Supporting Information

Biodegradable FeWO_x Nanoparticles for CT/MR Imaging-Guided Synergistic Photothermal, Photodynamic and Chemodynamic Therapy

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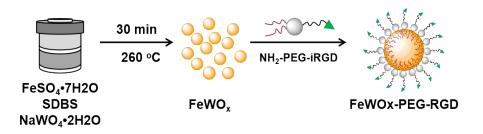


Figure S1. A scheme of the synthesis process of FeWO_x-PEG-RGD NPs.

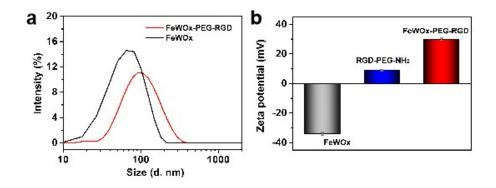


Figure S2. a) Dynamic light scattering (DLS) and b) zeta potential analysis of FeWO_x NPs and FeWO_x-PEG-RGD NPs.

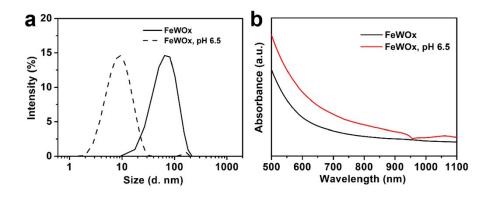


Figure S3. a) UV–vis absorption spectra of FeWO_x NPs before and after treatment in mimic TME for 24 h. b) DLS of FeWO_x NPs before and after treatment in mimic TME for 24 h.

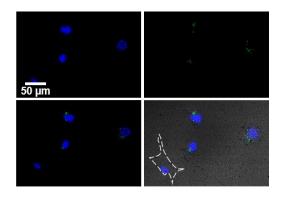


Figure S4. Confocal fluorescent microscopy images of FAM-labeled FeWO_x-PEG-RGD after incubating with MCF-7/NHDF cell lines (green = FAM-labeled FeWO_x-PEG-RGD, blue = nucleus; scale bar = 50 μ m). The NHDF cells were surrounded by write boxes.

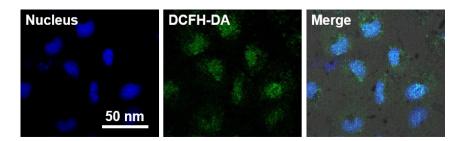


Figure S5. Confocal images of MCF-7 cells for intracellular ROS detection by ROS probe, DCFH-DA.

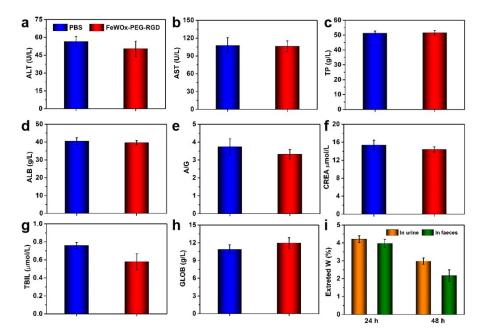


Figure S6. a-h) The serum biochemistry assay of FeWO_x-PEG-RGD NPs treated mice at a dose of 20 mg kg⁻¹ after intravenous injection 48 h. i) Accumulated W (in urine and stool) excretion out of the mice body after the injection of FeWO_x-PEG-RGD NPs through caudal vein for 24 h and 48 h.

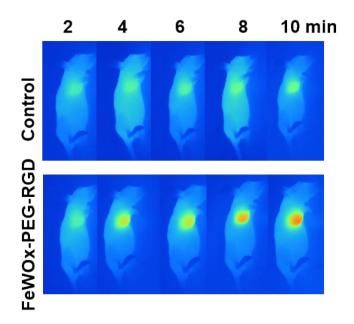


Figure S7. Infrared thermal images of tumor-bearing mice exposed to NIR laser (980 nm, 0.72 W cm⁻²) for 10 min after intravenously injected with PBS (control) or FeWO_x-PEG-RGD (20 mg kg⁻¹) for 24 h.

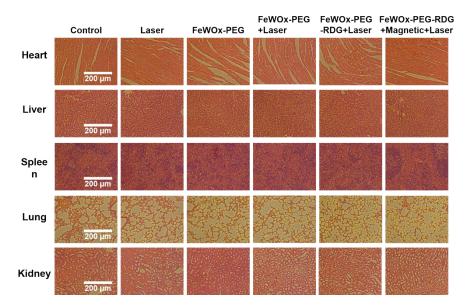


Figure S8. Histological data (H&E stained images) of the major organs (heart, liver, spleen, lung, and kidney) of mice received different treatments 16 days post injection (scale bars: 200 μm).