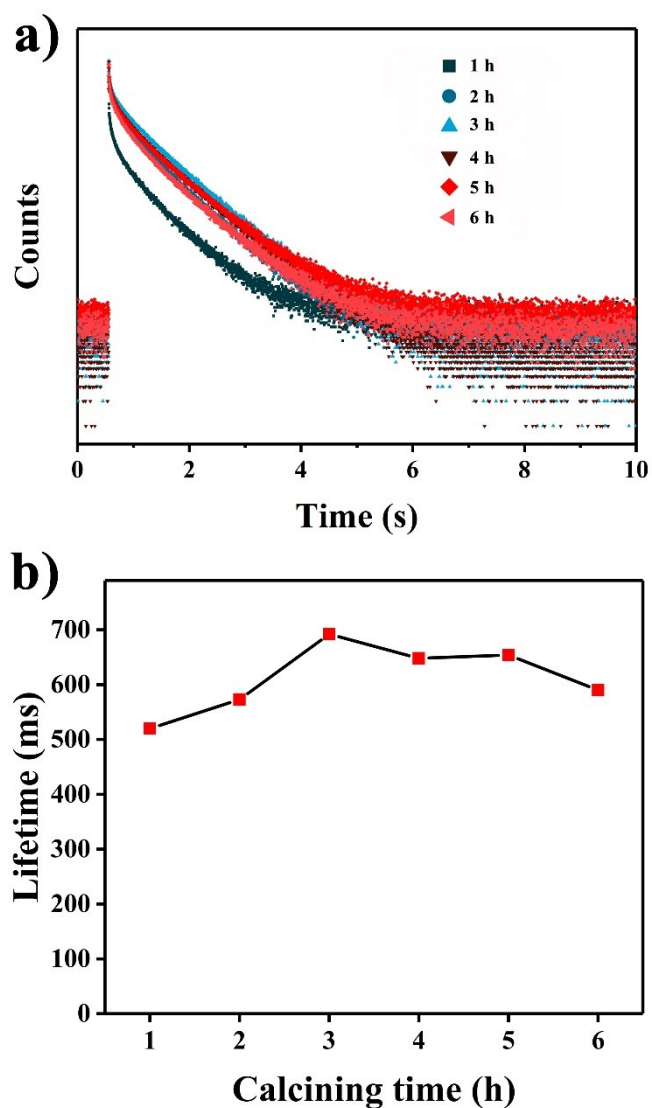


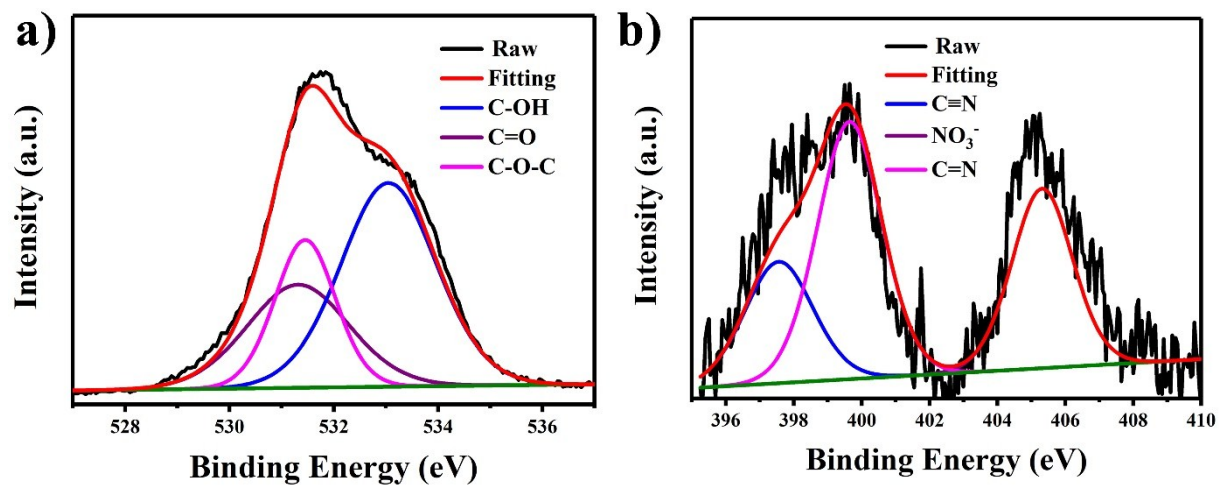
## **Supplementary Information**

### **Color tunable room temperature phosphorescent carbon dots based nanocomposites obtainable from multiple carbon sources via a molten salt method**

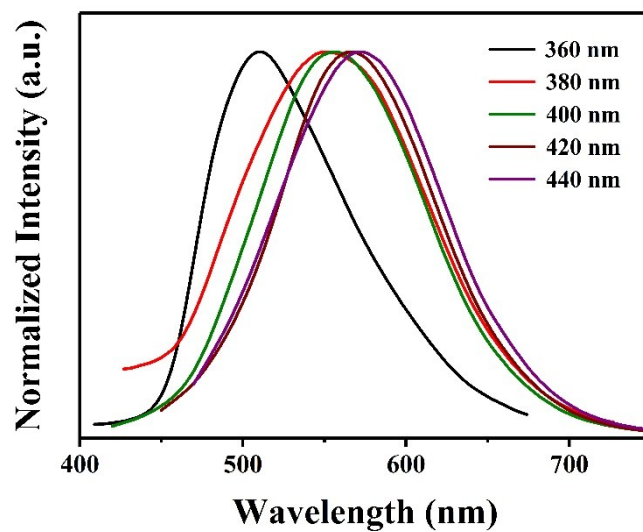
