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Supporting Information

FRET investigation toward DNA tetrahedron-based ratiometric analysis of

intracellular telomerase activity

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Fig. S1 Gel electrophoresis analysis of six samples formed by different DNA probes. Lane A is strand S1; lane B is strand S1+S2; lane C is strand S1+S2+S3; lane D is strand S1+S2+S3+S4; lane E is strand S1+S2+S3+S4+Flare+telomerase; lane F is strand S1+S2+S3+S4+Flare.



Fig. S2 Cytotoxicity of different concentrations of DNA nanoprobe exposed to the cells for 24 h.



Fig. S3 (a) Fluorescence spectra of Flare with different concentrations. (b) The relationship between peak intensity at 565 nm (Cy3) and Flare concentration. (c) The relationship between peak intensity at 662 nm (Cy5) and Flare concentration. (d) The relationship between the peak intensity ratio of donor to acceptor (F_d/F_a) and Flare concentration.



Fig. S4 (A) Fluorescence responses in the presence of different concentrations of telomerase from the Kit. (B) Calibration curve reflecting the relationship between F_d/F_a and telomerase concentration.



Fig. S5 Confocal images of DNA nanoprobe delivered into SW480 cells.

Table S1 DNA sequences used in this study.^{α}

Name	Sequence (from 5' to 3')
S1	ACATTCCTAAGTCTGAAGAAGAGCCGCCATAGTACCTCAGCCAACCC <u>TAACCCT</u>
	AA <u>CCCT</u> AACCCTAAAACTCTGCTCGACGGATTCCTTCCCACGTAGTGTCGTTATC
	ACCAGGCAGTTGATTGCGCAA
S2	ATGCGAGGGTCCAATACGCACCTGAGACGCGTTACGCATGACCCGAACTGGTCC
	CGTCTACTTACCGGCGACAGTCGTTAAGCTGAAATTGCGCAATCAACTGCCTGG
	TGATAACGACACTACGTGGGAA
S 3	ATTACAGCTTGCTACACCAACTAGAATGATCGGCTACAGACGAAGGCTGAGGTA
	CTATGGCGGCTCTTCTTCAGACTTAGGAATGTCCTCAGCTTAACGACTGTCGCCG
	GTAAGTAGACGGGACCAGTTC
S4	CGTCTGTAGCCGATCATTCTAGTTGGTGTAGCAAGCTGTAATCCGTCATGCGTAA
	CGCGTCTCAGGTGCGTATTGGACCCTCGCATCCAATCCGTCGAGCAGAGTT
Flare	Cy3-ATACAC <u>AGGG</u> GGCCAA <u>AGGGTT</u> CTCAAT-Cy5

 $^{\alpha}$ The underlined sequences are complementary with each other.