

Supporting Information

ROS-responsive multifunctional DCS-based micelle-hydrogel for simultaneously inhibiting post-melanoma-resection recurrence and promoting wound healing

Sha Liu,^{*a} Lina Wang,^c Yupeng Wang,^{*b} and Lina Geng^{*a}

^a Hebei Key Laboratory of Inorganic Nanomaterials, College of Chemistry and Material Science, Hebei Normal University, Shijiazhuang 050024, P. R. China.

^b Hebei Key Laboratory of Organic Functional Molecules, College of Chemistry and Material Science, Hebei Normal University, Shijiazhuang 050024, P. R. China.

^c Testing and Analysis Center, Hebei Normal University, Shijiazhuang 050024, P. R. China.

* Corresponding author.

Email: genglina1@hebtu.edu.cn, yupengwang@hebtu.edu.cn, liusha@hebtu.edu.cn

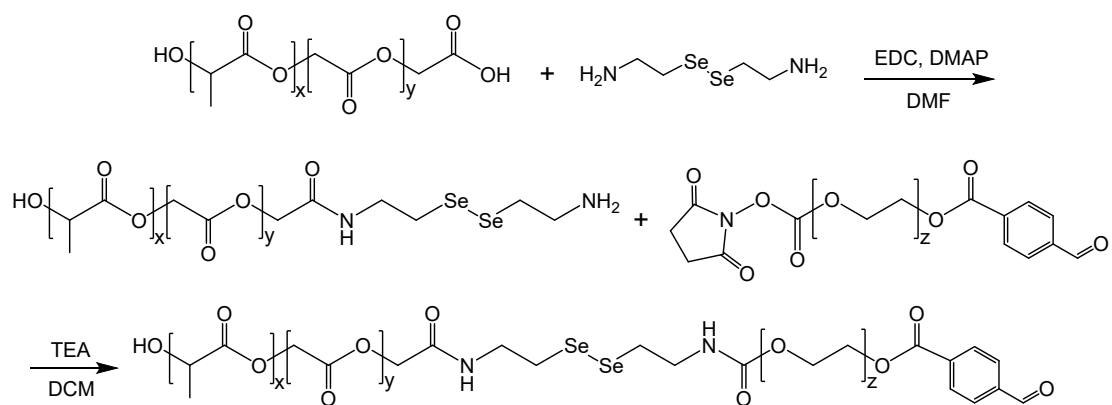


Fig. S1 Synthesis route of PLGA-Se-Se-PEG-CHO.

