SUPPLEMENTARY FIGURES



Supplementary Figure 1. Scatter diagram of flow cytometry analysis in each group in LLC subcutaneous tumor model (A) and MC38 subcutaneous tumor model (B).



Supplementary Figure 2. The staining coefficient of TPE-IQ-20. (A)The gray value of MTDR and TPE-IQ-20 were calculated using the Image J Plot Profile plugin. (B) The Mander's overlap coefficient (R = 0.8429741) and Pearson's correlation coefficient (Rr = 0.846065) were determined using the Image J co-localization plugin.



Supplementary Figure 3. Selecting optimal concentration for TPE-IQ-20. (A) The fluorescence intensities of cells incubated under the concentrations of 100 nM and 200 nM; λ_{ex} : 430 nm, λ_{em} : 560 nm, the red dotted line indicates the highest average fluorescence intensities of normal cell lines, ****P*<0.001 by unpaired Student's t-test. (B) Cell viabilities measured by CCK8 assay, ****P*<0.001 vs all other cell lines at 200 nM.



Supplementary Figure 4. Tumor targeting mechanism of TPE-IQ-20. (A) Schematic illustration of TPE-IQ-20 as tumor-targeting PS (Modified from [26]). (B) Representative images of JC-1 (2 μ M) staining (JC-1 monomer and JC-1 aggregate exhibited green and red fluorescence, respectively, scar bar = 50 μ m). (C) TPE-IQ-20 and JC-1 staining was analyzed by flow cytometry. (Black) Fluorescence intensity of different cells stained with TPE-IQ-20 (200 nM) for 20 min at FITC channel and (Grey) Mitochondrial membrane potential of different cells was assessed with JC-1 (2 μ M) at PE (red) channel and FITC (green) channel (red/green ratio was represented as the ratio of aggregated and monomeric JC-1.







Supplementary Figure 6. Comparative analysis of TPE-IQ-20 and 5-ALA to induce ROS in different cells. (A) Representative flow cytometry plots of DHE in different groups. (B) Histogram shows quantitative analysis of results of flow cytometry, ***P<0.001 vs 5-ALA.



Supplementary Figure 7. Comparison of biochemical (A) and blood cell (B) analysis in control and TPE-IQ-20 PDT group (The blood of each group was detected by Chemray 800 biochemical analyzer and BC-2800vet blood cell analyzer).



Supplementary Figure 8. TUNEL staining images (A) and quantification analysis by Image J (B) of the superficial and central LLC tumor (Green indicates the TUNEL-positive cells, Scar bar=50 μm, ****P*<0.001 vs central tumors).