

Supplementary Materials

Synthesis Mechanisms, Structural Models and Photothermal Therapy Applications of Top-Down Carbon Dots from Carbon Powder, Graphite, Graphene and Carbon Nanotubes

Wenquan Shi,^{a,b,+} Qiurui Han,^{a,+} Jiajia Wu,^a Chunyu Ji,^a Yiqun Zhou,^c Shanghao Li,^d
Lipeng Gao,^a Roger M. Leblanc^c and Zhili Peng,^{*a,b}

^aSchool of Materials and Energy, Yunnan University, Kunming 650091, People's Republic of China.

^bAdvanced Computing Center, Materials Genome Institute, Yunnan University, Kunming 650091, P. R. China

^cDepartment of Chemistry, University of Miami, 1301 Memorial Drive, Coral Gables, Florida 33146, United States.

^dMP Biomedicals, 9 Goddard, Irvine, CA 92618, USA.

***Corresponding authors:**

(Z. P.) Tel.: +86–871–65037399; E-mail: zhilip@ynu.edu.cn.

⁺These authors contributed equally to this work.

Keywords:

Carbon dots; carbon nano powders; graphite; graphene; carbon nano tubes; photothermal therapy

Table S1. Effects of reaction temperature, time and scale on the yields of GR-CDs

	Mass (mg)	H ₂ SO ₄ (mL)	HNO ₃ (mL)	Temperature (°C)	Time (h)	Yield (%)
1	250	9	3	110	7.5	10
2	250	9	3	75	15	3
3	500	18	6	110	15	11

Table S2. Volumes of NaOH solution consumed for titrations of CDs and the corresponding carboxyl contents determined for each CDs

Sample	V _{NaOH} consumed (mL)	-COOH (mmol/g)
C-CDs	1.7	6.8
	1.7	6.8
	1.5	6.0
G-CDs	1.1	4.4
	1.0	4.0
	1.1	4.4
GR-CDs	1.0	4.0
	1.2	4.8
	1.1	4.4
CNT-CDs	2.2	8.8
	2.1	8.4
	2.1	8.4

