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2018

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Cite this: *RSC Adv.*, 2018, **8**, 17254

Photoluminescence mechanism and applications of Zn-doped carbon dots†

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Heteroatom-doped carbon dots (CDs) with excellent optical characteristics and negligible toxicity have emerged in many applications including bioimaging, biosensing, photocatalysis, and photothermal therapy. The metal-doping of CDs using various heteroatoms results in an enhancement of the photophysics but also imparts them with multifunctionality. However, unlike nonmetal doping, typical metal doping results in low fluorescence quantum yields (QYs), and an unclear photoluminescence mechanism. In this contribution, we detail results concerning zinc doped CDs (Zn-CDs) with QYs of up to 35%. The zinc ion charges serve as a surface passivating agent and prevent the aggregation of graphene π - π stacking, leading to an increase in the QY of the Zn-CDs. Structural and chemical investigations using spectroscopic and first principle simulations further revealed the effects of zinc doping on the CDs. The robust Zn-CDs were used for the ultra-trace detection of Hg^{2+} with a detection limit of 0.1 μM , and a quench mechanism was proposed. The unique optical properties of the Zn-CDs have promise for use in applications such as *in vivo* sensing and future phototherapy applications.

Received 30th March 2018

Accepted 26th April 2018

DOI: 10.1039/c8ra02756k

rsc.li/rsc-advances

Introduction

Carbon dots (CDs) have become one of the most important basic carbon nano materials after the discovery of carbon nanotubes, nano diamonds, fullerenes and graphene.¹ Compared with semiconductor quantum dots, CDs with their lower toxicity, lower cost and better biocompatibility are widely used in many fields including in bioimaging,² photodynamic therapy,³⁻⁶ light-emitting diodes,⁷⁻¹⁰ and photocatalysis.¹¹⁻¹⁷ Numerical approaches have been developed to increase the photoluminescence quantum yield (PLQY or QY) of CDs using surface passivation, edge functionalization, and heteroatom-doping.¹⁸ As a result, sulfur-doped CDs have been developed with a QY of up to 67%¹⁹ and nitrogen-doped CDs have been created with a QY of up to 80%.² Nitrogen and sulfur co-doped

semi-crystalline CDs (S, N-CDs) with a QY of up to 55% have also been successfully produced.²⁰ When compared to non-metal doped CDs, metal-ion doped CDs have been poorly explored. Metallic elements can serve as electron donors in CDs, boost electron transfer and introduce novel functionalities into the CDs. For example, Wu *et al.* reported that Cu could serve as an electron transfer booster in CDs and be used in photocatalysis.²¹ Bourlinos *et al.* created Gd(III)-doped CDs that could be used as dual fluorescence-MRI probes for biomedical applications.²² Thus far, most metal-doped CDs have shown a low PLQY. An in-depth understanding of the PL mechanism of CDs is beneficial for tuning their fluorescence properties, which in turn can maximize their application.

Factors including the CD size effect, the surface passivation functionalization and the conjugate effect are considered to be important factors in tuning the PLQY of CDs. However, the explanations behind the mechanism responsible for PLQY remain contradictory.²³⁻²⁷ The photoluminescence (PL) properties of CDs vary with size, with small, medium, and large CDs emitting UV, visible light and near-infrared wavelengths, respectively.²⁸ Guo *et al.* reported that the tuneable PL mechanism of CDs was due to the variations in the surface state affected by the N content. Yang *et al.* summarized PL mechanisms in a review by comparing graphene quantum dots, carbon nanodots, and polymer dots. This comparison included a description of the quantum confinement effect or conjugated δ -domains, the surface state, the molecule state and the cross-link enhanced emission (CEE) effect.²⁹ Thus far, the mechanism

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8ra02756k



of metal-ion doped CDs has remained unclear. Metal doping may enhance both the electron donating/accepting abilities and can change CD surface charges, so further analysis of the QY of metal-doped CDs is crucial.

In this contribution, we analysed zinc-doped CDs to determine how the CD size and doping amount affected the PLQY of the CDs. The zinc ion charges serve as an important surface passivate agent and prevent the aggregation of the graphene π - π stacking. This led to the formation of sp³/sp² hybrid hybridization, which is a key step in increasing the QY of Zn-CDs. The prepared Zn-CDs had a QY of up to 35%, and the first principles calculations further proved this hypothesis. These findings can effectively guide scientists in the preparation of high QY metal-doped CDs for applications in the biomedical field.

Experimental section

Materials

Sodium citrate, citric acid, mercuric nitrate, hydrogen peroxide and metal salts used in the detecting application section were purchased from Aladdin Industrial Inc. Zinc chloride used for doping was obtained from Tianjin Guangfu Technology Development Co., Ltd. All solutions were prepared using deionized water obtained from equipment bought from Dongguanshi Nabaichuan Water-Treatment Equipment Co., Ltd.

Synthesis of the Zn-CDs

Zinc-doped carbon dots (Zn-CDs) were prepared using a facile one-step hydrothermal method, which was optimized in our previous work.³⁰ First, 25 mL of sodium citrate (0.1 M) was mixed with zinc chloride (0.05 M) and the mixture was poured into a 50 mL Teflon-lined stainless-steel autoclave. Then, samples of the product were placed in a drying cabinet at 185 °C for a reaction time varying from 1 h to 10 h (1 h, 2 h, 4 h, 6 h, 8 h, 10 h). The products were first filtered to remove carbon impurities, salts or other impurities. Then, the product solutions were further purified over a number of days using a dialysis membrane filter (0.22 mm) against ultrapure water, which was renewed every 10–12 h, until no Zn was detected in the ultrapure water.

