Serratia marcescens-derived fluorescent carbon dots as the platform toward multimode bioimaging and detection of *p*-nitrophenol

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Fig. S1 (A): Normalized fluorescence intensity of CDs-KMR3 in different NaCl concentrations ranging from 0 to 2 M. (B): The fluorescence intensity of CDs-KMR3 varying with the sample pH value from 2 to 10. (C): Normalized fluorescence intensity of CDs-KMR3 under UV (365 nm) irradiation for 3 h. (D): Effect of reaction time on

the detection of *p*-NP with CDs-KMR3.



Fig. S2 Hela cell viability from MTT assays with different CDs-KMR3 concentration after 24 h incubation.



Fig. S3 CLSM images of Hela cells incubated with CDs-KMR3 under different excitation wavelengths.