## **Supporting Information**

Name	Sequences (5'-3')	Base length
poly[dGdC]:poly[dGdC]	5'-GCGCGCGCGCGCGCGCGCGCGCGCGCG	34
(GC-DNA)	CGCGCGCGC -3'	
poly[dAdT]:poly[dAdT]	5'-ATATATATATATATATATATATATATATATAT	34
(AT-DNA)	ATAT -3'	
polydApolydT	5'- AAAAAAAAAAAAAAAAAAAAAAAAAA	22
	3'- TTTTTTTTTTTTTTTTTTTTTTTT-5'	

Table S1. The DNA sequences used in article.



Figure S1. Depiction of the SC-dots formation



Figure S2 Diameters statistics of SC-dots.



Figure S3: (A) XPS, (B)  $C_{1s}$ , (C)  $N_{1s}$  and (D)  $O_{1s}$  spectra of SC-dots.



Figure S4: XRD patterns for SC-dots.



Figure S5: (A) Plots of CD intensity versus concentration of SC-dots. CD intensity at 247 nm (black) and at 285 nm (red). (B) UV spectra of AT-DNA in the absence (black) or presence (red) of the 2  $\mu$ g/mL SC-dots.



Figure S6: (A) TEM image of G-dots; (B) Diameters statistics of G-dots.



Figure S7: CD spectra of GC-DNA in the absence (black) or presence (red) of 5  $\mu$ g/mL SC-dots in phosphate buffer containing 2 M NaCl.



Figure S8: Plots of CD intensity of ct-DNA versus concentration of SC-dots. CD intensity at 295nm (black) and at 252nm (red).



Figure S9: (A) CD spectra of 30  $\mu$ g ml<sup>-1</sup> ct-DNA (black), DNA solutions after addition with 5  $\mu$ g/mL G-dots (red) and after addition with 5 mM spermine (blue). (B) CD spectra of ct-DNA in the absence (black) or presence (red) of 5  $\mu$ g/mL SC-dots in phosphate buffer containing 4 M NaCl.



Figure S10: Agarose gel electrophoresis of ct-DNA with different SC-dots. The samples were electrophoresed on a 1% agarose gel for 30 min. Lane 1: DNA marker 2000 (TaKaRa company). Lane 2-10: SC-dots concentrations of 0, 0.5, 1, 2, 3, 5, 8,10, 20  $\mu$ g/mL, respectively. The gel electrophoresis was carried out at 37°C after incubation for 0.5h.



Figure S11: (A) CD spectra of 2  $\mu$ M polydApolydT (black), DNA solutions after addition of 3  $\mu$ g/mL SC-dots (red) and 5  $\mu$ g/mL SC-dots (blue). (B) UV spectra of polydApolydT duplex DNA in the absence (black) or presence (red) of 3  $\mu$ g/mL SC-dots.



Figure S12: Add different amounts of NaI solution from 5 to 50 mM in the 5  $\mu$ g ml<sup>-1</sup> SC-dots solution.



Figure S13: AFM images of (A) SC-dots, (B) ct-DNA, (C) ct-DNA with  $5\mu g/mL$  SC-dots, (D) ct-DNA with 10  $\mu g/mL$  SC-dots. The ct-DNA was condensed as a network after addition of 5  $\mu g/mL$  SC-dots, looking like forming DNA-wrapped complexes (Figure S15 C). By further increasing the amount of SC-dots higher to 10  $\mu g/mL$ , the condensed microsphere would be formed (Figure S15 D).



Figure S14: Add different amounts of NaI solution from 5 to 50 mM in the SC-dots/ct-DNA/EB solution with excitation wavelength of 400 nm (A) and 480 nm (B).



Figure S15: Iodide ions were used to quench the fluorescenc of SC-dots, thus no FRET occurred between SC-dots and EB. The concentration of NaI: 50 mM.