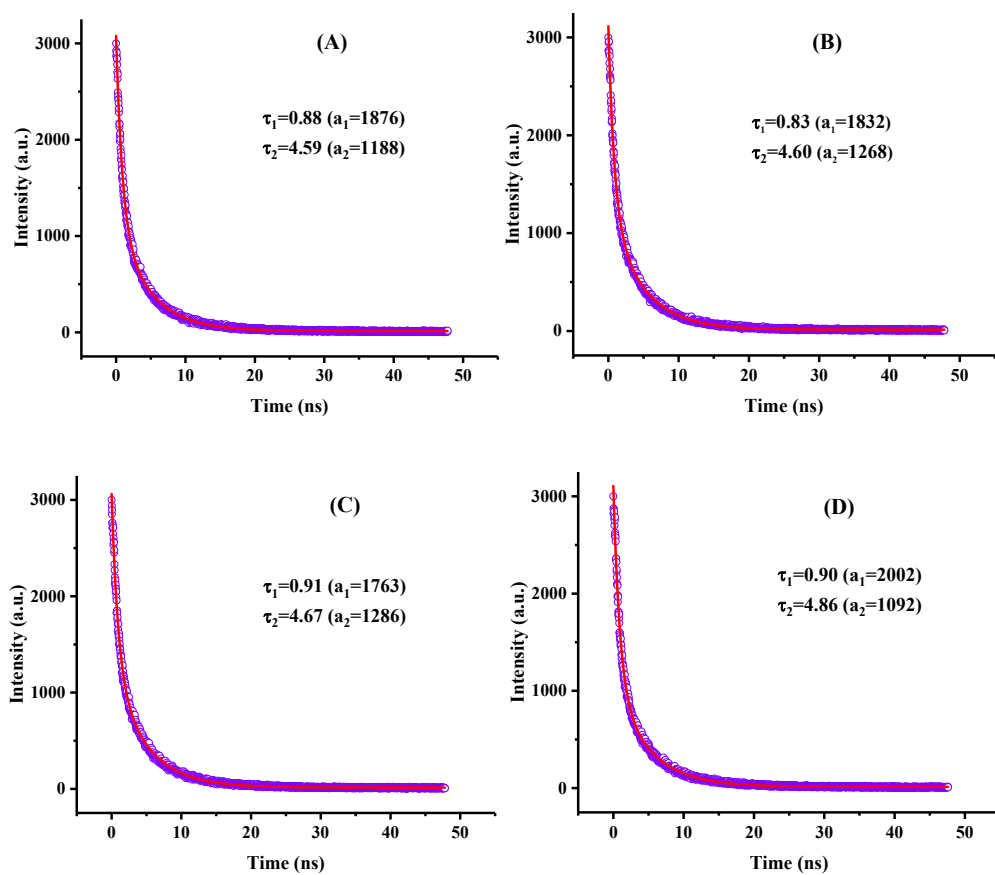


Figure S1. The photoluminescence decay curves of C-CDs (A), G-CDs (B), GR-CDs (C) and CNT-CDs (D) in 0.5 mg/mL aqueous solution, excited at 475 nm.

