

Homologous polydopamine ameliorates haemolysis of melittin for enhancing its anticancer efficacy

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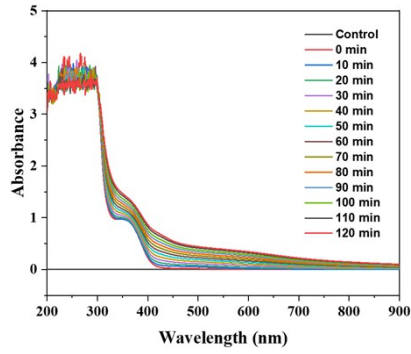


Fig. S1 UV-vis absorbance of MPF NPs solution at different reaction time points.

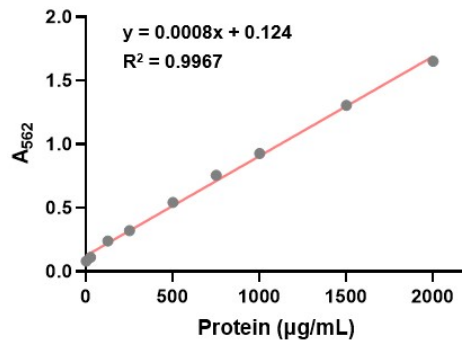


Fig. S2 The standard curve for MEL using BCA quantitative analysis method.

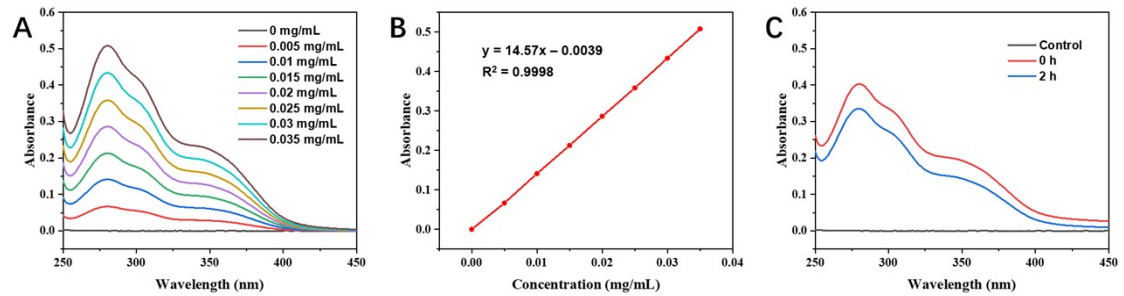


Fig. S3 (A) UV-vis absorbance of FA at different concentrations, and (B) the corresponding concentration standard curve. (C) UV-vis absorbance changes of FA in the reaction supernatant before and after the formation of MPF NPs.

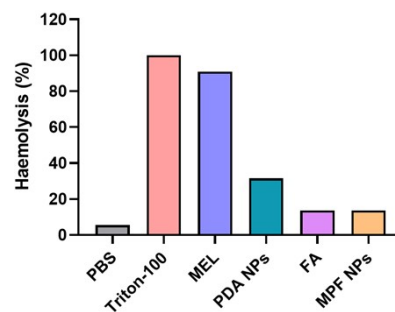


Fig. S4 Haemolysis analysis of different drugs, positive control (Triton-100), and

negative control (PBS).

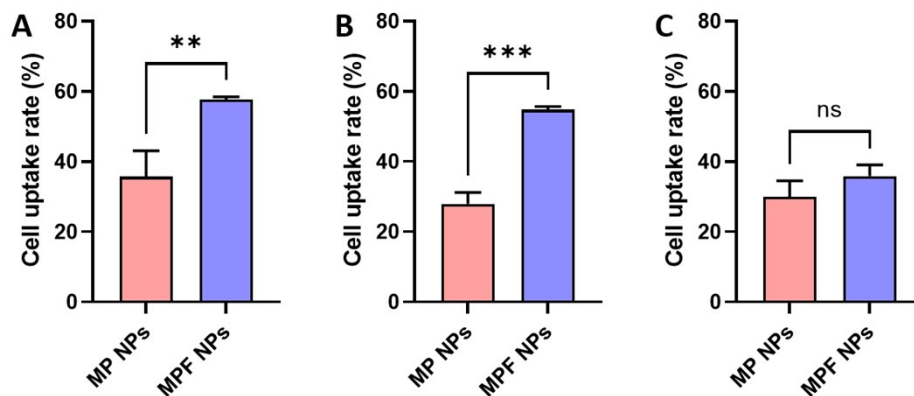


Fig. S5 Cell uptake rate detection of MP NPs and MPF NPs in 4T1, CT26 and HepG2 cells respectively.