Characterization

A FLS980 Fluorescence Spectrometer from Techcomp (China) Ltd was used to measure the photoluminescence quantum yield. The morphology of the prepared CDs was investigated using transmission electron microscopy (TEM) (Model JEM2100). The atomic force microscopy (AFM) images were measured using a Bruker Scan-Dimension-Icon System. The ultraviolet-visible (UV-Vis) absorption spectra of the samples were measured using a UV-Vis spectrometer (Jasco V-570). A Bruker Vertex 70v was employed for the measurement of the Fourier transform infrared (FTIR) spectra. The fluorescence measurements were carried out using a Cary Eclipse fluorescence spectrophotometer (Varian, Inc.). The solutions of the prepared CDs were placed in a quartz fluorescence cuvette. The

emission slits and excitation slits were set at 2.5 nm and 5 nm, respectively. Elemental analysis was performed using X-ray photoelectron spectroscopy (XPS) (ESCALAB 250 spectrometer). Raman spectroscopy was performed by employing a Horiba Jobin Yvon Xplora confocal Raman microscope at room temperature.

Simulation method

Density functional theory (DFT) was employed to study the electronic properties of the zinc-doped carbon dots (Zinc-CD). We chose the B3LYP hybrid method with the 6-311G basis set within the Gaussian 09 package to calculate the optimized structure of the Zn-CDs and their electronic properties. Three different sizes of Zn-CDs (C₁₈H₁₁Zn, C₂₄H₁₃Zn and C₃₀H₁₅Zn) were built representing the growing of the Zn-CDs at different reaction times ranging from a small size to a large size and the HOMO–LUMO of each model was calculated and discussed.

Detection of Hg²⁺

The typical steps of the detecting experiment were as follows: 20 μ L of the prepared Zn-CDs were placed into 2 mL of deionized water, and then the fluorescence intensity at 440 nm of the Zn-CDs (excited by light at 340 nm) was measured and recorded as the initial fluorescence intensity (F_0). Then, different concentrations of Hg²⁺ were introduced into the above sample solution. The fluorescence intensity was measured after 5 min and recorded as the fluorescence intensity of the end point (F_t). The change in the fluorescence intensity (ΔF) was defined as the difference between F_0 and F_t ($\Delta F = F_0 - F_t$). The linear relationship between the change in the fluorescence intensity ΔF ($\Delta F = F_0 - F_1$) and the concentration of Hg²⁺ was determined.

To explore the selectivity of the Hg²⁺ detection, 50 μ M solutions of various metal ions (including Hg²⁺, K⁺, Mg²⁺, Na⁺, Mn²⁺, Cu⁺, Cu²⁺, Al³⁺, Fe²⁺, Fe³⁺, Cl⁻, SO₄²⁻ and NO₃⁻) were prepared using deionized water. Then 20 μ L of each of the sample solutions were added to the aforementioned solution (2 mL), respectively, and the fluorescence intensity of the mixtures was measured and defined as F_t . The change in the fluorescence intensity, ΔF , was defined as: $\Delta F = F_0 - F_t$.

Results and discussion

Optimization of the synthesis conditions

The prepared Zn-CDs solution was observed to be a lavender colour under visible light and exhibited a bright blue luminescence when excited by light at 365 nm, as illustrated in Fig. 1a. The structure of the Zn-CDs was explored by transmission electron microscopy (TEM). Fig. 1a shows that the Zn-CDs were successfully synthesized by the one-step hydrothermal method, and the TEM showed the Zn-CDs to take the form of nanoparticles. The TEM images exhibited that there was an excellent separation of the particles, with a diameter range from 3–10 nm. The AFM images further demonstrated the formation of mostly monodispersed, quasi-spherical Zn-CDs with an average diameter of smaller than 3 nm, devoid of any aggregation (Fig. S2†).