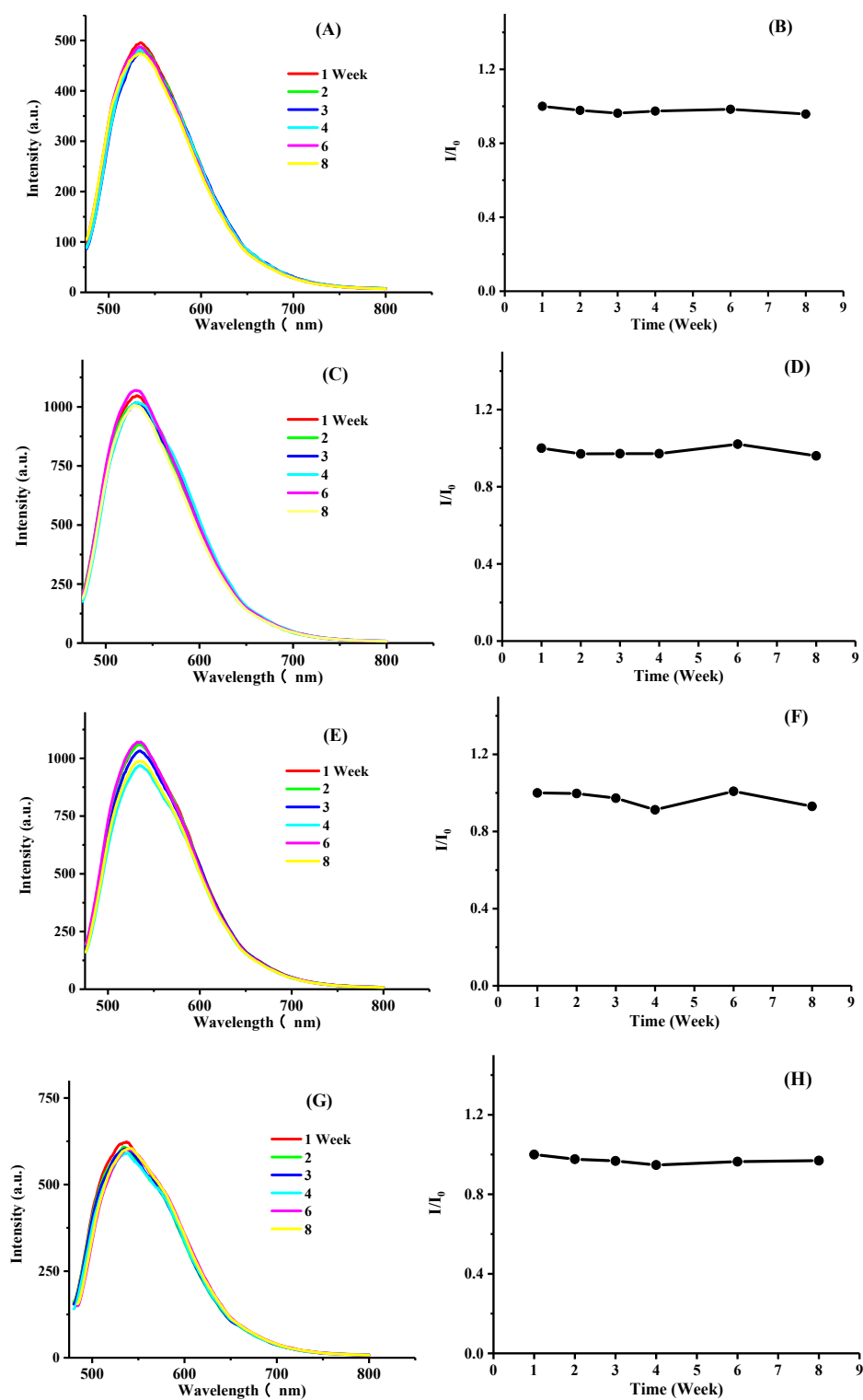


Figure S2. Photoluminescence stability tests of 0.5 mg/mL of C-CDs (A, B), G-CDs (C, D), GR-CDs (E, F) and CNT-CDs (G, H): solutions of CDs were placed in the ambient environment and their fluorescence emissions (excited at 460 nm) were monitored for two months.

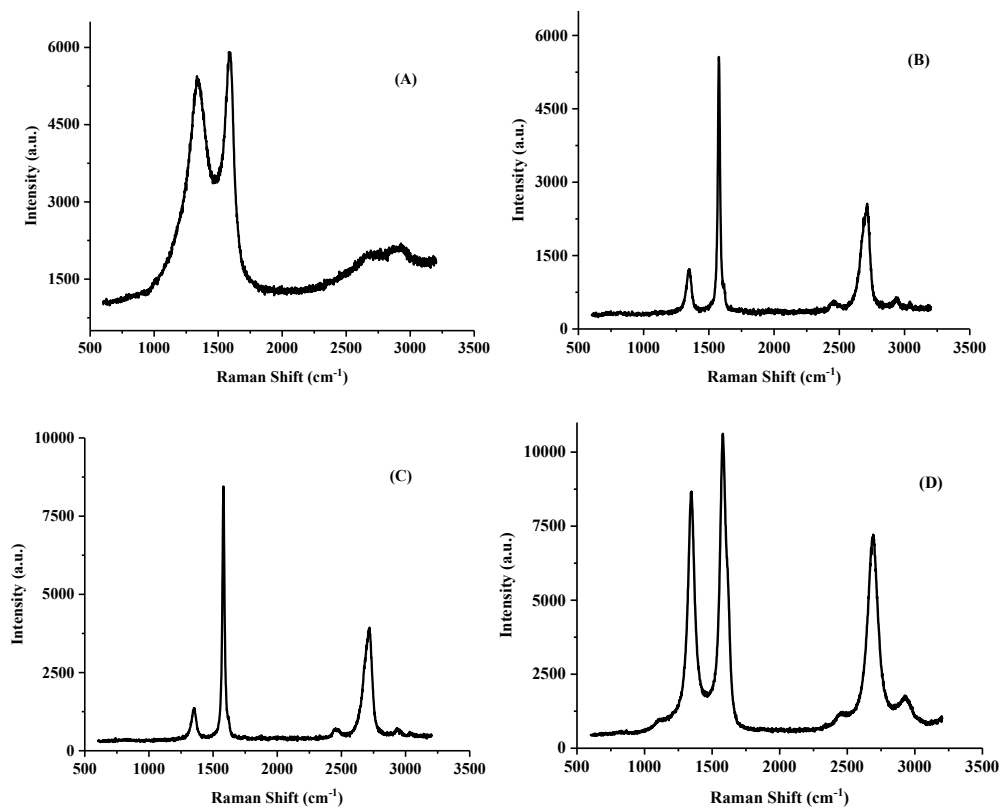


Figure S3. Raman spectra of carbon powders (A), graphite (B), graphene (C) and carbon nanotubes (D) used to synthesize CDs in this study.

