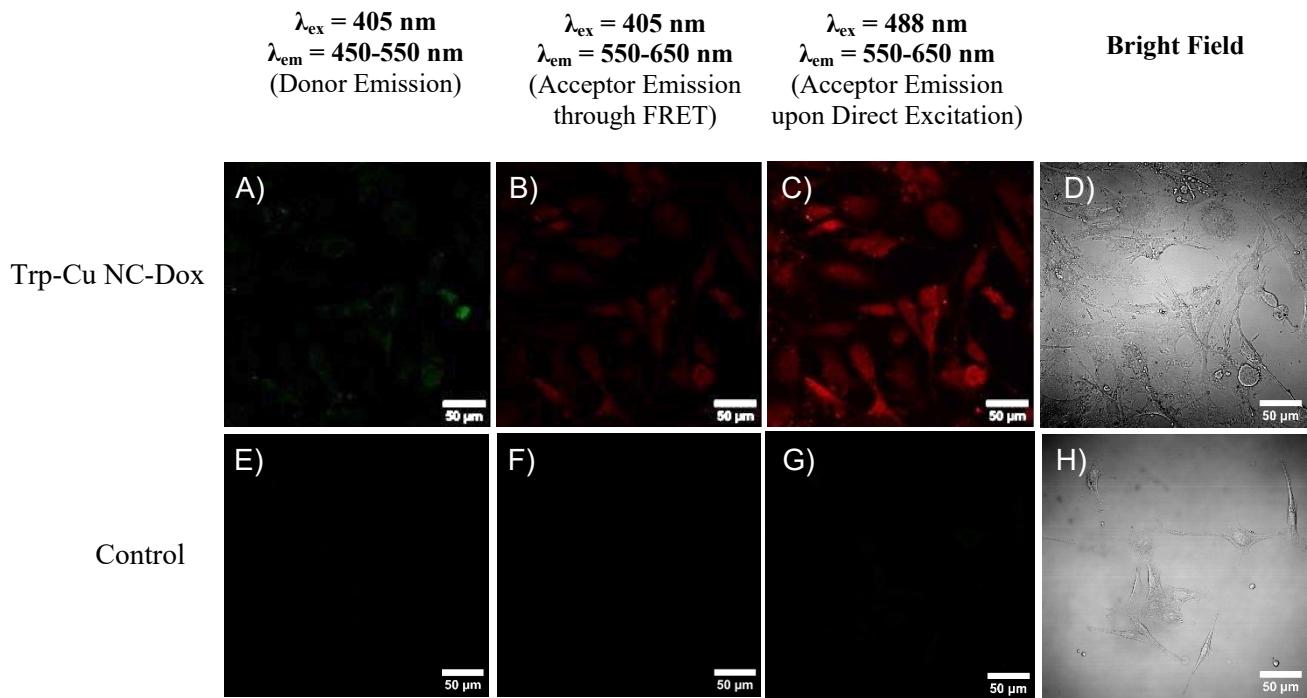
**Figure S2:** Time-resolved PL intensity decay curve ( $\lambda_{ex} = 405$  nm,  $\lambda_{em} = 480$  nm) of pure Trp-Cu NC (violet), Trp-Cu NC-Dox nano-conjugate before (orange) and after aging (red). The lifetime further decreased upon purification of the aged sample (cyan). The black curve represents the instrument response function (IRF).



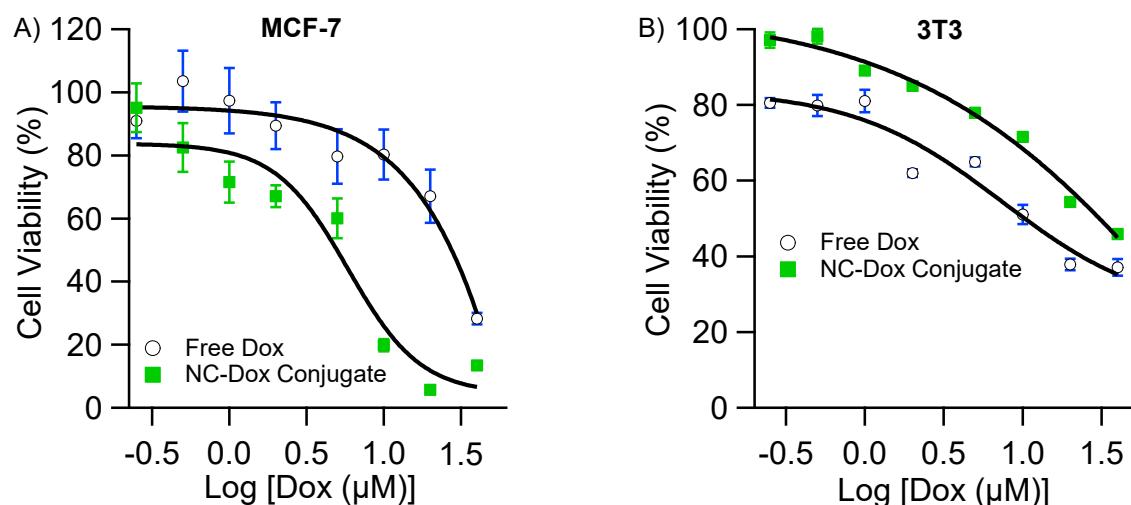
**Figure S3:** (A) FESEM image at low magnification (scale bar: 1  $\mu$ m) of purified Trp-Cu NC-Dox nanoconjugates. (B) HRTEM image at low magnification (scale bar: 100 nm) of purified Trp-Cu NC-Dox nanoconjugates.



**Figure S4:** Dynamic light scattering (DLS) spectrum of Trp-Cu NC-Dox nanocnjugate.



**Figure S5:** Confocal microscope images of 3T3 cells incubated (A-D) with and (E-H) without Trp-Cu NC-Dox nanoconjugate.



**Figure S6:** Cell viability (%) was plotted against the log of Dox concentration for free Dox (blue) and the Trp-Cu NC-Dox nanoconjugate (green) in (A) breast cancer (MCF-7) cells and (B) normal mouse fibroblast (3T3) cells. A sigmoidal fit is represented by the black line.