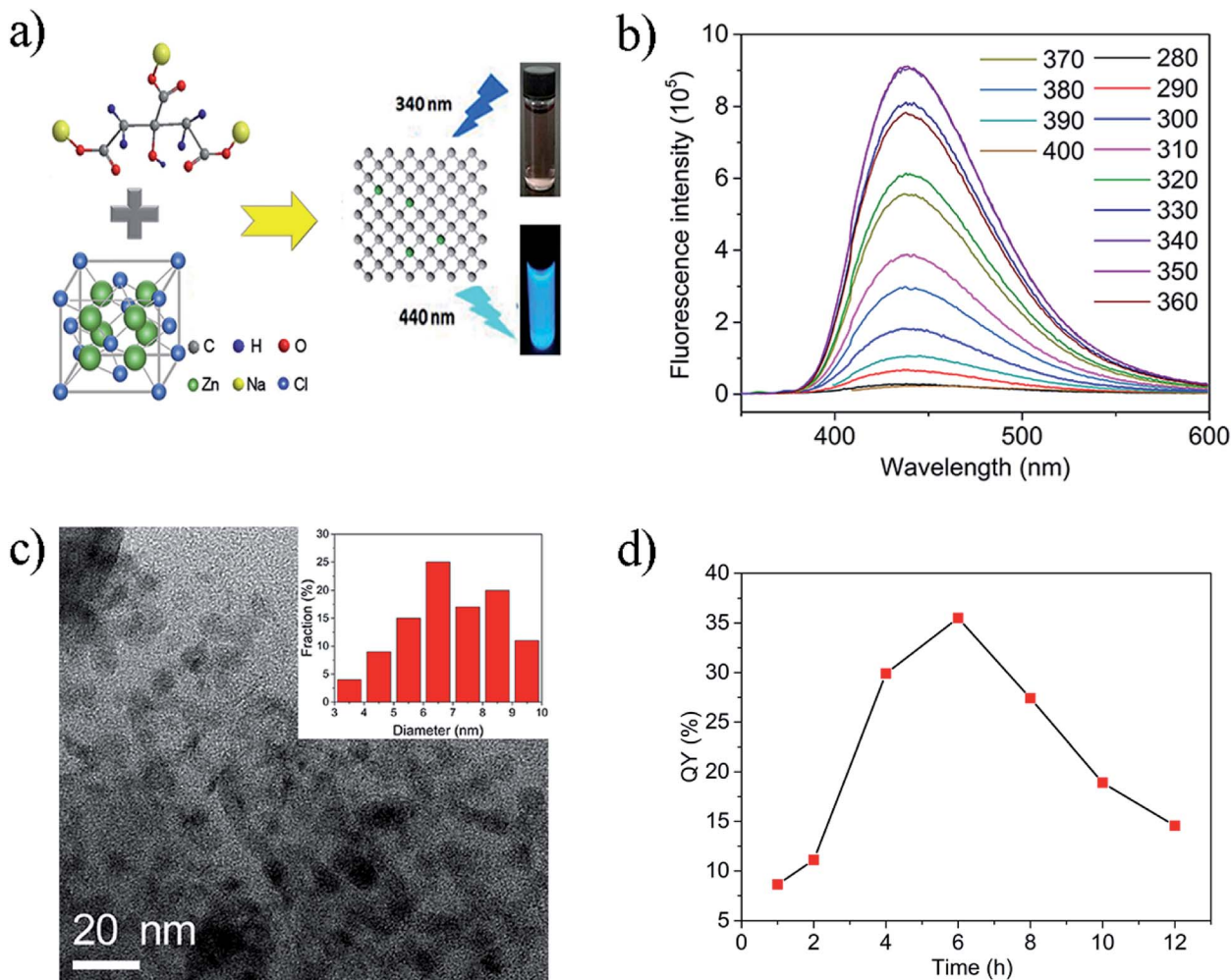


Fig. 1 (a) Schematic diagram of the synthesis of the Zn-CDs with blue luminescence (excited by light at 340 nm); (b) photoluminescence spectra of the Zn-CDs; (c) TEM images of the Zn-CDs and the diameter distribution of the Zn-CDs; (d) quantum yield of the Zn-CDs for reaction times varying from 1 h to 12 h (1 h, 2 h, 4 h, 6 h, 8 h, 10 h, 12 h).

The photoluminescence characteristics of the synthesized Zn-CDs were investigated in detail using a fluorescence spectrometer, which irradiated the samples at different wavelengths to analyze the emission properties. The prepared Zn-CDs had the strongest emission, centred at 440 nm (λ_{em}) when excited at a wavelength of 340 nm (λ_{ex}). This result is shown in Fig. 1b and S1.† The variation of λ_{ex} from 280 to 400 nm for the prepared samples only led to a rapid decrease in the intensity of λ_{em} , without a shift in the peak position, which was shown in our previous studies.³⁰ Fig. 1d shows that the quantum yield (QY) of the Zn-CDs varied relative to the heating time. The prepared Zn-CDs heated for 6 h had the highest quantum yield of 34.6%. Fig. S3† shows that the fluorescence lifetime of the Zn-CDs was measured to be 6.32 ns using the time-correlated single photon counting (TCSPC) method. The results showed that the lifetime of the Zn-CDs underwent almost no change as the reaction time increased, indicating little variety in the surface functional groups of the Zn-CDs.

In the C1s spectra (Fig. 2a), the two peaks at 288.5 eV and 284.5 eV are representative peaks of the C=O and C-C groups.

As the reaction time increased, the concentration of the C=O groups first increased and then decreased, indicating that the carbon dots experienced maximum oxidation after approximately 4 hours. This phenomenon was also observed in the O1s spectra (Fig. 2b). The carbonyl peak also exhibited an initial increase and then a decrease as the reaction continued, which was consistent with what was shown in the C1s spectra. Both results show that after undergoing a reaction for 2–4 h, the oxidation of the carbon dots reached its peak. This conclusion was further proved by the Raman spectra, shown in Fig. 2e. The D band is the indicator of the oxidation defect on the carbon dot surface and it exhibited a similar trend as seen in the C1s and O1s spectra. When normalized in the G band (1600 cm⁻¹), the D band increased at the beginning of the reaction (1, 2, 4 h), and decreased as the reaction continued (8, 10 h). According to the theory of passivation, a high D/G ratio implies that there is a high surface defect/passivation ratio, which can serve as an excitation energy trap and increase the photoluminescence. This provides a good explanation for the QY change seen in Fig. 1d. The effect of an extreme environment, such as highly



