

Supporting Information for

Synthesis and study of amorphous calcium phosphate dual-targeted drug-carrying platforms

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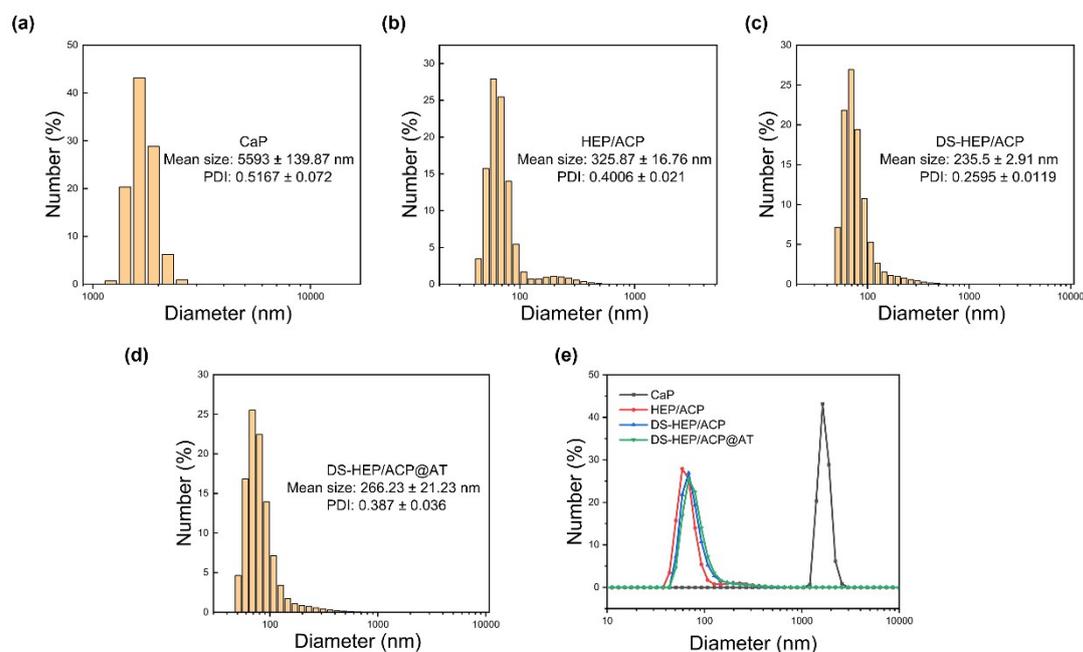


Fig. S1 Particle size distribution based on the number (a-d) and comparison graphs (e) of CaP, HEP/ACP, DS-HEP/ACP and DS-HEP/ACP@AT. Data are presented as the mean ± SD (n = 3)

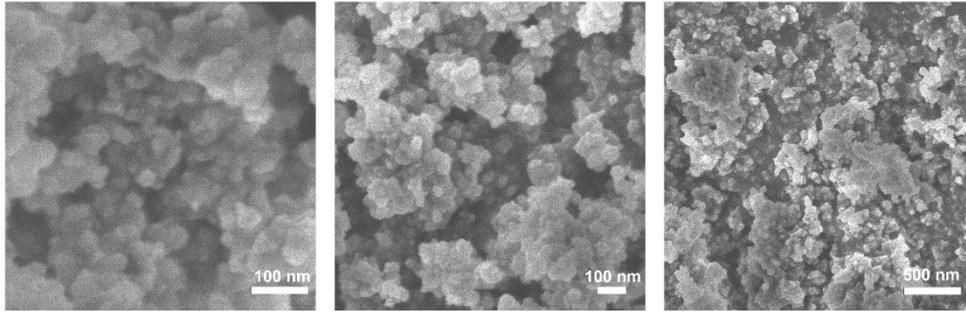


Fig.S2 SEM images of DS-HEP/ACP at different magnifications.

