Electronic Supplementary Information

A peptide-decorated and curcumin-loaded mesoporous silica nanomedicine for effectively overcoming multidrug resistance in cancer cell

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| Sample | zeta potential (mV) |
|---------------------|---------------------|
| MSN-41 | -18.42 |
| MSN-NH ₂ | 21.15 |
| MSN-alkyne | 2.12 |
| MSN-Pep | 12.43 |
| DOX/CUR@MSN-Pep | 12.05 |

Table S1. Zeta potentials of different nanocarriers in PBS



Figure S1. FT-IR spectra of MSN, MSNs-NH₂, MSNs-alkyne, and MSN-Pep.



Figure S2. UV-VIS absorption spectra of MSNs, MSNs–NH₂, MSNs–alkyne, and MSN-Pep.



Figure S3. BET nitrogen adsorption/desorption isotherms (A) and BJH pore size distribution (B) of MSN and MSN-Pep.



Figure S4. (A) TEM image of blank MSN. Scale bar: 20 nm. (B) Size distribution ofblankMSNmeasuredbyDLS.



Figure S5. (A) The absorption spectra of DOX with different concentrations. (B) The standard curve of DOX absorbance value at 488 nm. The obtained standard curve is y=0.01902x+0.00106 (y: absorbance value at 488 nm; x: concentration of DOX, $R^2=0.99879$). (C) The absorption spectra of CUR with different concentrations. (D) The standard curve of CUR absorbance value at 425 nm. The obtained standard curve is y=0.02389x+0.00586 (y: absorbance value at 425 nm; x: concentration of DOX, $R^2=0.99767$).



Figure S6. Confocal fluorescence images of MCF-7 cells incubated with free DOX, DOX+CUR, DOX@MSN-Pep and DOX/CUR@MSN-Pep for 4 h. Scale bar: 10 μm.



Figure S7. Confocal fluorescence images of MCF-7/ADR cells incubated with DOX/CUR@MSN-Pep (without RGDS), DOX/CUR@MSN-Pep for 4 h and the block group pretreated with excessive free RGD, followed by incubation with DOX/CUR@MSN-Pep for 4 h. Scale bar: 10 µm.



Figure S8. Confocal fluorescence images of MCF-7/ADR cells stained with LysoTracker Deep Red (green) and Hoechst 33342 (blue) after cells were incubated DOX/CUR@MSN-Pep for 1, 2, 4, 6, 12 and 24 h, respectively. Scale bar: 10 μm.



Figure S9. MTT assays of MCF-7 (A) and MCF-7/ADR (B) cells after incubation with CUR. The IC₅₀ values for MCF-7 and MCF-7/ADR cells are measured to be 22.93 μ M and 41.68 μ M, respectively. Data are means ± SD (n = 5).



Figure S10. MTT assays of MCF-7 and MCF-7/ADR cells after exposure to (A) blank MSN, (B) MSN–NH₂, (C) MSNs–alkyne and (D) MSN-Pep nanocarrier at various concentrations. Data are means \pm SD (n = 5).