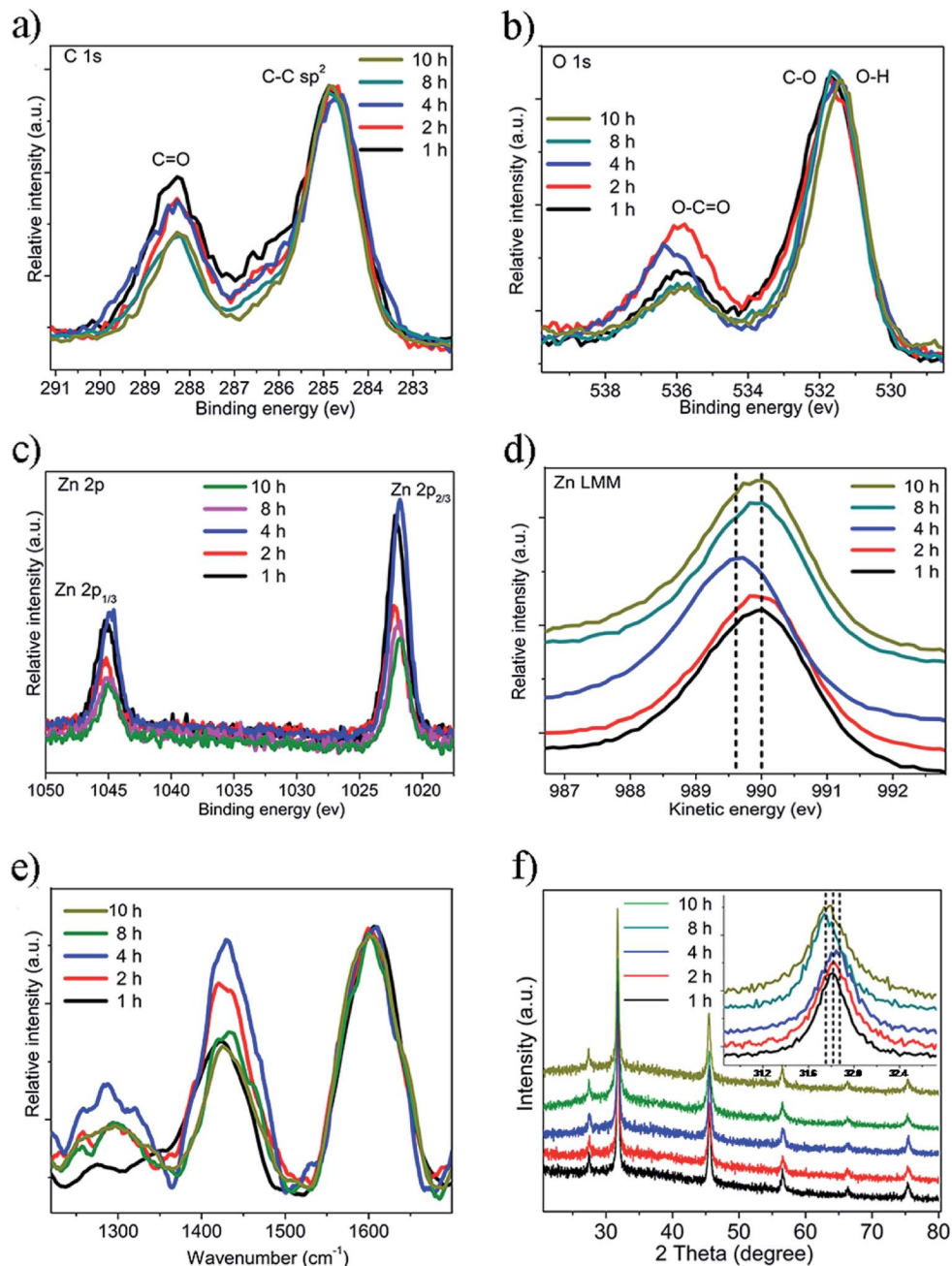


Fig. 2 (a) C1s, (b) O1s, (c) Zn2p, and (d) Zn LMM XPS spectra of the Zn-CDs; (e) Raman spectra and (f) PXRD patterns of the Zn-CDs.

oxidizing solutions (hydrogen peroxide), was also tested and the Zn-CDs manifested high stability and steady fluorescence against highly oxidizing H₂O₂ (Fig. S4†).

The Zn peak in the Zn LMM spectra shifted to a higher binding energy in the first part of the reaction and then shifted back to a lower binding energy (Fig. 2d). This indicated that during the growth phase of the carbon dots, that Zn was firstly oxidized and then reduced. These results combined with the O1s and C1s spectra showed a high possibility that the high oxidation state of the Zn was connected to the carbonyl group. It was observed that 2 theta in the Zn PXRD patterns exhibited a slight shift during the reaction time of 2–4 hours, indicating

the change in the lattice distance, which may have resulted from the high oxidation state of the Zn (Fig. 2f). The crystal structure of Zn-CDs was a lattice structure, which indicated an existence of crystallization characteristics of graphitic carbon due to a sp² hybridization (Fig. 3a–c). The lattice fringe spacing is shown in Fig. 3c, of which the interplanar distance of 0.22 nm corresponded to the (002) plane of Zn-CDs. It was observed that the images of the particles were darker, which indicated the growth of the crystalline Zn-CDs (Fig. 3d). Fig. 3b shows the areas partitioned by dotted lines had the existence of both the lattice structure and absolute amorphous structure. The existence of the amorphous structure indicated a sp³/sp² hybrid



hybridization, indicating the formation of a 3D structure. Considering that the high oxidation state of the Zn concurred with the high oxidation of the carbon dots, it is reasonable to assume that Zn may play an important role as the active binding site to connect neighbouring small carbon dots through chelation between Zn and the carbonyl group. This process is believed to indirectly promote the growth of large carbon dots (Fig. 3d). The change in the peaks in the UV-Vis absorption spectra (Fig. S5†) provided strong evidence that the cyclization degree of the Zn-CDs rose as the reaction progressed. The IR spectra in Fig. S6† show the existence of OH groups inside the

Zn-CDs, which corresponds to what is shown in the XPS results of O1s. It is assumed that the better solubility of the Zn-CDs in water was due to the polar OH groups on the surface of the CDs, which could form intermolecular hydrogen bonds in the solution.

Simulations

To further understand the Zn doping process and its effect on the CDs, we proposed a theoretical lattice model of the Zn-CDs. Both the structural and electronic properties were investigated as