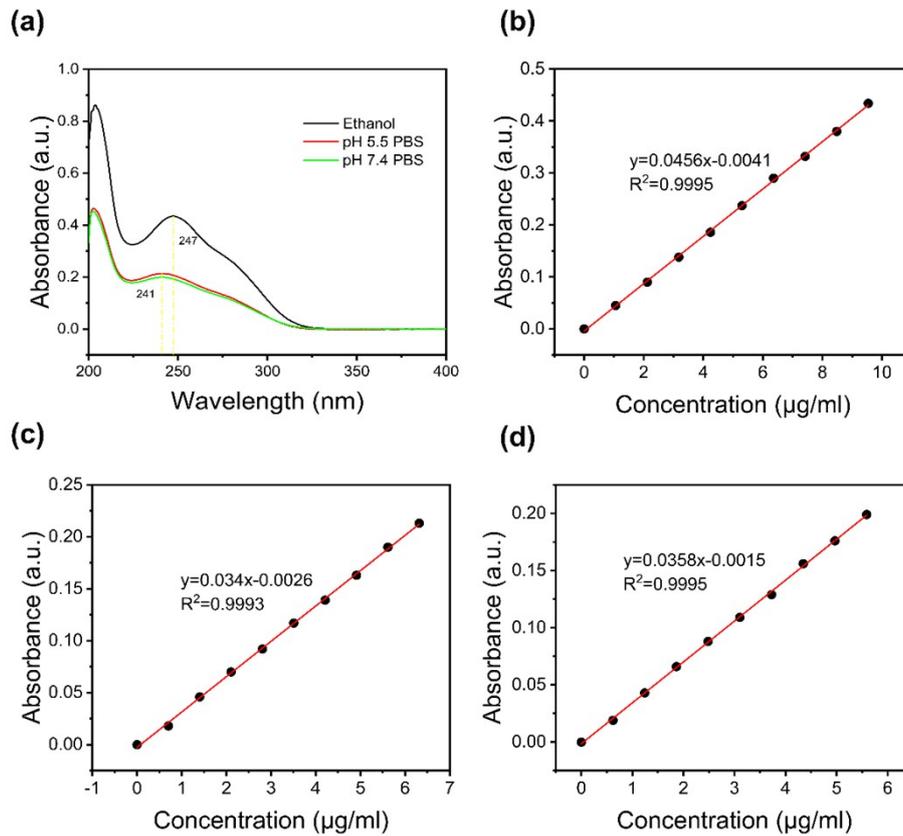


Fig.S3 (a) UV-vis absorption spectra of AT under different solvents, the next largest absorption peaks appeared at 247 nm and 241 nm for ethanol and PBS as solvents, respectively. The standard curve between absorption intensity and concentrations of AT for ethanol (b), PBS at pH 5.5 (c), and PBS at pH 7.4 (d) as solvent, respectively.

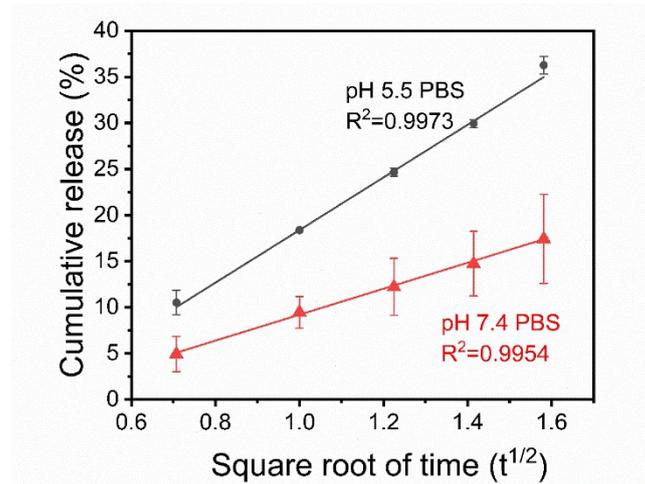


Fig.S4 The cumulative amount of released drug versus the square root of release time for the DS-HEP/ACP drug delivery system in both PBS solutions with pH 5.5 and pH 7.4 in the first 2.5 h at 37 °C. Data are presented as mean ± SD (n ≥ 3).

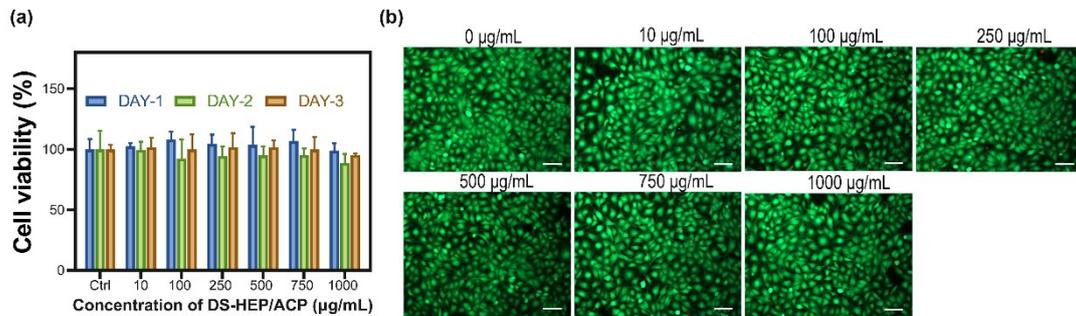


Fig.S5 (a) Cytotoxicity test results of HUVECs cocultured with different concentrations of DS-HEP/ACP for 1, 2 and 3 days. (b) Live/dead staining results of HUVECs cocultured with different concentrations of DS-HEP/ACP for 1 day. Scale bar: 100 µm. Data in (a) are mean ± SD (n ≥ 3).

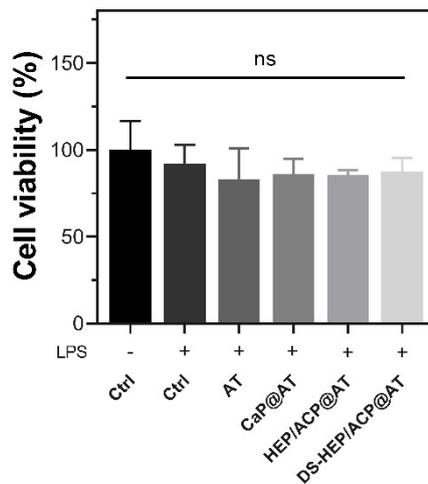


Fig.S6 Cytotoxicity test results of RAW264.7 cocultured with different drug-encapsulated samples for 1 day. Data are presented as mean ± SD (n ≥ 3).

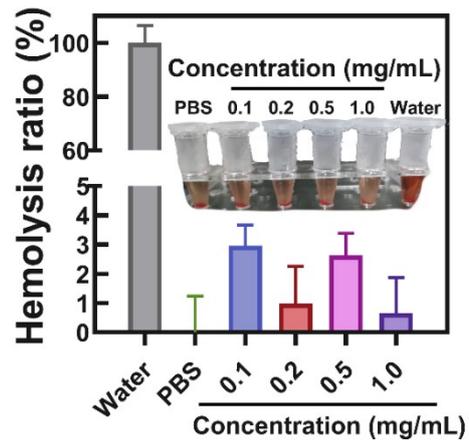


Fig.S7 Hemolysis analysis of blood samples after exposure to various concentrations of DS-HEP/ACP. Data are presented as mean \pm SD (n=3).