



Supporting Information

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Far-Red/Near-Infrared Conjugated Polymer Nanoparticles for
Long-Term In Situ Monitoring of Liver Tumor Growth

Jie Liu, Kai Li,* and Bin Liu*

Supporting Information

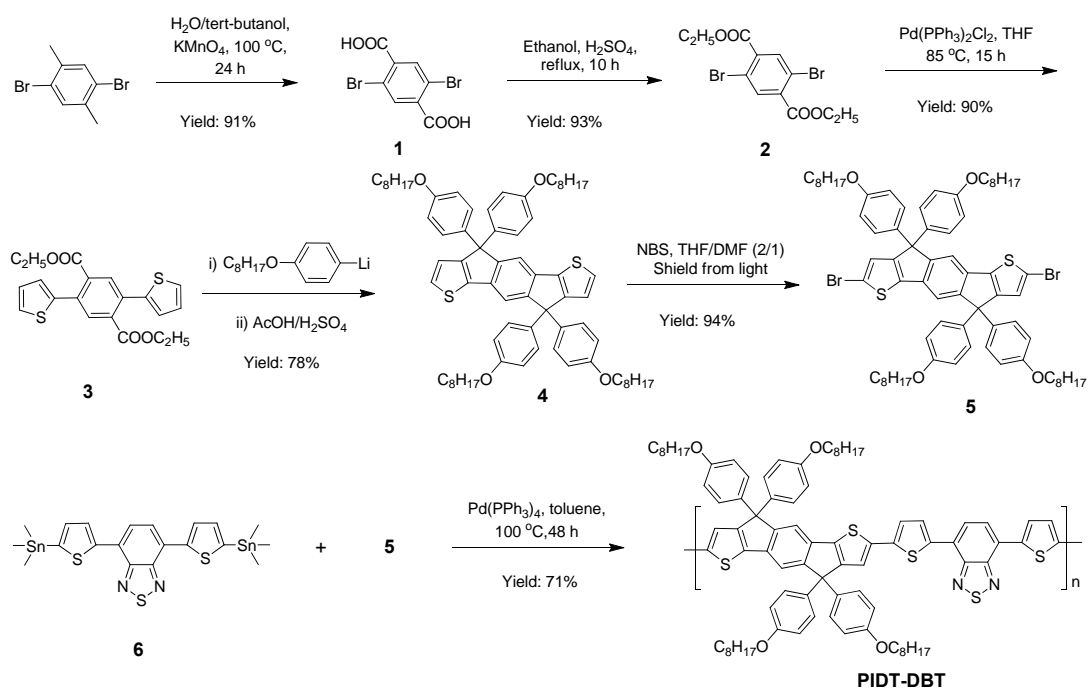
Far-Red/Near-Infrared Conjugated Polymer Nanoparticles for Long-Term In-Situ Monitoring of Liver Tumor Growth*Jie Liu, Kai Li,* and Bin Liu****Calculation of PIDT-DBT-Tat NP concentration:**

The PIDT-DBT-Tat NPs stock solution (5 mL) was freeze-dried to yield 1.2 mg of powders. As the PIDT-DBT-Tat NPs have good colloidal stability in water without precipitation, the density of the NP suspension could be estimated as $\sim 1 \text{ g/cm}^3$. The average size of PIDT-DBT-Tat NPs determined from TEM is $\sim 49 \text{ nm}$. Thus, the concentration of the PIDT-DBT-Tat NPs in stock can be calculated from the following equation:

$$\begin{aligned} & \text{Total number of PIDT - DBT - Tat NPs in 5 mL of suspension} \\ = & \frac{\text{Total Volume of PIDT - DBT - Tat NPs}}{\text{Average Volume of Each NP}} = \frac{\frac{1.2 \times 10^{-3} \text{ g}}{1 \text{ g/mL}}}{\frac{4}{3} \pi \times (24.5 \times 10^{-7})^3 \text{ mL}} = 1.9 \times 10^{13} \end{aligned}$$

Finally, the concentration of PIDT-DBT-Tat NPs in stock solution was calculated as

$$\text{following: } [\text{PIDT - DBT - Tat NP}] = \frac{2.3 \times 10^{13}}{2 \times 10^{-3} \text{ L}} = 16 \text{ nM}$$



Scheme S1. Synthetic route towards PIDT-DBT.