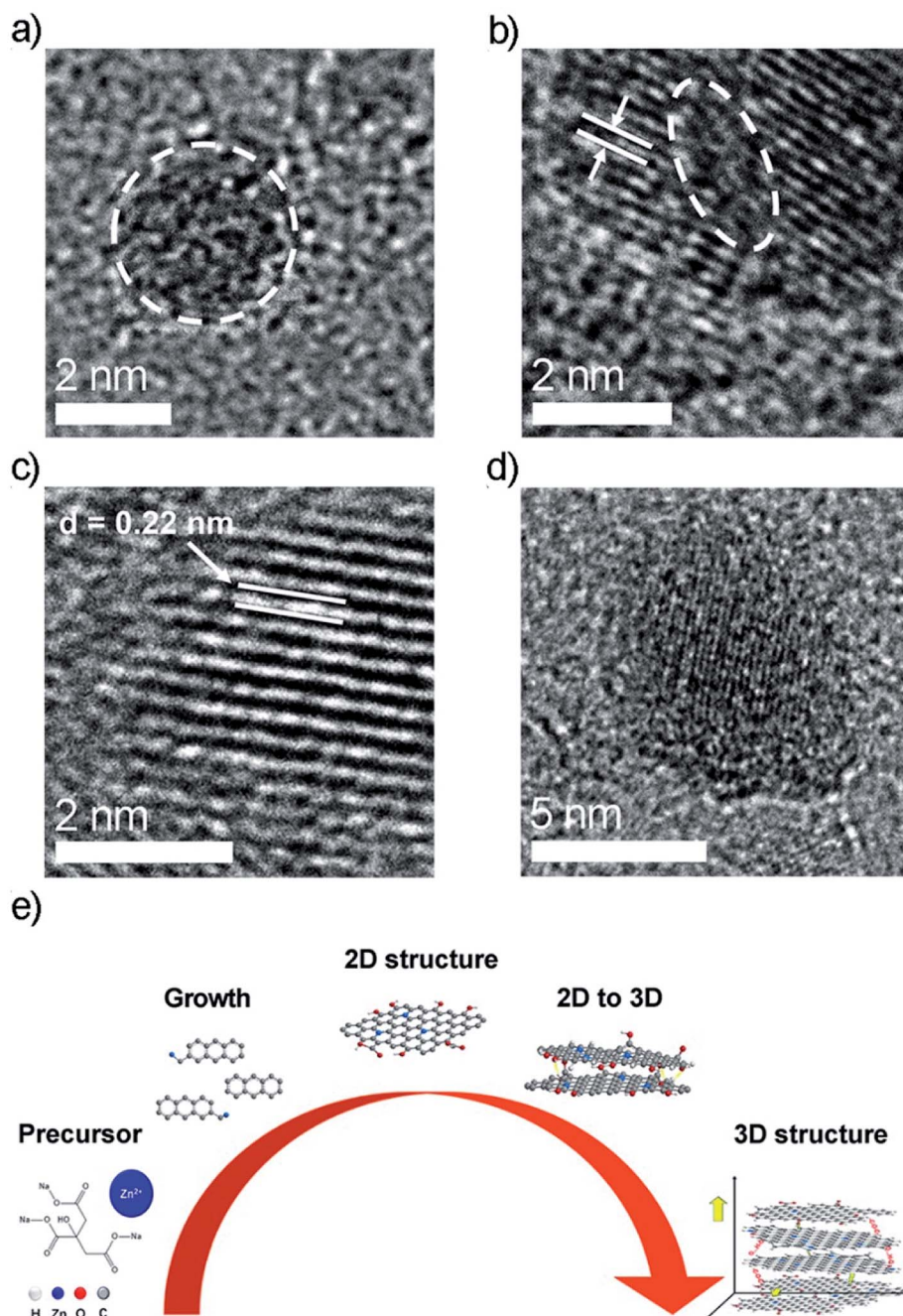


Fig. 3 (a–d) TEM images of the Zn-CDs with reaction times of 1 h, 2 h, 4 h and 10 h, respectively. The cross lattices show the 3D structure of the Zn-CDs. (e) A schematic diagram showing the growth of the Zn-CDs from a sp^2 to a sp^3 structure.



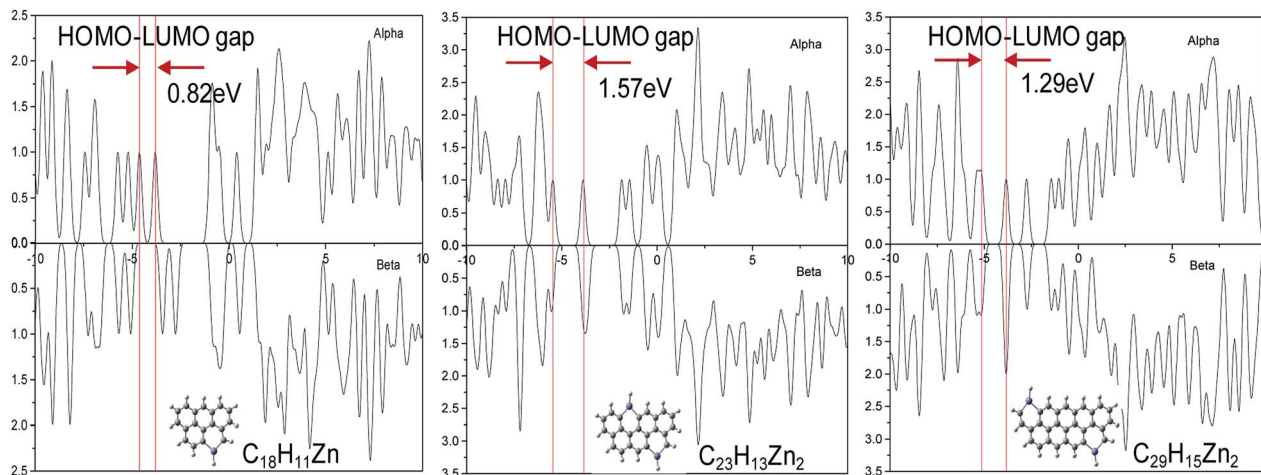


Fig. 4 Density of states for the different sizes of Zn-CDs.

functions of the location of impurity/dopant atoms incorporated into the CDs. Our results clearly showed that the location of the Zn atoms incorporated into the CD lattice plays an important role in regulating the properties of the doped CDs. The HOMO–LUMO energy gap of the three Zn-CD materials were calculated and were found to have values of 0.82 eV, 1.57 eV and 1.29 eV as the size of the Zn-CDs increased, as shown in Fig. 4. Among these three Zn-CDs, the concentration of oxidized Zn first increased and then decreased. The results clearly revealed that when comparing pristine-CDs with the Zn-CDs, that the doped Zn ions tune the HOMO–LUMO values and the energy gap. In three different sizes of pristine-CDs, the HOMO–LUMO energy gaps were found to be ~ 1.75 eV, as listed in Table 1. The doped Zn adjusts the HOMO–LUMO gap of the $C_{23}H_{13}Zn_2$ to a value of 1.57 eV, in which the concentration of the oxidized Zn group is neither lower nor higher than 1.57 eV. This value may correspond to the preferred radiation from the singlet excitation state S_1 to the ground state S_0 , therefore generating the high fluorescence QY. The simulation results revealed that doped Zn ions play an important role as the CDs grow from a small size to a medium size. They may increase charge transfer/charge recombination and thus increase the QY of the CDs with a certain concentration. However, the doping would cause a loss in function as the CDs kept growing as more graphene would π – π stack, and thus cause the QY of the CDs to decrease.