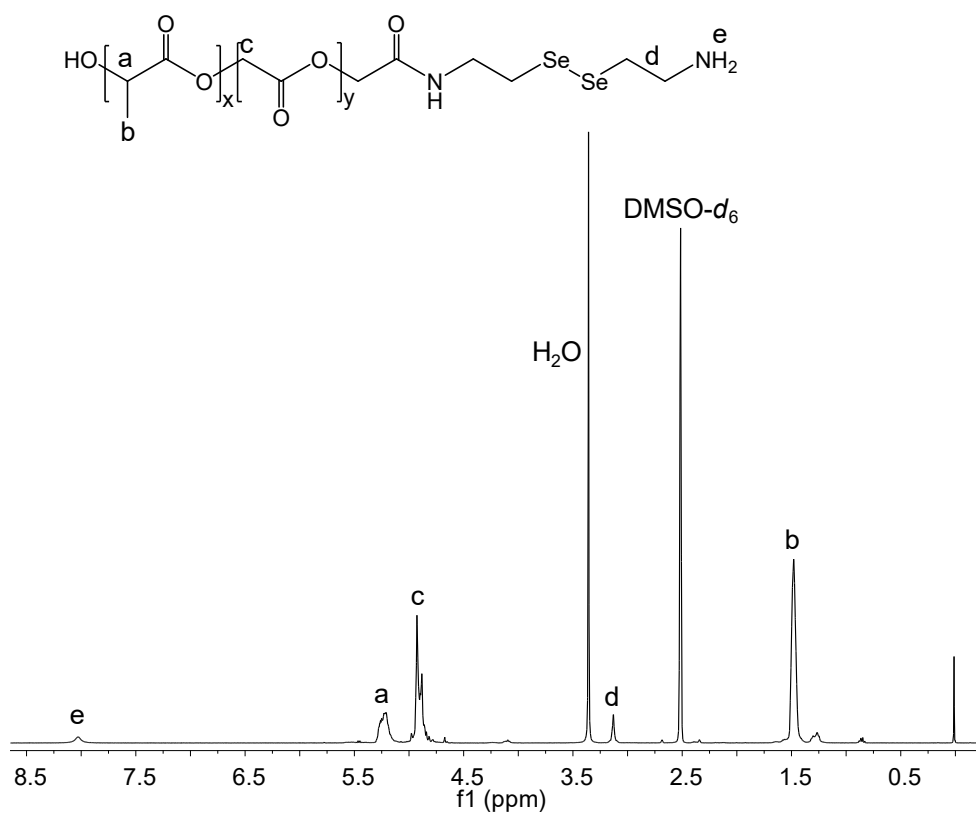


Fig. S2 ^1H NMR spectra of PLGA-Se-Se- NH_2 in $\text{DMSO-}d_6$. 8.20 (e, NH_2), 5.24 (a, CH), 4.93 (c, CH_2), 3.15 (d, CH_2CH_2), 1.49 (b, CH_3).

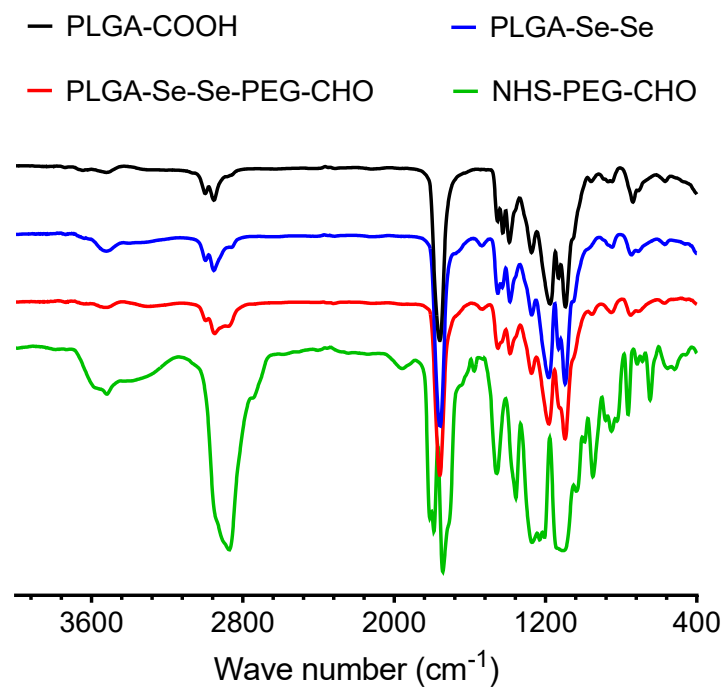


Fig. S3 FTIR spectra of PLGA-COOH, PLGA-Se-Se, NHS-PEG-CHO and PLGA-Se-Se-PEG-CHO.