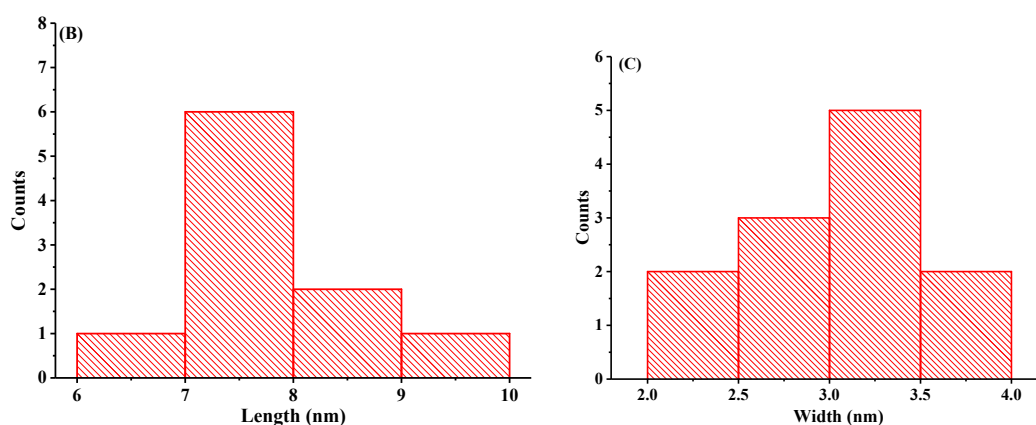
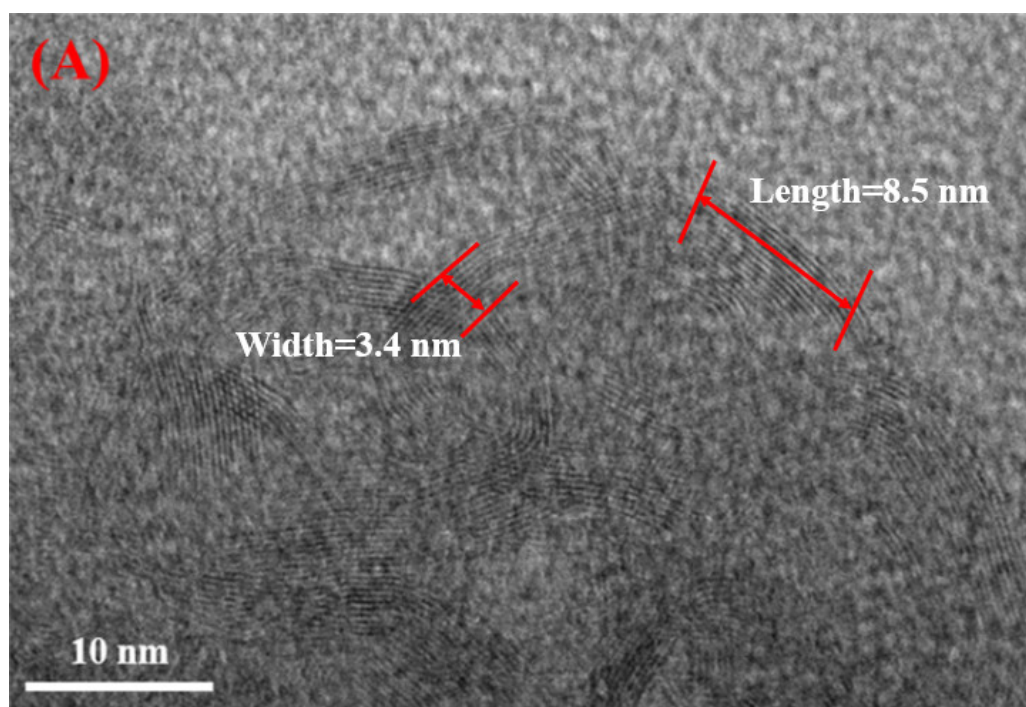


