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Supplementary Information



Figure S1. EPR spectra of T2C₁₂, T2C₁₂-T80₁₀ and T2C₁₂-T80 (all at 100 μ M) in micellar form in HEPES-Buffered saline solution at pH 7.4.



Figure S2. Field Emission Scanning Electron Microscopy (FESEM) image of $T2C_{12}$ -T80 100 μ M acquired with a Magnification of 323k x, 5 keV of energy and 100 pA of probe current using a Tescan FEG-SEM S9000.

Measurements are acquired with a Schotty emitter, the probe set at 100 pA and the electron beam energy at 5 keV. The analysis was performed with an in-beam SE detector. Microanalysis was performed using OXFORD - Ultim Max detector - AZTEC software.



Figure S3. Zeta-Potential graphs of $T2C_{12}$ -T80₁₀ 100 μ M (A) and $T2C_{12}$ -T80 100 μ M (B) measured by DLS.



Figure S4. Calibration curve obtained using different concentrations of 4-Oxo-TEMPO in Hepes-Buffered saline solution. The measurements were fitted with a linear fit function ($y = a + b^*x$) using OriginPro 2018.



Figure S5. Residual nitroxide radical (%) of T2C₁₂-T80 micelle (100 μ M) in the presence of Ascorbic Acid 1mM at room temperature over 120 min.



Figure S6. EPR spectra of $T2C_{12}$ -T80 100 μ M after 3 hours of incubation with 3T3, Hs766T, and HepG2 cells lysates obtained from 250 x 10³ cells each.



Figure S7. EPR Intensity of $T2C_{12}$ -T80 100 μ M incubated, at 37°C and under stirring at 400 rpm, with cytosolic extracts of 3T3, Hs766T, and HepG2 cells, obtained from 250 x 10³ cells, with and without the addition of BNPP 600 μ M.



Figure S8. EPR spectra of Hs766T cells pellet incubated with 4-Oxo-TEMPO 100 μ M and T2C₁₂-T80 100 μ M for 24 hours.



Figure S9. Vitality percentage (measured by MTT assay) of Hs766T and 3T3 cell lines, measured after 24 h of incubation at different concentrations of $T2C_{12}$ (left) and $T2C_{12}$ -T80 (right). Graphs show the mean ± SD of % vitality evaluated on 3 independent experiments.