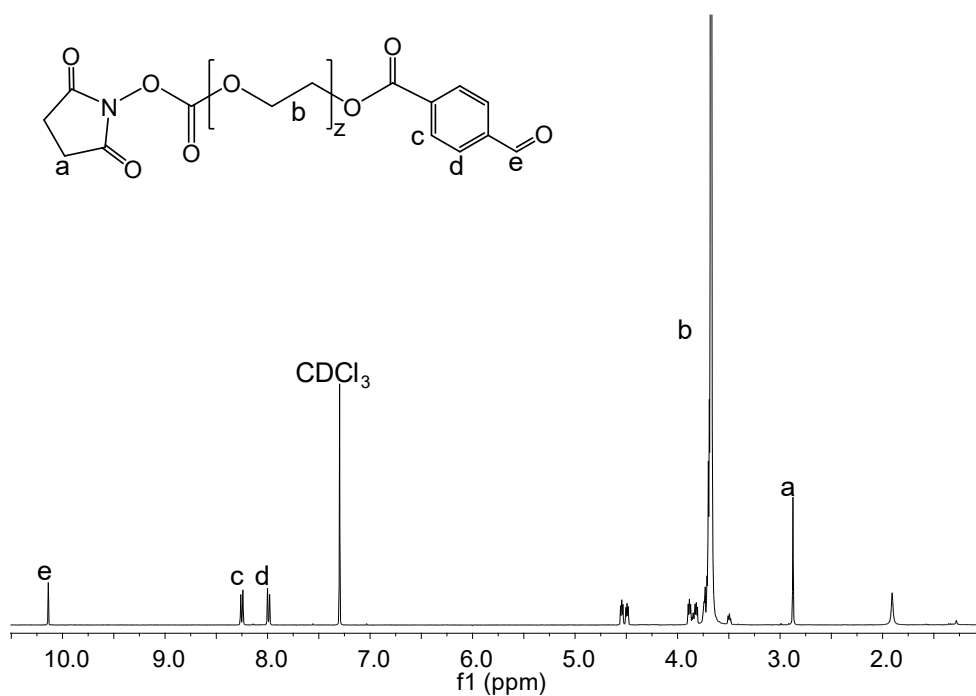


Fig. S4 ^1H NMR spectra of NHS-PEG-CHO in CDCl_3 . 10.14 (e, CHO), 8.26(c, CH), 8.00 (d, CH), 3.66 (b, CH₂CH₂), 2.87 (a, CH₂).