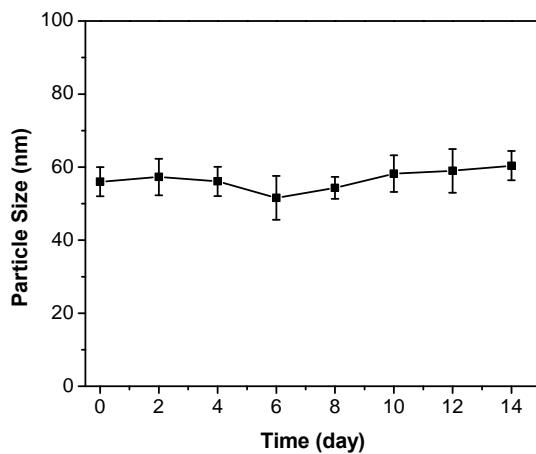


Figure S1. Average hydrodynamic diameter change of PIDT-DBT-Tat NPs when incubating in phosphate-buffered saline at 37 °C for 14 days.

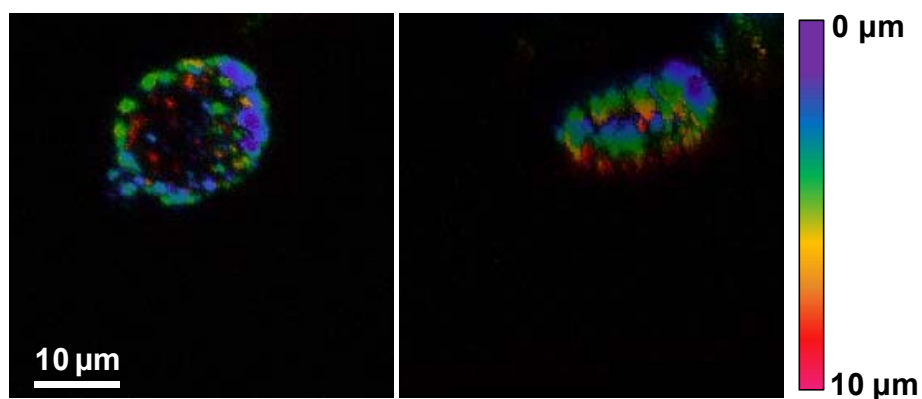


Figure S2. Color-coded projections of z-stacks of confocal images of HepG2 cells upon incubation with PIDT-DBT-Tat NPs overnight. ($\lambda_{\text{ex}} = 590 \text{ nm}$, 600-800 nm bandpass filter).

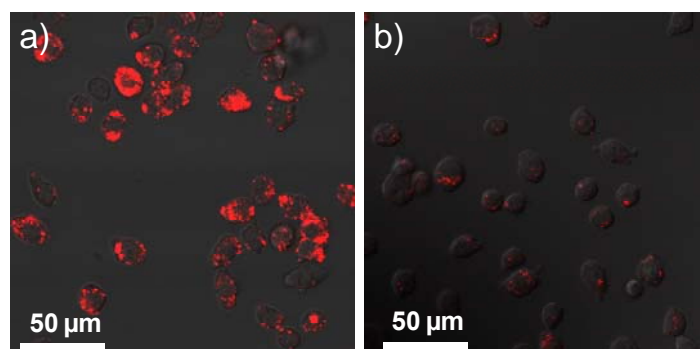


Figure S3. CLSM fluorescence/transmission overlay images of HepG2 cancer cells after incubation with 2 nM (a) PIDT-DBT-Tat NPs and (b) PIDT-DBT NPs at 37 °C for 24 h. ($\lambda_{\text{ex}} = 561 \text{ nm}$, 710/50 nm bandpass filter).