Chan Wang, Yueyue Chen, Tantan Hu, Yong Chang, Mei Wang and Qijun Song\*



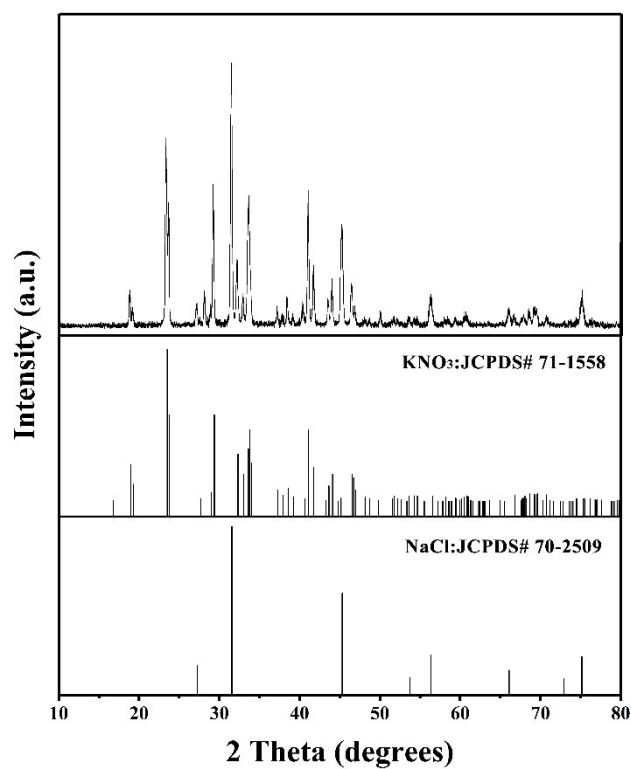
**Figure S1.** Corresponding lifetimes of the CDs@MS nanocomposites under different calcining time at the same synthesis temperature of 350 °C (1h: 520 ms; 2h: 573 ms; 3h: 701 ms; 4h: 648 ms; 5h: 654 ms; 6h: 590 ms).



