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Fluorescence enhancement of carbon dots by graphene for highly sensitive detection of tetracycline hydrochloride†

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In this study, fluorescence of self-assembled multilayers (SAMs) of carbon dots (CDs) was found to be enhanced by graphene. The number of polyelectrolyte layers can be tuned to control the distance between CDs and graphene in SAMs. The enhanced fluorescence efficiency was found to be dependent on graphene concentration, degree of graphene reduction, and the distance between CDs and graphene surface. When graphene concentration and polyelectrolyte bilayer number were set to 0.005 mg mL⁻¹ and 3 layers, respectively, fluorescence intensity of CDs could be increased up to 3.2 times. Tetracycline hydrochloride (Tc) could be detected by the established SAMs in the presence of graphene as the sensitivity was 2 orders higher than that of SAMs in the absence of graphene. Limit of detection of this sensing system was 0.9284 nM, which is 1–3 orders of magnitude lower than those of most of reported fluorescence sensors. This method could be successfully applied to detect trace Tc in milk samples.

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Introduction

Tetracycline hydrochloride (Tc) is a well-known antibacterial drug, widely used in pharmaceuticals, food and environmental fields.^{1,2} Although Tc is beneficial for treatment of some diseases, its residues in food may lead to some allergic reactions. According to European Union law, the maximum amount of tetracycline hydrochloride in milk must not exceed 100 ng mL⁻¹ (ca. 225 nM).³ Therefore, it is desirable to establish a highly sensitive and simple method to detect Tc.

As an important and effective analytical tool, fluorescence analysis has been widely used in the fields of life sciences, medicine and environmental monitoring.^{4–6} However, fluorescence quantum yields of some fluorescent materials are low, resulting in poor detection sensitivity, thus limiting their applications in fluorescence analysis. Plasmon-enhanced fluorescence (PEF) effect can greatly enhance fluorescence intensity of probes, leading to significantly increased sensitivity of detection, thus expanding the scope of applications.^{7,8}

In recent years, studies on PEF materials mainly focused on metal nanoparticles, such as AgNPs and AuNPs.^{9,10} However, these materials have several shortcomings, such as poor light stability, low biocompatibility and relatively large optical loss. These shortcomings limit the application scope of PEF sensors. Therefore, it is necessary to explore novel materials that are

cheap, stable and biocompatible for use in surface enhanced fluorescence.

Graphene, a rising star in the world of materials, is widely used in various fields because of its outstanding electronic and optical properties.¹¹ One of these properties is that under certain conditions, graphene can produce surface plasmon, similar to that of metal nanoparticles.¹² In addition, graphene surface plasmon has the following excellent properties: it has extremely high wave momentum; it is a strong longitudinal local plasmon; it has infrared to terahertz band work area; and it is highly adjustable compared to metal surface plasmon.^{13,14} It should be pointed out that there many reports have focused on enhanced Raman spectroscopy, but reports involving enhanced fluorescence spectroscopy are rare.^{15–17} Until now, only a few examples involving the study of the phenomenon of enhanced fluorescence by graphene have been reported, including studies on fluorescence enhancement of GeSi quantum dots, SnO₂ nanostructures and ZnO nanostructures by graphene.^{18–20}

Self-assembled multilayers (SAMs) can be easily prepared and converted to devices, and therefore, SAMs have been widely applied in biochemical analysis.^{21,22} Our group has reported that solid-liquid interfacial detection based on fluorescent self-assembled multilayers can provide high sensitivity.^{23–25} SAMs can be employed to easily control the distance between graphene and fluorophores with a polyelectrolyte.

Recently, carbon dots (CDs) have attracted considerable attention due to easy preparation, high photochemical stability and good biocompatibility, compared with traditional semiconductor quantum dots and organic dyes.^{26,27} Moreover, CDs have also been applied in the sensing of ions (such as Hg²⁺, Cu²⁺, and ClO⁻),^{28,29} temperature,³⁰ biological molecules (for

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instance, dopamine, bilirubin, and L-/D-arginine)^{31–33} and proteins.³⁴

In this study, we synthesized poly(diallyldimethylammonium) chloride functionalized graphene (PDDA-G) with different reduction degrees to study its fluorescence enhancement efficiency. CDs were selected as fluorescent probes. PDDA and sodium polystyrenesulfonate (PSS) polyelectrolyte assembling layers were used to control the distance between CDs and PDDA-G surface (Scheme 1). Experimental results show that graphene can enhance fluorescence intensity of CDs-based SAMs. Efficiency of fluorescence enhancement of CDs was found to strongly depend on the distance between CDs and PDDA-G surface, graphene concentration, and degree of graphene reduction. This SAMs sensor could be further applied to detect trace Tc with limit of detection (LOD) of 0.9284 nM.