Figure S4. TEM image of broken carbon nanotubes (A), and the histograms showing the distributions of length (B) and width (C) of these pieces

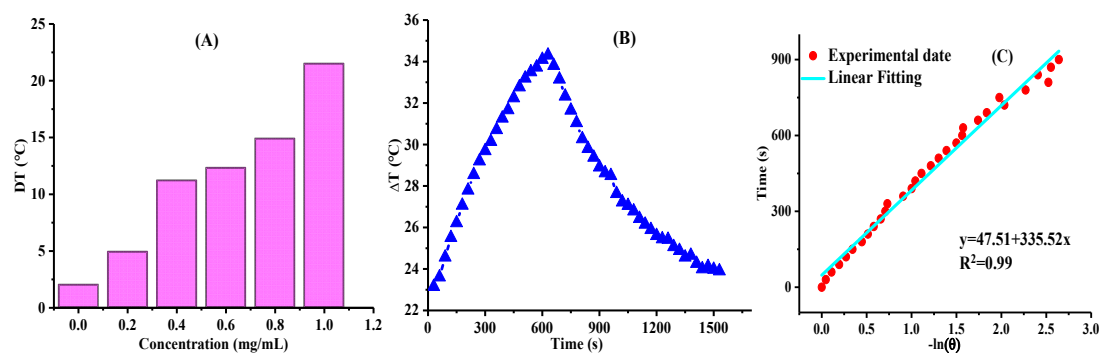


Figure S5. Photothermal properties of C-CDs: (A) Temperature change (ΔT) diagram of water and C-CDs solutions with concentrations range from 0.2 to 1.0 mg/mL after laser irradiation for 10 minutes; (B) Photothermal effect of 1 mL of C-CDs solution with concentration of 0.4 mg/mL after laser irradiation for 10 minutes; (C) Linear relationship between cooling time and $-\ln(\theta)$ obtained from Figure S5B.

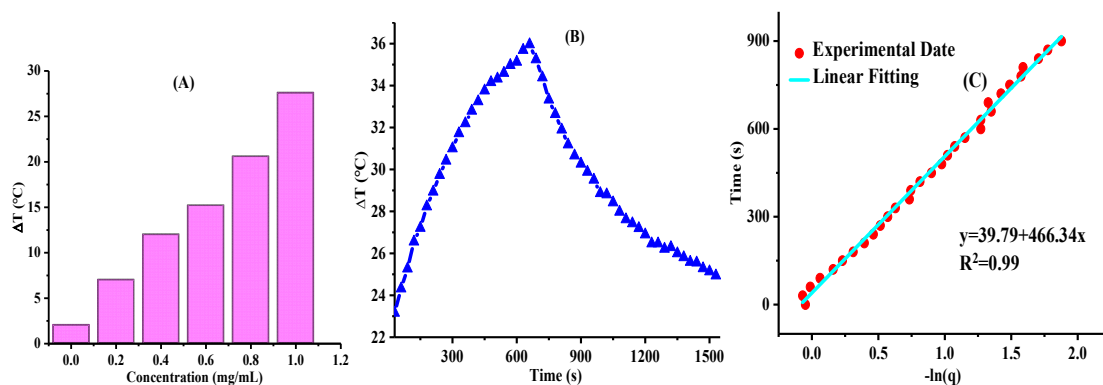


Figure S6. Photothermal properties of G-CDs: (A) Temperature change (ΔT) diagram of water and G-CDs solutions with concentrations range from 0.2 to 1.0 mg/mL after laser irradiation for 10 minutes; (B) Photothermal effect of 1 mL of G-CDs solution with concentration of 0.4 mg/mL after laser irradiation for 10 minutes; (C) Linear relationship between cooling time and $-\ln(\theta)$ obtained from Figure S6B.