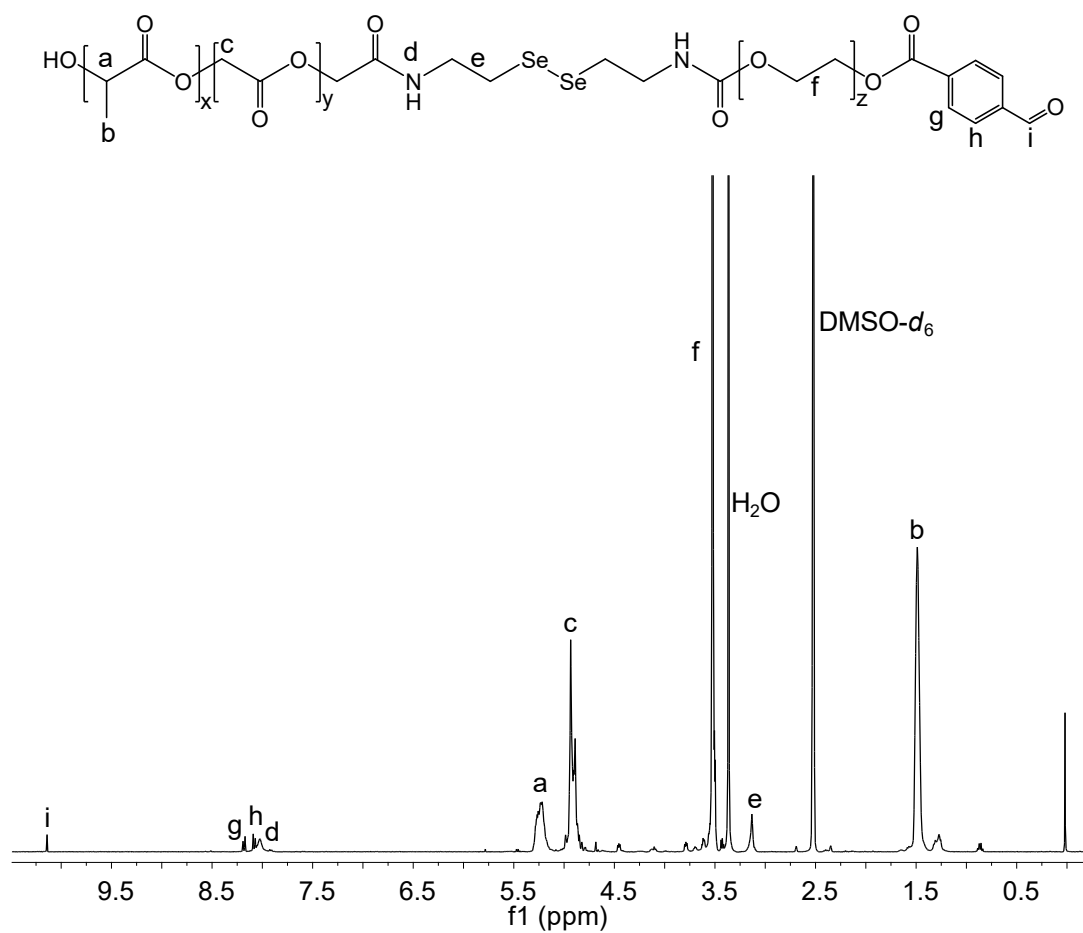


Fig. S5 ^1H NMR spectra of PLGA-Se-Se-PEG-CHO in DMSO- d_6 . 10.14 (i, CHO), 8.20 (g, CH), 8.09 (h, CH), 8.03 (d, NH), 5.24 (a, CH), 4.93 (c, CH₂), 3.52 (f, CH₂CH₂), 3.13 (e, CH₂CH₂), 1.49 (b, CH₃).

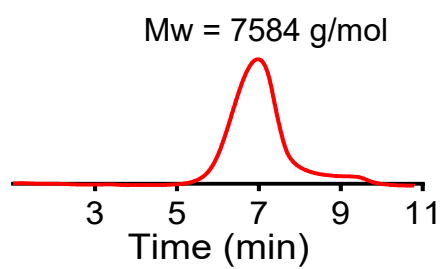


Fig. S6 GPC profiles of PLGA-Se-Se-PEG-CHO.

