Supporting Information

Dual Enzyme-like Co-FeSe₂ Nanoflowers with GSH Degradation Capability for NIR II-Enhanced Catalytic Tumor Therapy

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Fig. S1. SEM images of different synthetic condition: (a) V_{OM} : V_{DDT} =1:2; (b) V_{OM} : V_{DDT} =1:1; (c) V_{OM} : V_{DDT} =2:1.

Sample	Mass ratio (%)	Atomic ratio (%)
Со	1.72	2.08
Fe	25.49	32.42
Se	72.79	65.50

Table S1 EDS result of Co-FeSe₂



Fig. S2. EDS spectrum of Co-FeSe₂ nanozymes.



Fig. S3. (a) TEM of modified Co-FeSe₂ and (b) FT-IR spectra of DSPE-PEG, Co-FeSe₂, DSPE-PEG modified Co-FeSe₂.



Fig. S4. DLS of modified Co-FeSe₂ dispersed in (a) PBS and (b) complete medium within 7 days.



Fig. S5. Fitting plots of time versus $-\ln\theta$ during the cooling period.



Fig. S6. (a) UV-Vis of modified Co-FeSe₂. (b) Temperature change profile of modified Co-FeSe₂ (1064 nm, 0.8 W/cm²).



Fig. S7. Raw data of western blot experiment. Group i) Control, ii) NIR, 1064 nm, 0.8 W/cm², 10 min, iii) Co-FeSe₂, 200 μg/mL, iv) Co-FeSe₂-NIR.



Fig. S8. CLSM imaging of mitochondrial morphology in 4T1 cells after different treatments, scale bar: 50 µm. Group i) Control, ii) NIR, 1064 nm, 0.8 W/cm², 10 min, iii) Co-FeSe₂, 200 µg/mL, iv) Co-FeSe₂-NIR



Fig. S9. HE staining images of the excised organs (heart, liver, spleen, lung, kidney) of BALB/c tumor-bearing mice at 14th day after treatment.