Detection of Hg^{2+}

To explore the potential application of Zn-CDs, fluorescence probes adopting Zn-CDs were studied for their detection of Hg^{2+} . The environmental hazards of mercury can eventually impact upon plants, animals, and human health due to its bioaccumulation. Mercury in the environment can be transferred through the food chain and greatly threaten human health. Even very low concentrations can cause diseases including kidney failure, permanent brain damage, and various motion and cognitive disorders.^{31,32} The most frequently used technologies for Hg^{2+} detection, ultraviolet-visible spectroscopy (UV-Vis) and inductively coupled plasma mass spectrometry

(ICP-MS), usually demands expensive equipment and complicated procedures. Here, we report a fluorescence-based detection method, which could be employed for Hg^{2+} detection with good sensitivity, instant response, high selectivity, and simple operation.

In this report, synthesized Zn-CDs were found to be efficient for detecting Hg^{2+} . Once 10 μM of Hg^{2+} (20 μL) was introduced to the Zn-CDs, an absolute drop in the fluorescence intensity was observed (Fig. 5a). Through an alternate and efficient electron transfer process, Hg^{2+} caused the efficient quenching of Zn-CDs due to its ability to facilitate electron/hole recombination annihilation. When the Hg^{2+} solution was introduced, a substantial decay was observed. Fig. 5b and c show the trend in the decay of the fluorescence intensity of the Zn-CDs as the concentrations of Hg^{2+} increased. Each spectrum was recorded after 5 min to keep an equilibration reaction time. A linear relationship was found between the Hg^{2+} concentration and the fluorescence intensity of the Zn-CDs. The resultant calibration curve for the Hg^{2+} had excellent linearity in a concentration range of 1 to 60 μM , with a coefficient of determination of 0.9894 (Fig. 5c). The limit of detection was 100 nM. To explore the quenching selectivity of the Zn-CDs for Hg^{2+} as a sensing probe, the detection was carried out after introducing various metal ions. In Fig. 5d, it can be seen that Hg^{2+} had the highest influence on the fluorescence intensity of the Zn-CDs among all of the metal ions. Moreover, the changes in the lifetime (Fig. 5e) and absorption (Fig. 5f) provided evidence that the change in the surface state of the Zn-CDs was brought about by Hg^{2+} .

Table 1 The HOMO–LUMO energy gaps of the Zn-CDs

Zn-CDs/CDs	E_{HOMO} (eV)	E_{LUMO} (eV)	ΔE_{gap} (eV)
$C_{18}H_{11}Zn$	−4.61	−3.79	0.82
$C_{19}H_{11}$	−4.34	−2.59	1.75
$C_{23}H_{13}Zn_2$	−3.89	−5.46	1.57
$C_{25}H_{13}$	−4.37	−2.64	1.73
$C_{29}H_{15}Zn_2$	−5.12	−3.83	1.29
$C_{31}H_{15}$	−4.41	−2.66	1.75