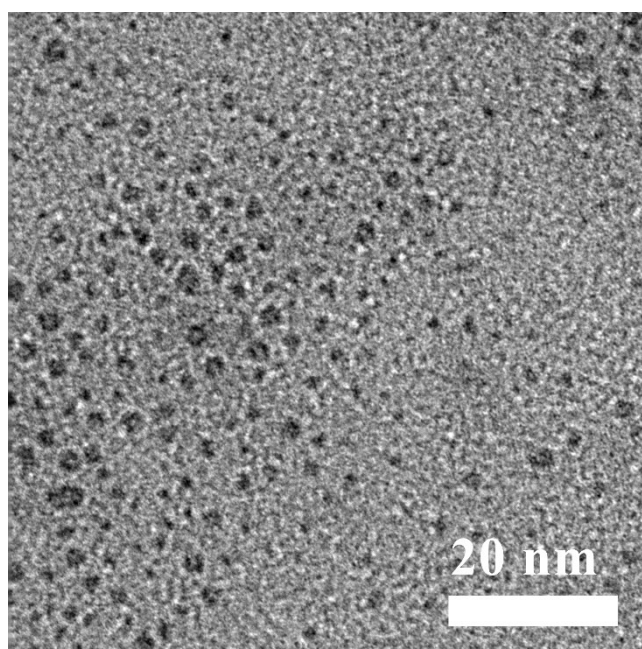
**Figure S2.** The high-resolution XPS spectra of O 1s and N 1s, respectively.



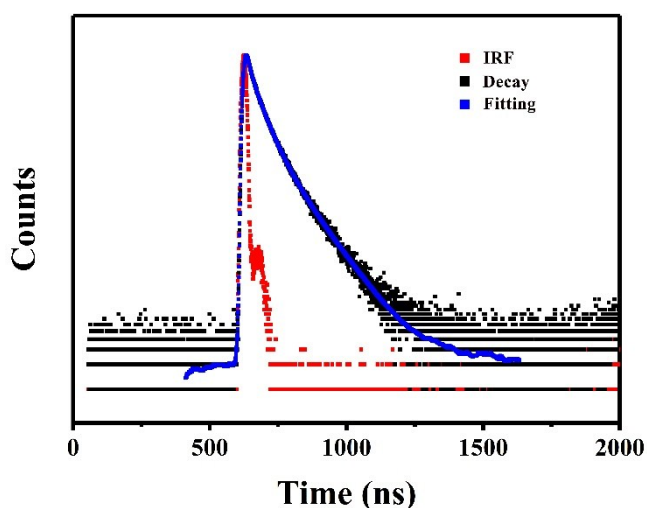
**Figure S3.** Normalized phosphorescence spectra of the CDs@MS nanocomposites under different excitation wavelengths.



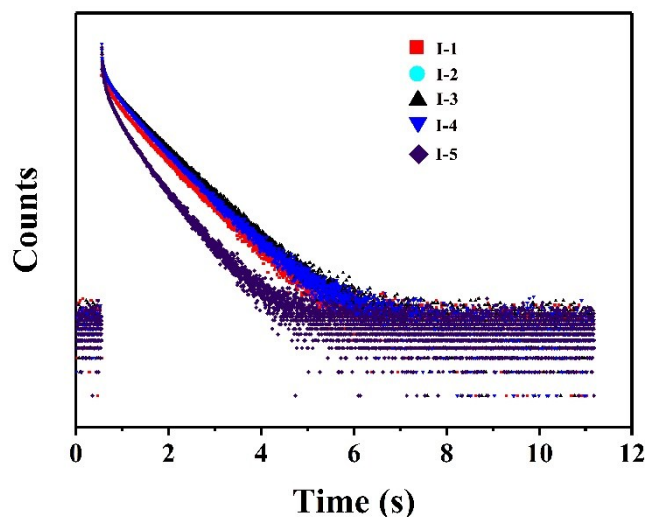
**Figure S4.** PXRD patterns of the CDs@MS nanocomposites from 1,2,4-triaminobenzene and inorganic salts ( $\text{KNO}_3$  and  $\text{NaCl}$ ) calcinated at  $350\text{ }^\circ\text{C}$  for 3 h.



