

Supplementary Material

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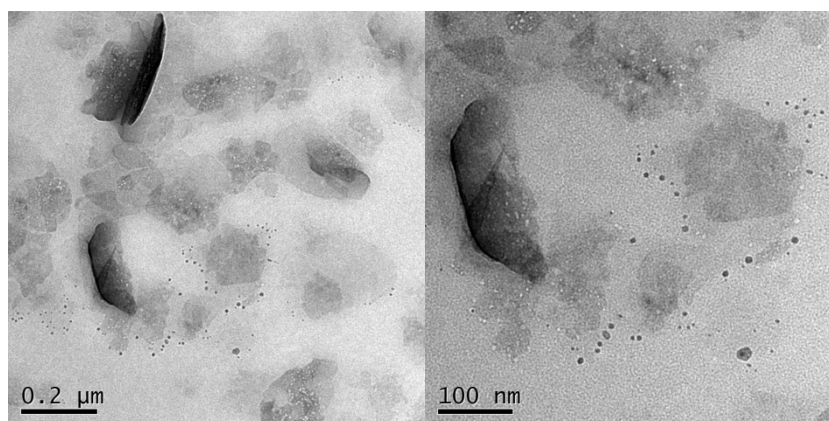


Figure S1. Other size TEM images (0.2 μm , 100 nm) of C-CQDs. It is obvious that the size of C-CQDs is mainly distributed below 10 nm, and combining the results of DLS can indicate that the size of C-CQDs is between 7 and 9 nm.

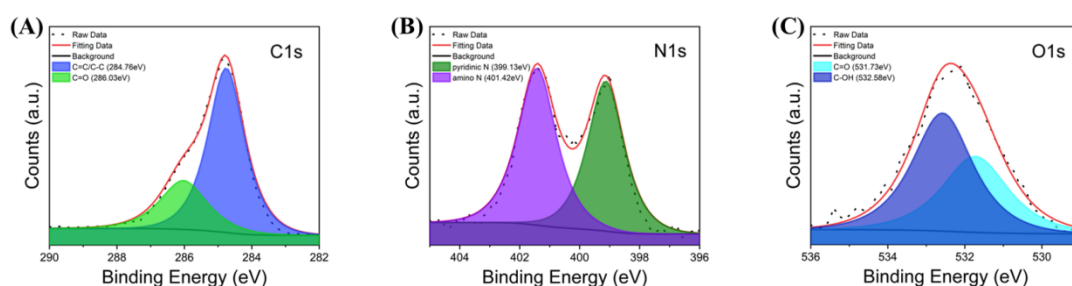


Figure S2. C 1s(A), N 1s(B), O 1s(C) XPS spectrum of C-CDs. The high-resolution C 1s spectrum (A) can be deconvoluted into two peaks at 284.76 and 286.03 eV, which corresponded to C=C/C-C and C=O bonds, respectively. The high-resolution N 1s spectrum (B) shows two peaks at 399.13 and 401.42 eV, which indicated the presence of pyridinic N and amino N, respectively. The two main bands at 531.73 and 532.58 eV in the high-resolution O 1s spectrum (C) could be identified as C=O and C-OH, respectively.

Table S1. Analytical results for the detection of TCs in real samples (n = 5). Milk has been processed. C-CQDs were prepared one day earlier and used after 24h of placement. To reduce the error, detection was performed immediately after the addition of antibiotics.

