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Supporting Information

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Tumor Microenvironment Activable Self-Assembled DNA Hybrids for pH and Redox Dual-Responsive Chemotherapy/PDT Treatment of Hepatocellular Carcinoma

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Tumor micro-environment activable self-assembled DNA hybrids for pH and redox dual-responsive chemotherapy / PDT treatment of Hepatocellular carcinoma

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Figure S1. (A) Lane 1 and 8 is the DNA marker, lanes 2 to 4 are RQD, CD and TD, respectively. Lane 5, 6 is RQD + CD (without Ce6) and CD (without Ce6) + TD, respectively. Lane 7 was the self-assembled final products containing RQD, CD (without Ce6) and TD.



Figure S2. The mean fluorescence intensity (MFI) of Ce6 in CD or Ce6-fDNA as indicated (n = 3).

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Figure S3. Fluorescence emission spectrum of Ce6-fDNA probe (excitation at 405 nm) in the absence of DTT (10 mM).



Figure S4. The mean fluorescence intensity (MFI) of Ce6 in Ce6-fDNA with (black) or without DTT (red) as indicated (excitation 405 nm, emission 670 nm) (n = 3).

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Figure S5. Absorbance of 9, 10-dimethylanthracene (ABDA, 100 μ M) after photodecomposition by ROS generation upon NIR laser irradiation (670 nm (0.2 W/cm²)) in the presence of the Ce6-fDNA only in TM buffer.



Figure S6. The mean fluorescence intensity (MFI) of Dox in free DOX and in Ce6-fDNA^{DOX} as indicated (excitation 488 nm, emission 595 nm) (n = 3).



Figure S7. Fluorescence spectra of DOX were used to determine the release of doxorubicin from the Ce6-fDNA^{DOX} triggered by low pH value.

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Figure S8. HepG2 cells incubated with Ce6-fDNA^{DOX} probe or Ce6-fNDNA^{DOX} (Ce6, 2.05 μ M; Dox, 39.5 μ M) for 2 h, 2h + 2h, 2h + 4h, 2h + 6h and 2h + 24h, respectively (n =5).