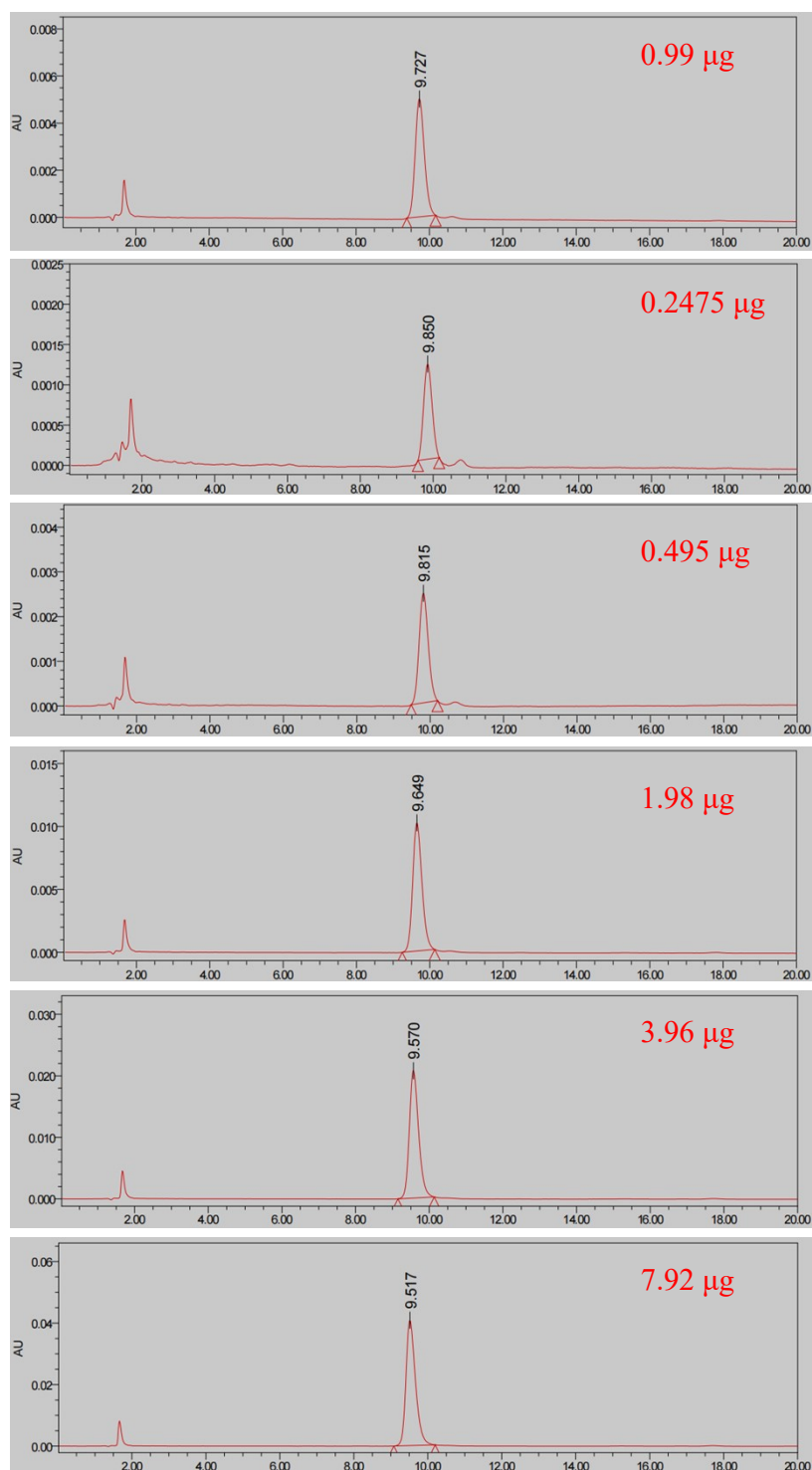


Fig. S7 The HPLC curve of different contents of BMS-202.

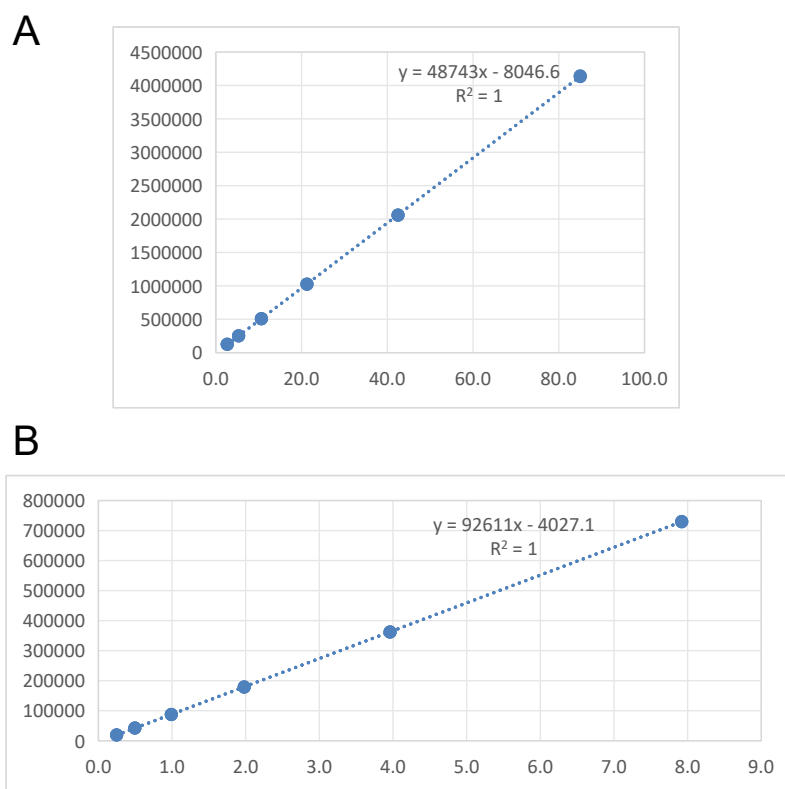


Fig. S8 (A) The standard curve of apigenin concentration. (B) The standard curve of BMS-202 concentration.