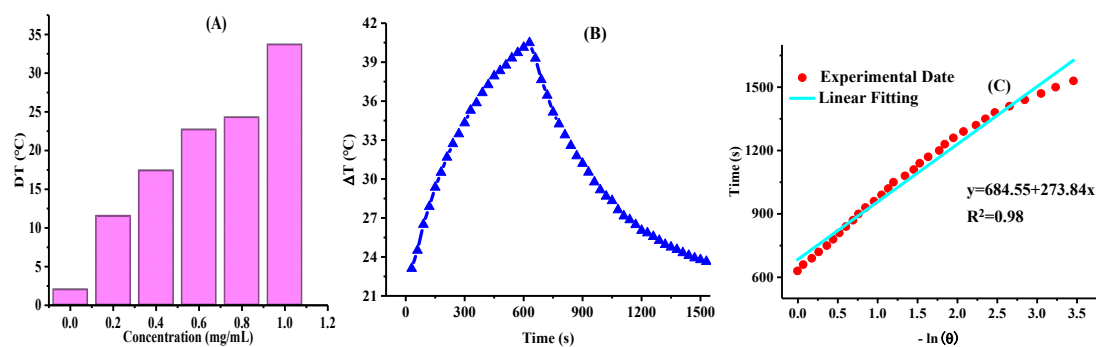


Figure S7. Photothermal properties of GR-CDs: (A) Temperature change (ΔT) diagram of water and GR-CDs solutions with concentrations range from 0.2 to 1.0 mg/mL after laser irradiation for 10 minutes; (B) Photothermal effect of 1 mL of GR-CDs solution with concentration of 0.4 mg/mL after laser irradiation for 10 minutes; (C) Linear relationship between cooling time and $-\ln(\theta)$ obtained from Figure S7B.

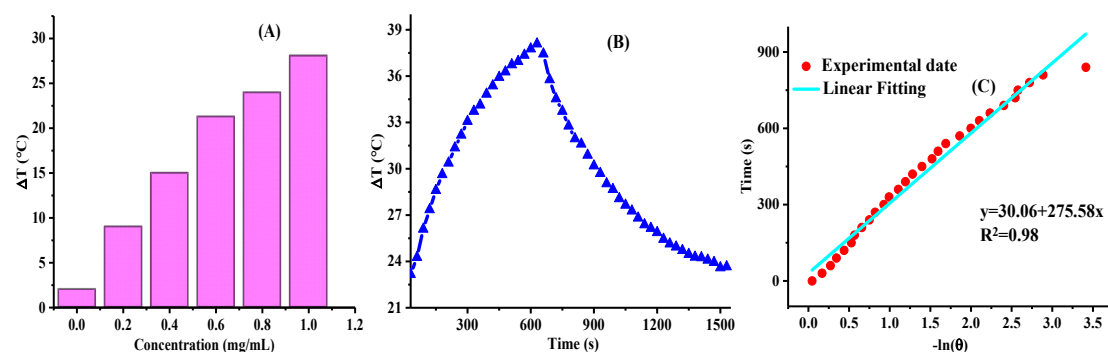


Figure S8. Photothermal properties of CNT-CDs: (A) Temperature change (ΔT) diagram of water and CNT-CDs solutions with concentrations range from 0.2 to 1.0 mg/mL after laser irradiation for 10 minutes; (B) Photothermal effect of 1 mL of CNT-CDs solution with concentration of 0.4 mg/mL after laser irradiation for 10 minutes; (C) Linear relationship between cooling time and $-\ln(\theta)$ obtained from Figure S8B.