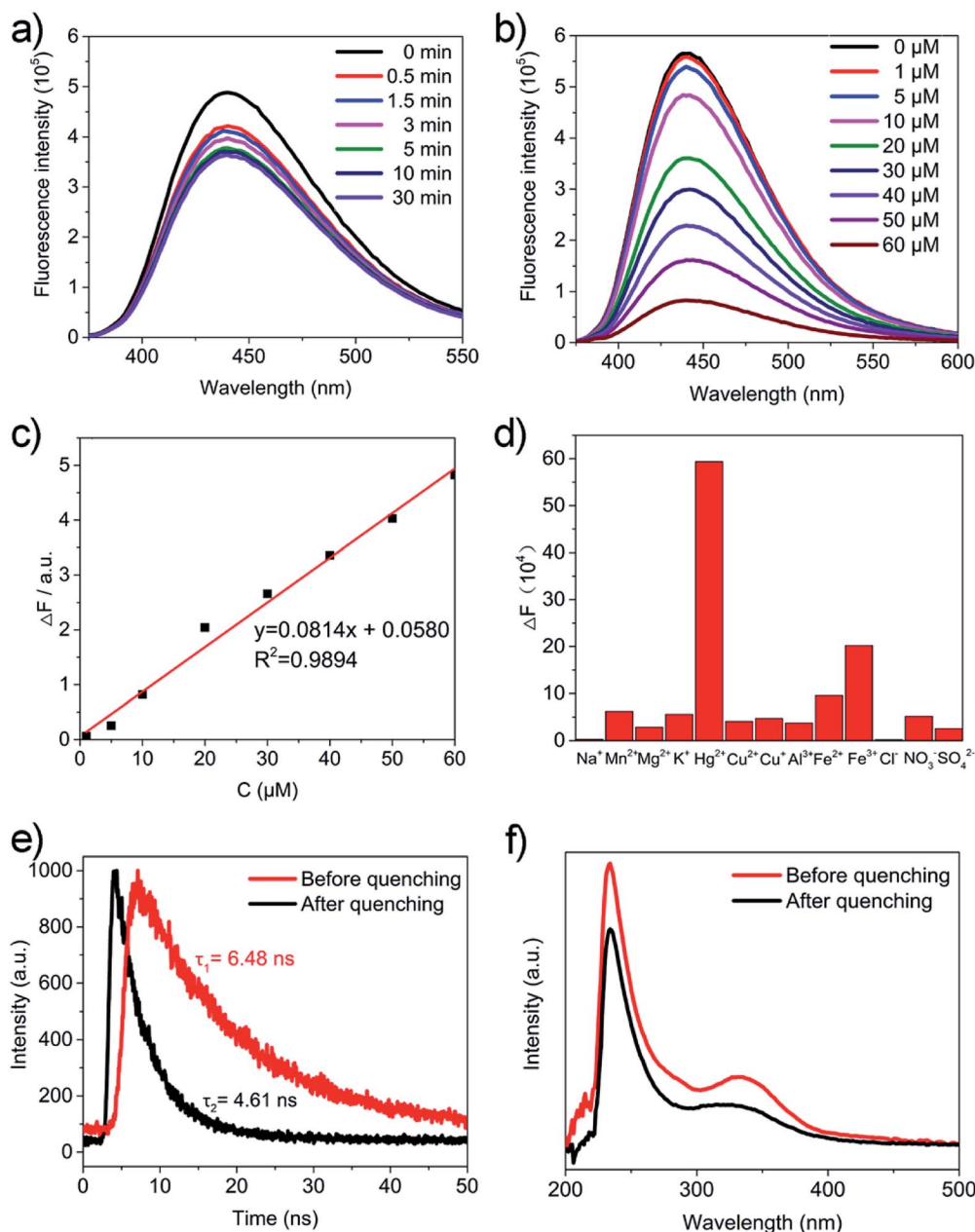


Fig. 5 (a) Time-dependent fluorescence changes of the Zn-CDs in the presence of Hg^{2+} ($10 \mu\text{M}$); (b) PL emission spectra of the Zn-CDs solution with different concentrations of Hg^{2+} ($0, 1, 5, 10, 20, 30, 40, 50, 60 \mu\text{M}$); (c) ΔF calibration curve of the Zn-CDs solution versus the concentration of Hg^{2+} ; (d) the ΔF at 440 nm for the Zn-CDs in the presence of various metal ions and anions ($50 \mu\text{M}$); (e) lifetimes and (f) absorption spectra before and after the quenching of Zn-CDs by Hg^{2+} .

There was a complexation as Hg^{2+} was introduced, which caused electron or energy transfer.^{33,34} Thus, the Zn-CDs were found to have a good specificity for the detection of Hg^{2+} . Based on the quenching effect of mercury ions on the fluorescence of CQDs, our group previously used S, N-doped CDs for the selective and sensitive sensing of Hg^{2+} in the aqueous phase with a limit of detection (LOD) of 100 pM .²⁰ As for metal-doped CDs, Liu *et al.* reported Mg, N-doped CDs for the sensing of Hg^{2+} with a LOD of $0.2 \mu\text{M}$,³⁵ and Yuan *et al.* reported Ge-doped CDs with a LOD of $0.075 \mu\text{M}$.³⁶ However, the Zn-CDs we have reported showed a LOD of $0.1 \mu\text{M}$, which is well placed among the LOD values of the aforementioned materials. It shows the great

potential of the Zn-CDs for application in sensing and detecting. Moreover, the PL properties of the Zn-CDs in $0.1 \text{ M H}_2\text{O}_2$ and water (Fig. S6†) were investigated and found to be quite stable. The results show that the Zn-CDs have excellent potential for use in the protection and monitoring of the environment.

Conclusions

In this report, we first determined how the heating time affects the growth of Zn-CDs. The degree of graphitization had a non-negligible effect on the fluorescence mechanism



and PLQY of the Zn-CDs. The XPS results showed the formation of C–Zn bonds in the Zn-CDs, strongly proving that the successful doping of Zn contributed to the emission of the bright blue fluorescence. Moreover, the Zn-CDs showed a good selectivity for Hg²⁺ with a detection limit of 0.1 μM. This shows that Zn-CDs could be employed in environmental monitoring. This work concerning the growth mechanism of Zn-CDs provides guidance for the controllable preparation and functional design of CDs. This study assists in determining the factors responsible for the advantageous synthesis, heteroatom doping, and functionalization of carbon dots for use in the environmental and biomedical fields.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank the National Key Research and Development Plan (No. 2016YFC0303701), the Beijing Nova Program Interdisciplinary Studies Cooperative project (No. Z181100006218138), the Beijing Science and Technology project (No. Z161100001316010), the State Key Laboratory of Petroleum & Petrochemical Pollution Control and Treatment (No. PPC2017015), the Science Foundation of the China University of Petroleum, Beijing (2462018BJC004) for support. This study is supported by the National Institute of Food and Agriculture, the U.S. Department of Agriculture, and McIntire Stennis under accession number 1009735.