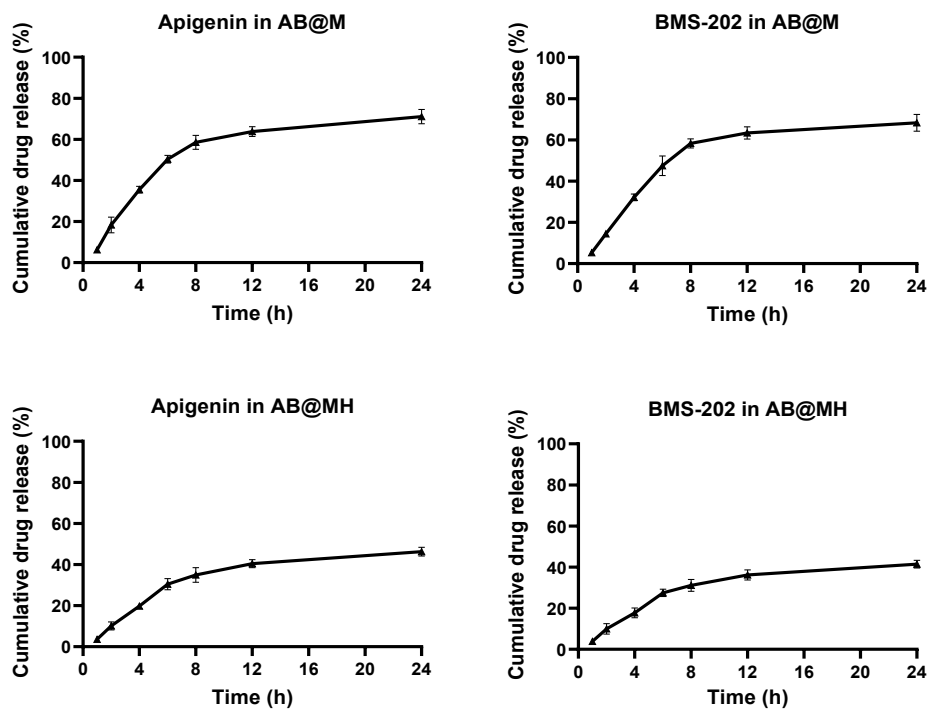


Fig. S9 Release profiles of apigenin or BMS-202 from AB@M or AB@MH in 1 mM H₂O₂.

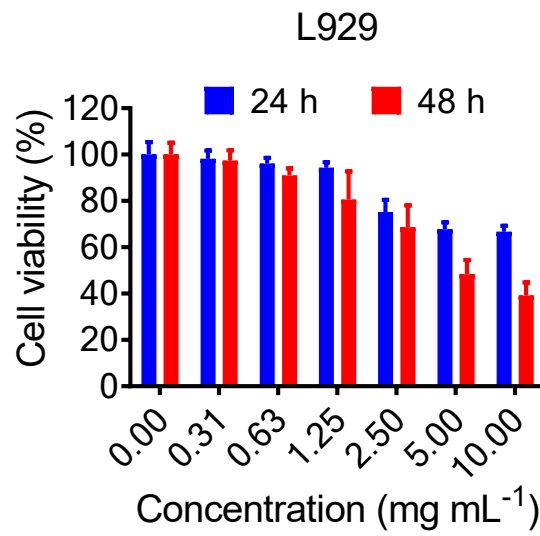


Fig. S10 L929 cell viability after treating with different concentrations of AB@M for 24 h and 48 h, respectively.

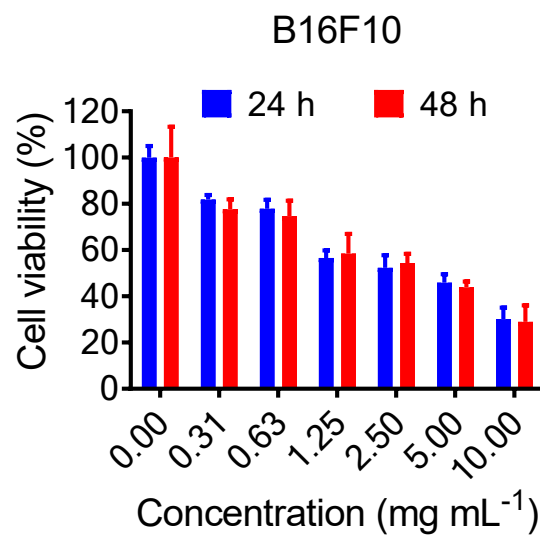


Fig. S11 B16F10 cell viability after treating with different concentrations of AB@M for 24 h and 48 h, respectively.

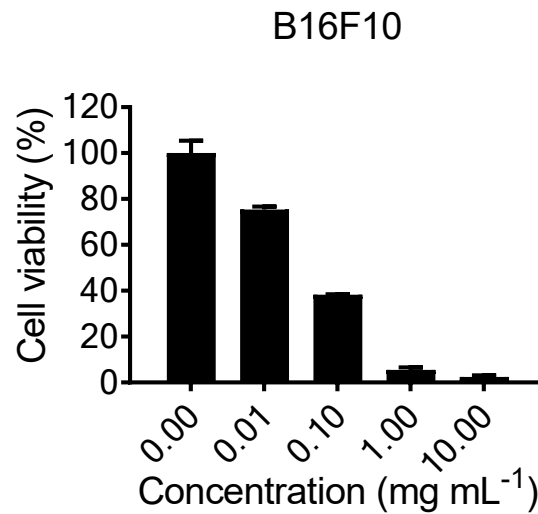


Fig. S12 B16F10 cell viability after treating with different concentrations of Triton X-100 for 24 h.