Tetracycline	Added ($\mu\text{g/mL}$)	Found ($\mu\text{g/mL}$)	Recovery (%)	RSD (%)
CTC	2.00	1.97	98.5	0.88
	5.00	4.99	99.8	0.94
	10.00	10.21	102.1	0.83
OTC	2.00	2.04	100.5	1.02
	5.00	5.04	100.8	0.94
	10.00	9.98	99.8	0.88
TET	2.00	1.94	97.0	0.82
	5.00	4.91	98.2	0.87
	10.00	9.97	99.7	0.97

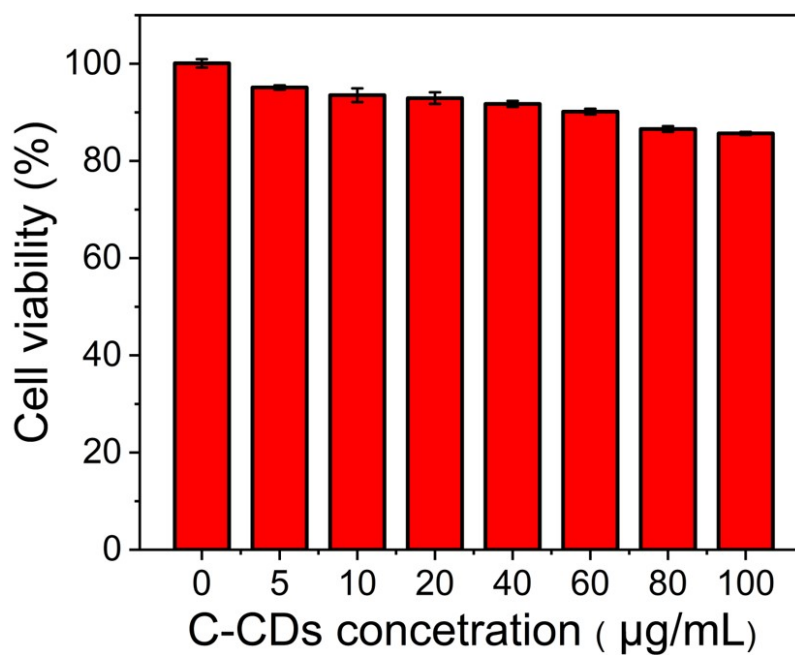


Figure S3. The cell viabilities of HeLa cells under different concentrations of the C-CDs.

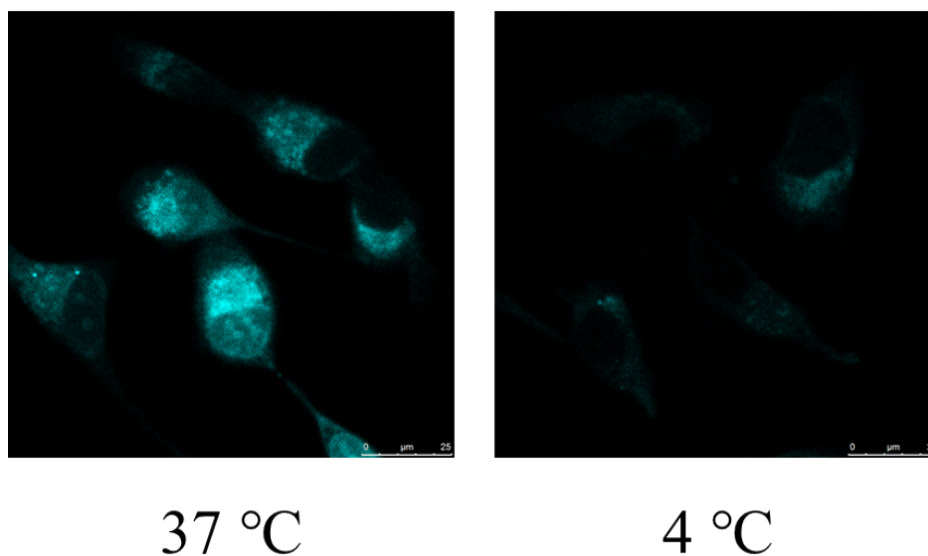


Figure S4. Laser scanning confocal microscope images of HeLa cells under 37°C and 4°C. It can be seen that the fluorescence intensity of C-CQDs in the cells decreased significantly, which indicated that the uptake of C-CQDs depends on cellular energy.