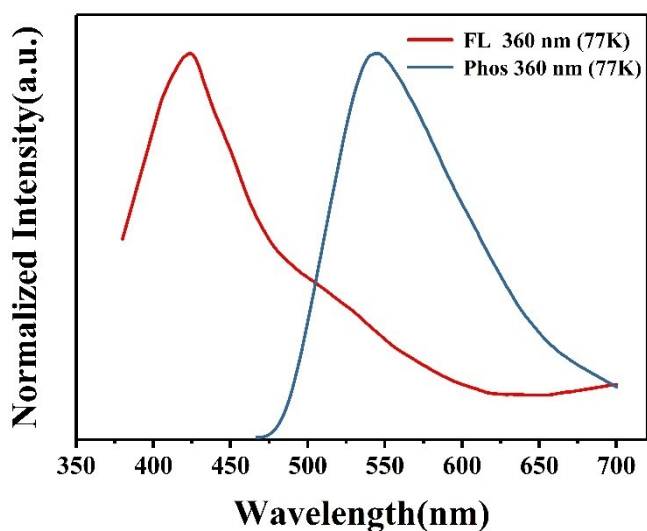
**Figure S5.** TEM image of CDs. (The CDs@MS nanocomposites were dissolved in water, and the salts were removed by a 500Da dialysis bag for 48 h. The morphology of the corresponding CDs nanoparticles in the dialysis bag were characterized by the transmission electron microscope (TEM), the resultant CDs were orderly arranged and possessed almost the average size of approximately 2.5 nm, which was in accordance with the result in Figure 4b.)



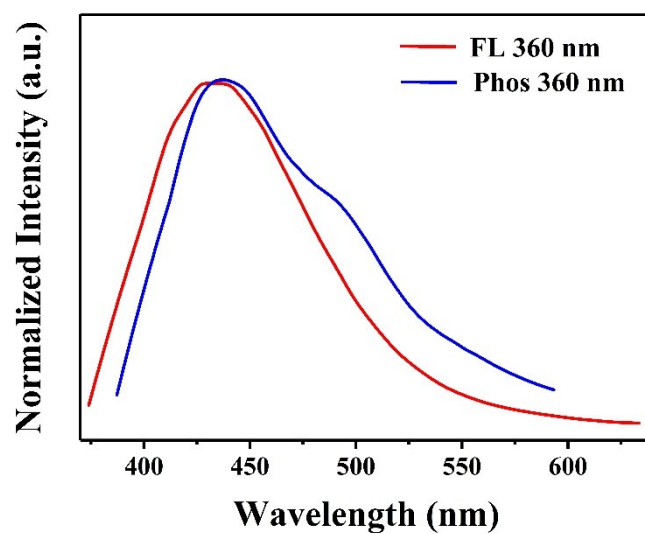
**Figure S6.** Fluorescence decay curve of the CDs@MS water dispersion.



