Electronic supplementary information

Hydrophobically Modified Carbon Dots as Multifunctional

Platform for Serum-Resistant Gene Delivery and Cell Imaging

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Table S1. The QY of PEI-CD and Ole-CDs.

	PEI-CD	Ole0.5-CD	Ole1-CD	Ole1.5-CD	Ole2-CD	Ole2.5-CD
YQ (%	21.49%	16.00%	12.00%	11.20%	7.70%	4.95%

Table S2. Mean DLS PDI of **Ole-CDs** under the concentration of 20 mg/L. Data represent mean \pm SD (n = 3).

	Ole0.5-CD	Ole1-CD	Ole1.5-CD	Ole2-CD	Ole2.5-CD
PDI	0.322 ± 0.017	0.381 ± 0.075	0.293 ± 0.013	0.299 ± 0.026	0.305 ± 0.054



Fig. S1. Fluorescence microscopy images of pEGFP transfected by PEI-CD and X-CD in A549 cells. Scale bar = $100 \mu m$.



Fig. S2. ¹H NMR spectrum of PEI-CD and X-CD in CDCl₃.



Fig. S3. ¹H NMR spectrum of PEI-CD and Ole-CDs in CDCl₃(A) and D₂O (B).



Fig. S4. FT-IR spectrum of PEI-CD and X-CD.



Fig. S5. FT-IR spectrum of PEI-CD and Ole-CDs.



Fig. S6. Mean DLS particle sizes (red) and surface charge (blue) of **Ole-CDs** under the concentration of 20 mg/L. Data represent mean \pm SD (n = 3).



Fig. S7. The full spectra (A & E) and high resolution XPS spectra of C 1s (B & F), N 1s (C & G), O 1s (D & H) for **PEI-CD** (upper) and **Ole1.5-CD** (lower).



Fig. S8. The maximum PL excitation and emission spectra of Ole-CDs aqueous solution, (A) Ole0.5-CD, (B) Ole1-CD, (C) Ole2-CD, (D) Ole2.5-CD.



Fig. S9. Electrophoretic gel retardation assays of pDNA binding for **X-CD**/pDNA complexes under various mass ratios.



Fig. S10. Confocal fluorescence images of A549 cells incubated with **Ole-CDs** under the concentration of 10 mg/L. Scale bar = $20 \mu \text{m}$.



Fig. S11. Fluorescence microscopy images of pEGFP-transfected Hela, 7702 and HepG2 cells by **Ole-CDs** and PEI 25 kDa. Scale bar =200 µm.



Fig. S12. BSA adsorption assays, photographs show the PEI solution(0.5 mg/mL) turned turbid by addition of BSA(1mg/mL), while Ole-CD solution(0.5 mg/mL) is still clear.