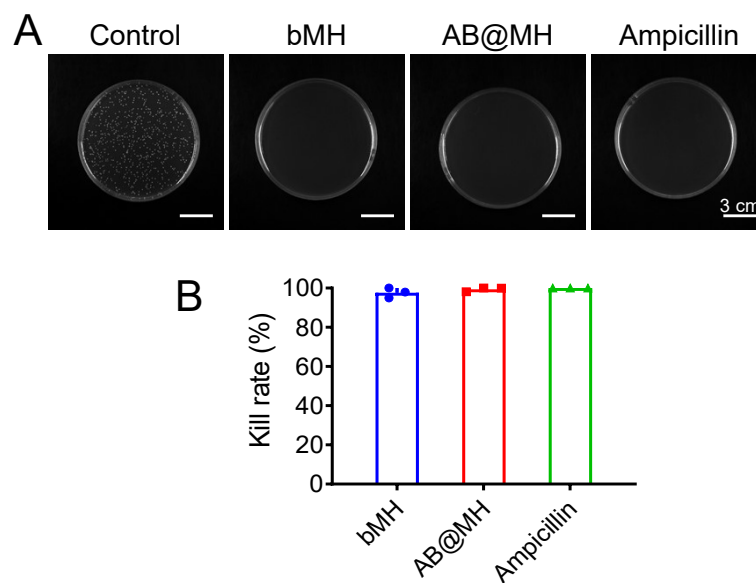


Fig. S13 (A) The photographs of bacterial colony counts of *E. coli* with 10000x dilution after culturing with different micelle-hydrogels. Ampicillin (100 $\mu\text{g mL}^{-1}$) treatment was as a positive control. (B) The kill rate of bMH, AB@MH and Ampicillin for *E. coli*.

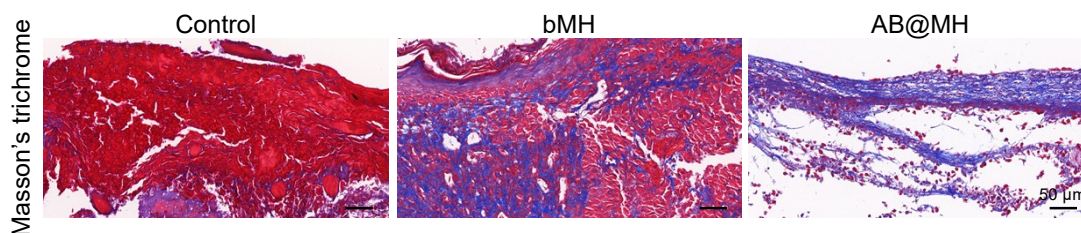


Fig. S14 Representative images of wound tissues stained with Masson's trichrome staining after different treatment at day 15 on tumor resection model.

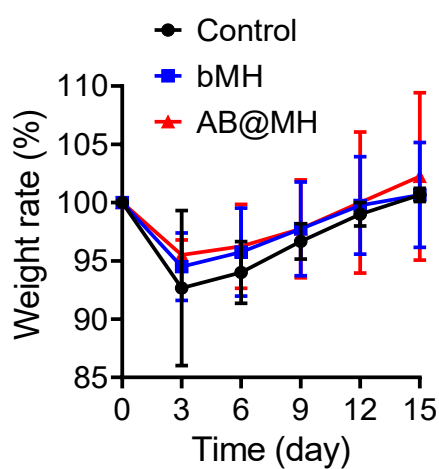


Fig. S15 Quantitative analysis of weight area at the indicated days in comparison with the original weight.

Table S1. The encapsulation efficiency (EE%) and loading capacity (LC%) of apigenin and BMS-202 in AB@M.

AB@M	EE%	LC%
apigenin	92.7 ± 0.6	2.25 ± 0.02
BMS-202	82.6 ± 1.6	0.201 ± 0.004