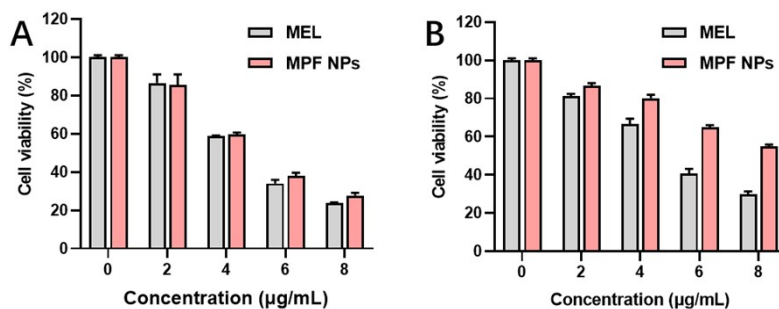


Fig. S6 Cell viability analysis of (A) CT26 and (B) HepG2 cells after treated with MEL and MPF NPs.

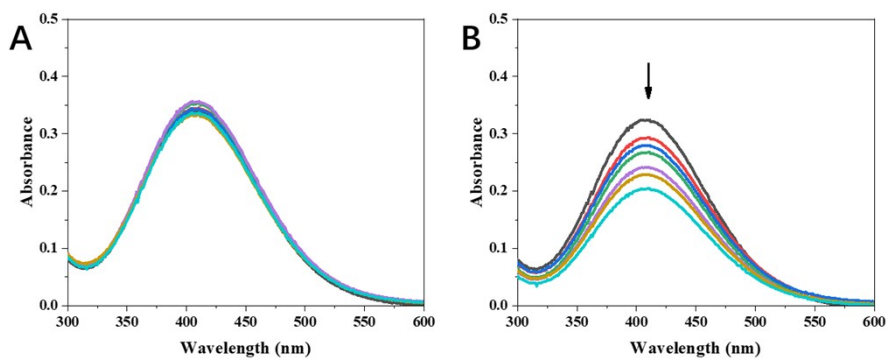


Fig. S7 The UV-vis absorbance changes of hydrogen peroxide probe at 415 nm after treated with (A) distilled water and (B) MPF NPs.