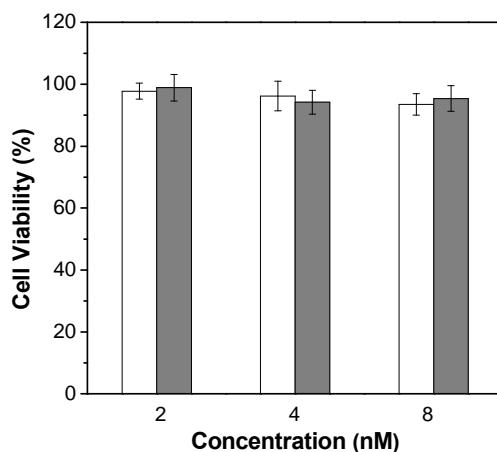


Figure S4. Metabolic viability of HepG2 cells (blank) and NIH/3T3 cells (gray) after incubation with 2, 4 and 8 nM PIDT-DBT-Tat NPs for 48 h.

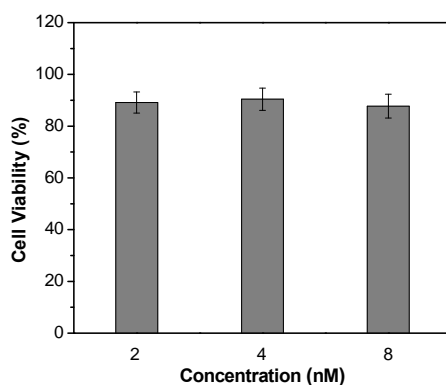


Figure S5. Metabolic viability of NIH/3T3 cells after incubation with 2, 4 and 8 nM PIDT-DBT-Tat NPs for 72 h.

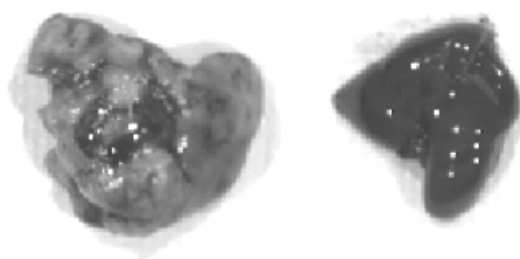


Figure S6. Bright-field image of the liver collected from mouse transplanted with 4×10^6 of PIDT-DBT-Tat NP-labeled HepG2 cells (left) and mouse without transplantation (right) after 42-day injection.

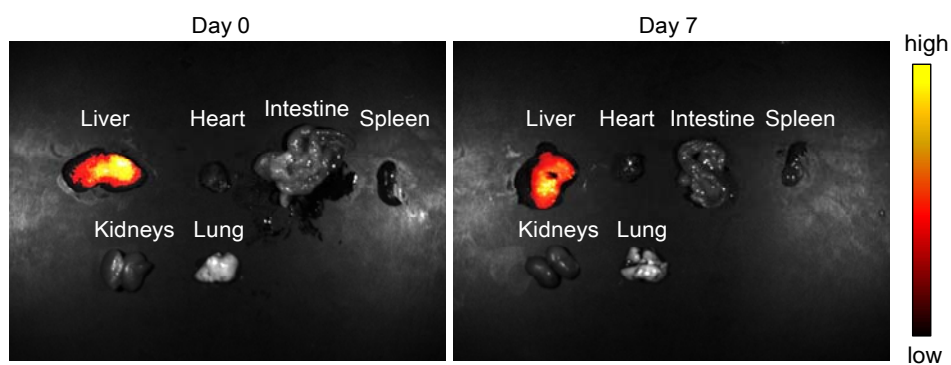


Figure 7. Representative *ex vivo* fluorescence images of the organ tissues collected from mouse transplanted with 4×10^6 of PIDT-DBT-Tat NP-labeled HepG2 cells after 0 and 7 days post injection.