References

- 1 S. N. Baker and G. A. Baker, *Angew. Chem., Int. Ed. Engl.*, 2010, **49**, 6726–6744.
- 2 S. Zhu, Q. Meng, L. Wang, J. Zhang, Y. Song, H. Jin, K. Zhang, H. Sun, H. Wang and B. Yang, *Angew. Chem., Int. Ed. Engl.*, 2013, **125**, 4045–4049.
- 3 J. Ge, M. Lan, B. Zhou, W. Liu, L. Guo, H. Wang, Q. Jia, G. Niu, X. Huang, H. Zhou, X. Meng, P. Wang, C. S. Lee, W. Zhang and X. Han, *Nat. Commun.*, 2014, **5**, 4596–4603.
- 4 B. Zhang, J. Wang, J. Yu, X. Fang, X. Wang and D. Shi, *Bioconjugate Chem.*, 2017, **28**, 330–335.
- 5 J. Wang, Y. Zhong, X. Wang, W. Yang, F. Bai, B. Zhang, L. Alarid, K. Bian and H. Fan, *Nano Lett.*, 2017, **17**, 6916–6921.
- 6 W. Yang, W. Guo, W. Le, G. Lv, F. Zhang, L. Shi, X. Wang, J. Wang, S. Wang, J. Chang and B. Zhang, *ACS Nano*, 2016, **10**, 10245–10257.
- 7 F. Yuan, Z. Wang, X. Li, Y. Li, Z. a. Tan, L. Fan and S. Yang, *Adv. Mater.*, 2017, **29**, 1604436–1604441.
- 8 D. H. Kim and T. W. Kim, *Nano Energy*, 2017, **32**, 441–447.
- 9 L. Cui, X. Wang, N. Chen, B. Ji and L. Qu, *Nanoscale*, 2017, **9**, 9089–9094.
- 10 L. F. Cui, X. P. Wang, N. Chen, G. F. Zhang and L. T. Qu, *J. Mater. Chem. A*, 2017, **5**, 14508–14513.
- 11 S. Fang, Y. Xia, K. Lv, Q. Li, J. Sun and M. Li, *Appl. Catal., B*, 2016, **185**, 225–232.
- 12 Q. Han, B. Wang, J. Gao and L. Qu, *Angew. Chem., Int. Ed. Engl.*, 2016, **55**, 10849–10853.
- 13 X. Y. Kong, W. L. Tan, B.-J. Ng, S.-P. Chai and A. R. Mohamed, *Nano Res.*, 2017, **10**, 1720–1731.
- 14 P. Wang, X. Zhang, J. Zhang, S. Wan, S. Guo, G. Lu, J. Yao and X. Huang, *Nat. Commun.*, 2017, **8**, 14580–14588.
- 15 R. Wang, K.-Q. Lu, Z.-R. Tang and Y.-J. Xu, *J. Mater. Chem. A*, 2017, **5**, 3717–3734.
- 16 J. Chen, J. S. Wei, P. Zhang, X. Q. Niu, W. Zhao, Z. Y. Zhu, H. Ding and H. M. Xiong, *ACS Appl. Mater. Interfaces*, 2017, **9**, 18429–18433.
- 17 H. Li, X. He, Z. Kang, H. Huang, Y. Liu, J. Liu, S. Lian, C. H. Tsang, X. Yang and S. T. Lee, *Angew. Chem., Int. Ed. Engl.*, 2010, **49**, 4430–4434.
- 18 C. J. Reckmeier, J. Schneider, A. S. Sussha and A. L. Rogach, *Opt. Express*, 2016, **24**, A312–A340.
- 19 Q. Xu, P. Pu, J. Zhao, C. Dong, C. Gao, Y. Chen, J. Chen, Y. Liu and H. Zhou, *J. Mater. Chem. A*, 2015, **3**, 542–546.
- 20 Q. Xu, Y. Liu, C. Gao, J. Wei, H. Zhou, Y. Chen, C. Dong, T. S. Sreeprasad, N. Li and Z. Xia, *J. Mater. Chem. C*, 2015, **3**, 9885–9893.
- 21 W. Wu, L. Zhan, W. Fan, J. Song, X. Li, Z. Li, R. Wang, J. Zhang, J. Zheng, M. Wu and H. Zeng, *Angew. Chem., Int. Ed. Engl.*, 2015, **54**, 6540–6544.
- 22 A. B. Bourlinos, A. Bakandritsos, A. Kouloumpis, D. Gournis, M. Krysmann, E. P. Giannelis, K. Polakova, K. Safarova, K. Hola and R. Zboril, *J. Mater. Chem.*, 2012, **22**, 23327–23330.
- 23 Q. Xu, T. Kuang, Y. Liu, L. Cai, X. Peng, T. Sreenivasan Sreeprasad, P. Zhao, Z. Yu and N. Li, *J. Mater. Chem. B*, 2016, **4**, 7204–7219.
- 24 F. Zhao, L. Wang, Y. Zhao, L. Qu and L. Dai, *Adv. Mater.*, 2017, **29**, 1604972–1604978.
- 25 Q. Yang, Z. Ma, H. Wang, B. Zhou, S. Zhu, Y. Zhong, J. Wang, H. Wan, A. Antaris, R. Ma, X. Zhang, J. Yang, X. Zhang, H. Sun, W. Liu, Y. Liang and H. Dai, *Adv. Mater.*, 2017, **29**, 1605497–1605505.
- 26 S. Sun, L. Zhang, K. Jiang, A. Wu and H. Lin, *Chem. Mater.*, 2016, **28**, 8659–8668.
- 27 X. Li, Y. Liu, X. Song, H. Wang, H. Gu and H. Zeng, *Angew. Chem., Int. Ed. Engl.*, 2015, **54**, 1759–1764.
- 28 L. Guo, J. Ge, W. Liu, G. Niu, Q. Jia, H. Wang and P. Wang, *Nanoscale*, 2016, **8**, 729–734.
- 29 S. Zhu, Y. Song, X. Zhao, J. Shao, J. Zhang and B. Yang, *Nano Res.*, 2015, **8**, 355–381.
- 30 Q. Xu, Y. Liu, R. Su, L. Cai, B. Li, Y. Zhang, L. Zhang, Y. Wang, Y. Wang, N. Li, X. Gong, Z. Gu, Y. Chen, Y. Tan, C. Dong and T. S. Sreeprasad, *Nanoscale*, 2016, **8**, 17919–17927.
- 31 H. H. Harris, I. J. Pickering and G. N. George, *Science*, 2003, **301**, 1203.
- 32 E. M. Nolan and S. J. Lippard, *Chem. Rev.*, 2008, **108**, 3443–3480.
- 33 Y. Dong, R. Wang, H. Li, J. Shao, Y. Chi, X. Lin and G. Chen, *Carbon*, 2012, **50**, 2810–2815.



- 34 L. Wang, B. Li, F. Xu, X. Shi, D. Feng, D. Wei, Y. Li, Y. Feng, Y. Wang, D. Jia and Y. Zhou, *Biosens. Bioelectron.*, 2016, **79**, 1–8.
- 35 T. Liu, N. Li, J. X. Dong, H. Q. Luo and N. B. Li, *Sens. Actuators, B*, 2016, **231**, 147–153.
- 36 Y. H. Yuan, R. S. Li, Q. Wang, Z. L. Wu, J. Wang, H. Liu and C. Z. Huang, *Nanoscale*, 2015, **7**, 16841–16847.