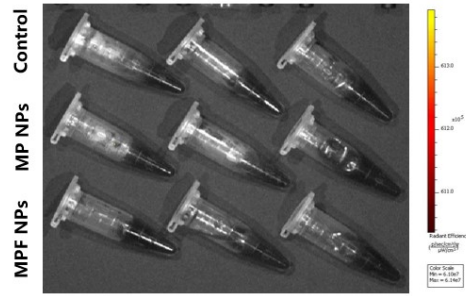
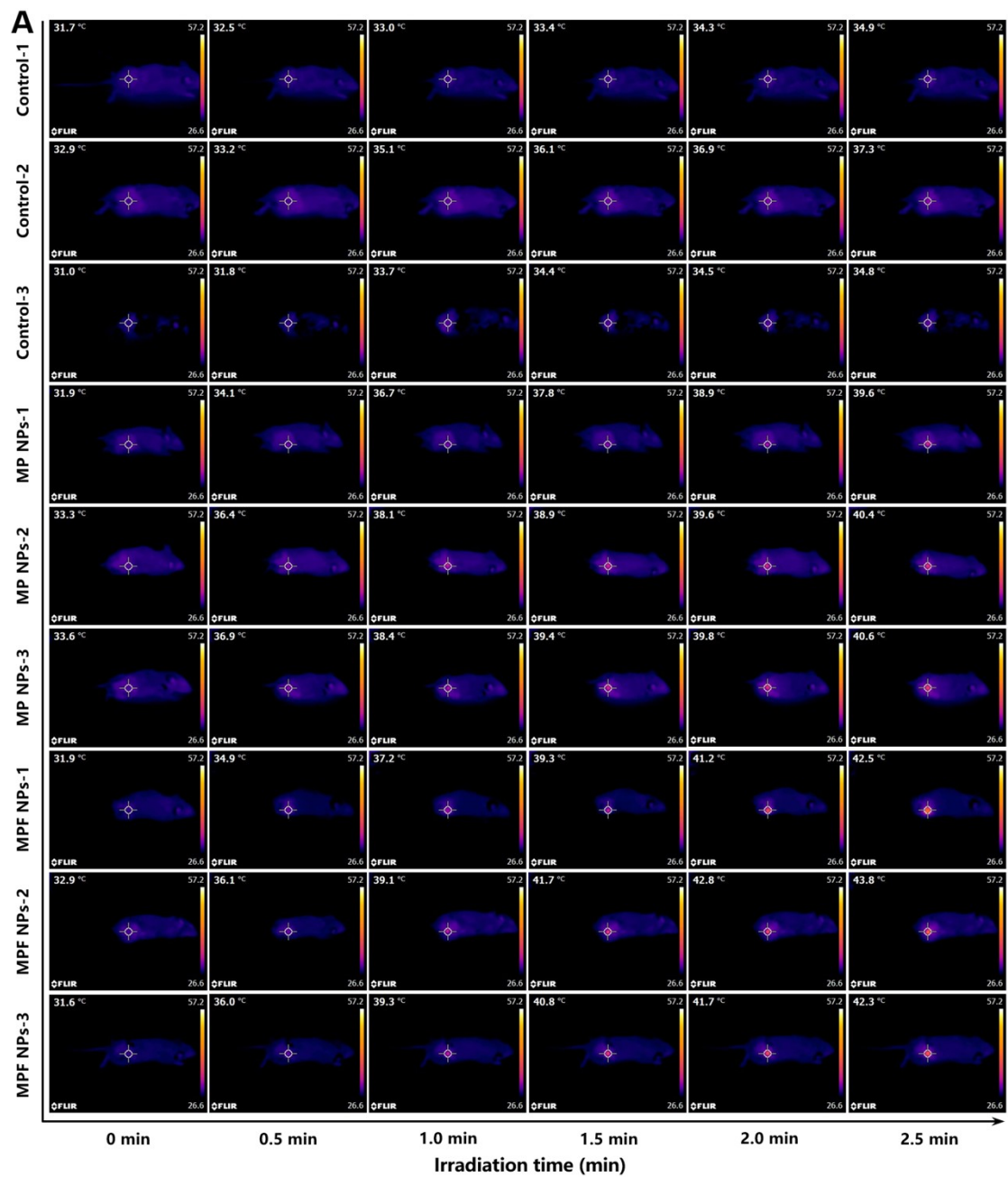


Fig. S8 Fluorescent distribution imaging of blood obtained from 4T1-bearing mice after different drugs treatment.



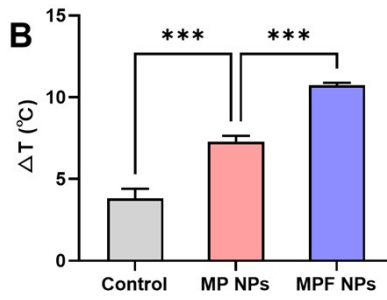


Fig. S9 (A) Photothermal imaging of MP NPs and MPF NPs in tumour-bearing mice under laser irradiation (808 nm, 0.5 W cm⁻²) and (B) corresponding temperature changes comparison.

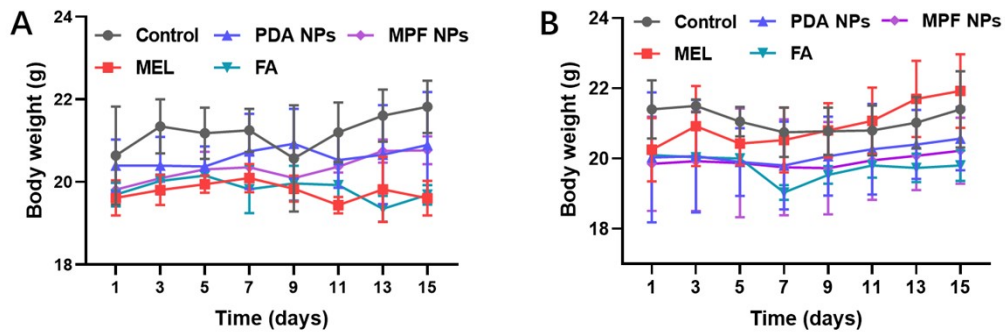


Fig. S10 Body weight changes of (A) CT26-bearing mice and (B) HepG2-bearing mice during 14 days' treatment.