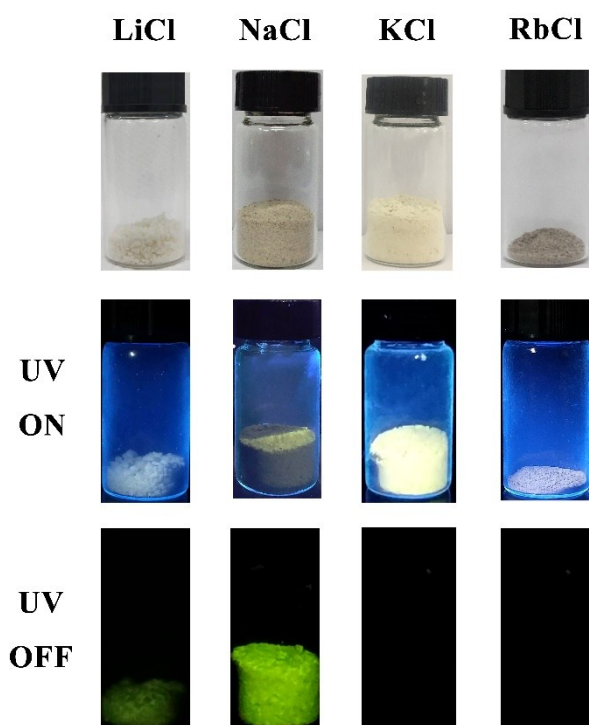
**Figure S7.** RTP decay spectra of the CDs@MS nanocomposites synthesized from 1,2,4-triaminobenzene via MS method by different inorganic salts systems calcinated at 350 °C for 3 h. (I-1: inorganic salts systems:  $\text{KNO}_3$  and  $\text{NaCl}$ , lifetime: 701 ms; I -2: inorganic salts systems:  $\text{KNO}_3$  and  $\text{KCl}$ , lifetime: --; I-3: inorganic salts systems:  $\text{NaNO}_3$  and  $\text{NaCl}$ , lifetime: 685 ms; I-4: inorganic salts systems:  $\text{NaNO}_3$ , lifetime: 469 ms; I -5: inorganic salts systems:  $\text{NaNO}_3$  and  $\text{KCl}$ , lifetime: 692 ms.)



**Figure S8.** Low temperature (77 K) FL and phosphorescence spectra of the CDs-based nanomaterials from 1,2,4-triaminobenzene and inorganic salts ( $\text{KNO}_3$  and  $\text{KCl}$ ) calcinated at 350 °C for 3 h.

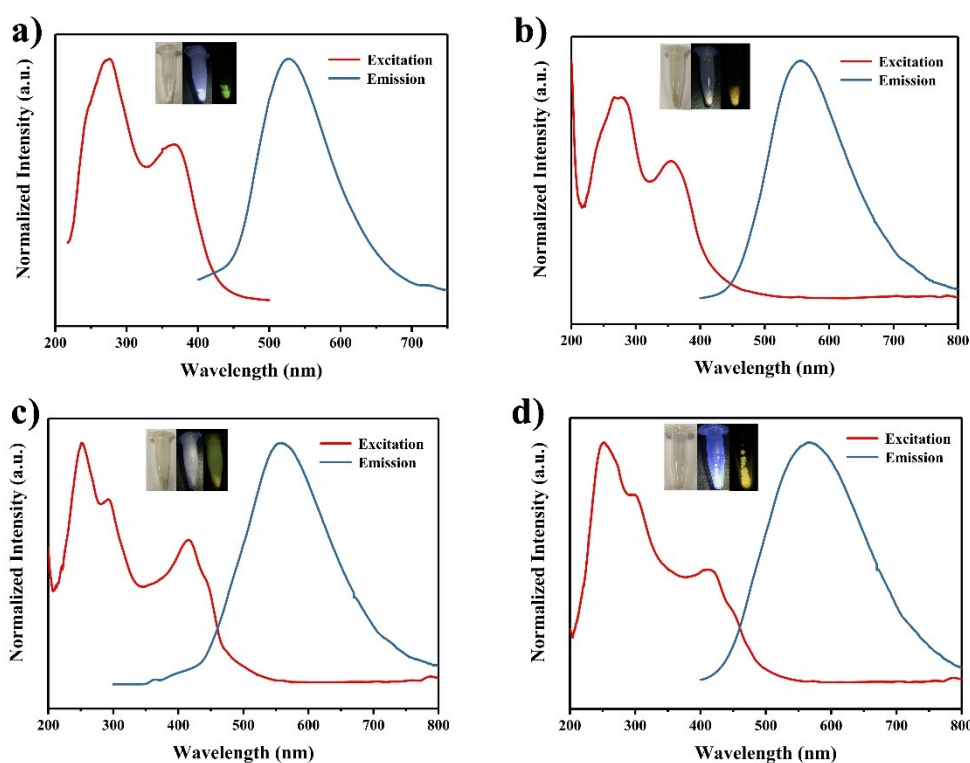


**Figure S9.** Low temperature (77 K) FL and phosphorescence spectra of the CDs-based nanomaterials from 1,2,4-triaminobenzene and inorganic salts ( $\text{KNO}_3$  and  $\text{LiCl}$ ) calcinated at  $350\text{ }^\circ\text{C}$  for 3 h.

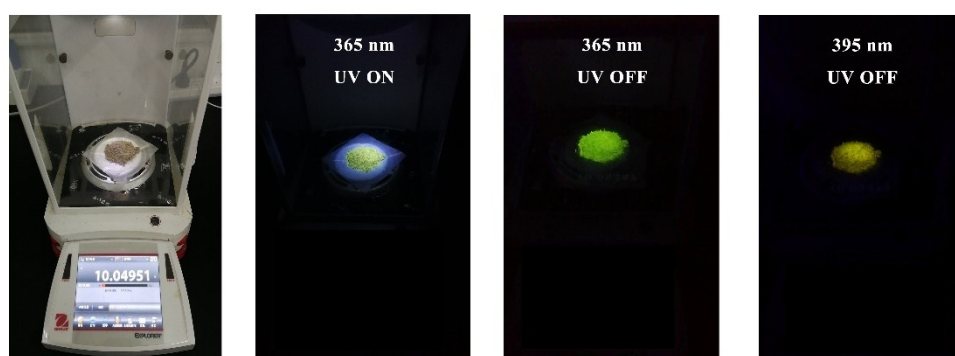


**Figure S10.** Photographs of CDs@MS nanocomposites synthesized from 1,2,4-triaminobenzene via MS method by different inorganic salts systems calcinated at 350 °C for 3 h under daylight, 365 nm UV lamp ON and OFF, respectively.

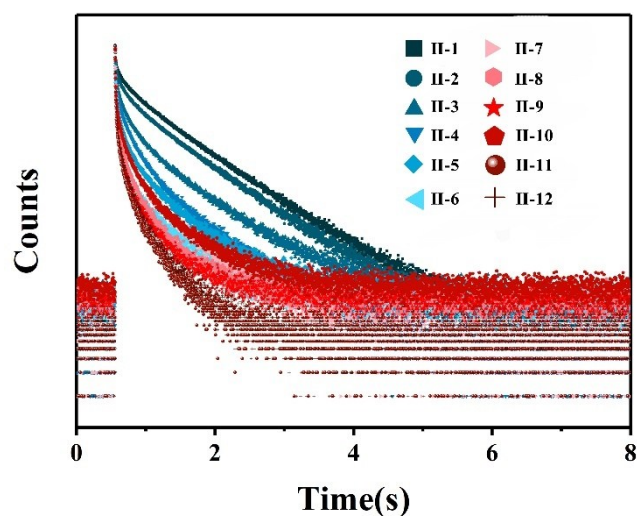




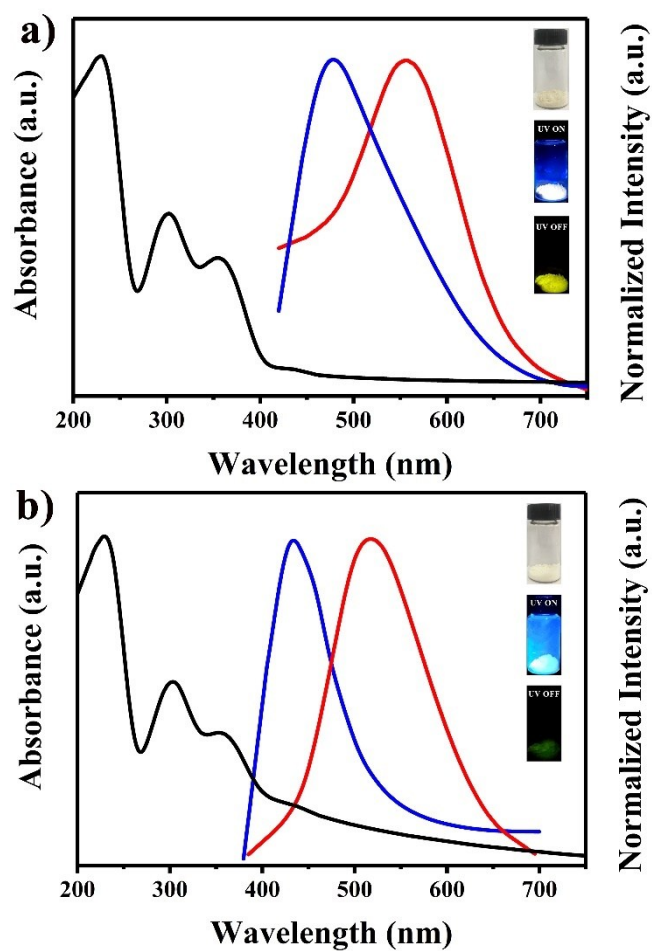
**Figure S11.** Phosphorescence spectra of the CDs@MS nanomaterials synthesized from 1,2,4-triaminobenzene via MS method by doping  $KNO_3$  with a)  $MgCl_2$ , b)  $CaCl_2$ , c)  $SrCl_2$  and d)  $BaCl_2$ , respectively. Insets: Images of the CDs@MS nanocomposites under daylight, corresponding UV lamp ON and OFF.



**Figure S12.** Schematic illustration of large-scale preparation of the CDs@MS nanocomposites from 1,2,4-triaminobenzene and inorganic salts ( $KNO_3$  and  $NaCl$ ) calcinated at  $350\text{ }^\circ\text{C}$  for 3 h.



**Figure S13.** RTP decay spectra of the CDs@MS nanocomposites synthesized by different precursors via MSM from same inorganic salts ( $\text{KNO}_3$  and  $\text{NaCl}$ ) calcinated at  $350\text{ }^\circ\text{C}$  for 3 h. (II -1:carbon source:1,2,4-triaminobenzene, lifetime: 701 ms; II -2:carbon source: p-phenylenediamine, lifetime: 619 ms; II -3:carbon source: ethylenediaminetetraacetic acid, lifetime: 496 ms; II -4:carbon source: benzimidazole, lifetime: 432 ms; II -5:carbon source: m-phenylenediamine, lifetime: 428 ms; II -6:carbon source: L-lysine, lifetime: 380 ms; II -7:carbon source: chitosan, lifetime: 351 ms; II -8:carbon source: 4-aminobenzoic acid, lifetime: 335 ms; II -9:carbon source: citric acid, lifetime: 224 ms; II -10:carbon source: glucose, lifetime: 210 ms; II -11:carbon source: ascorbic acid, lifetime: 206 ms; II -12:carbon source: dopamine hydrochloride, lifetime: 203 ms.)



**Figure S14.** The UV-vis absorbance (black line), FL emission (deep blue line) and phosphorescence emission (red line) of the a) CD<sub>s-II-9</sub> nanocomposites excited under 400 nm and b) CD<sub>s-II-3</sub> nanocomposites excited under 365 nm. Insets: Images of the CD<sub>s</sub>@MS nanocomposites under daylight, corresponding UV lamp ON and OFF.

**Table S1.** Comparison of RTP property from different CDs-based RTP materials.

CDs-based RTP materials	Phosphorescent emission wavelength (nm)	Phosphorescent lifetime (ms)	Reference
CDs@zeolite	430	350	22
m-CDs-PVA	506	456	24
CQDs/PU	450-500	8.7	26
PCDs	494	658.11	27
CDs/PVA	500	380	34
CDs-LDH	525	386.8	37
ACDs/PVA	450	450	38
P-CDs	538	1390	39
URTP CDs	535	1460	40
CDs@MS	566	886	this work