Experimental

Chemicals and reagents

Poly(diallyldimethylammonium)chloride (PDDA, MW 100 000–200 000) and tetracycline hydrochloride (Tc) were purchased from Aladdin (Shanghai, China). Sodium polystyrenesulfonate (PSS, MW 70 000) was obtained from Acros (USA). Procaine hydrochloride, clindamycin hydrochloride, cefotaxime sodium, 1,3,5-triazine, and trimethoprim were supplied by Macklin (Shanghai, China). Graphene oxide (GO) was purchased from Nanjing Xianfeng Nano Co. Ltd (Nanjing, China). Other reagents were of analytical grade and used without further purification. Ultrapure water was prepared by a Milli-Q water purification system (USA). Carbon dots (CDs) were synthesized according to the reported literature.³⁵

Apparatus

All fluorescence spectra were recorded on a HITACHI F-7000 fluorescence spectrometer (Hitachi, Japan). Absorption spectra were obtained from a UV-2600 UV-Vis spectrophotometer (Shimadzu, Japan) and a Cary 5000 UV-Vis-NIR spectrophotometer (Agilent, USA). Fluorescence images were obtained from a Leica TCS SP8 laser scanning confocal microscope (Leica, Germany). Fluorescence quantum yield measurements were conducted on a FLS-920 steady-state/transient fluorescence spectrometer (Edinburgh Instruments, UK). Fluorescence lifetime decay curves were obtained on a FS5 fluorescence spectrometer (Edinburgh Instruments, UK). Raman spectra were measured with an InVia confocal Raman spectrometer (Renishaw, UK). Thickness of the

graphene film was recorded by atomic force microscopy (AFM, Mutimode-8J, America). The potentials of PDDA-G and CDs were recorded by a Zetasizer Nano ZS (Malvern Instruments Ltd., UK). Assembly process was monitored in real time using a quartz crystal microbalance (QCM, Q-Sense Biolin Scientific, Sweden). Morphology of SAMs was probed by using a scanning electron microscope (SEM, FEI Inspect F50, America). Morphology of CDs was recorded by using a transmission electron microscope (TEM, H-7650, Japan). PDDA-LG was synthesized using a Xe lamp (CHF-XM-500 W, China) as the light source.

Construction of quartz/PDDA-G/(PSS/PDDA)_n/CDs SAMs

Quartz substrates were hydroxylated as previously reported.²³ Then, PDDA-G with different concentrations was poured onto the hydroxylated substrates and placed into an oven for 1 h. Then, the substrates with PDDA-G were dipped into 1 g L⁻¹ PSS and 1% (v/v) PDDA (0.1 M NaCl), sequentially, for 1 h. This process was repeated *n* times. Finally, these substrates were immersed into a CDs solution for 3 h and quartz/PDDA-G/(PSS/PDDA)_n/CDs SAMs were obtained.

Results and discussion

Characterization of PDDA-G

The obtained PDDA-G was characterized by UV-Vis, Raman and FTIR. UV-Vis absorption spectra of GO and PDDA-G were recorded, as shown in Fig. 1a. Compared with GO, the π - π^* transition of C=C of PDDA-G shifts from 230 nm to 270 nm and the shoulder peak at ca. 300 nm disappears. Raman spectral results are presented in Fig. 1b. PDDA-G and GO show similar Raman spectra. The peak (D peak) at ca. 1350 cm⁻¹ is the structural disorder peak and the graphite lattice defect peak, which is caused by vibration of sp³ hybrid carbon atoms. The G peak at ca. 1600 cm⁻¹ is the typical Raman peak of bulk crystalline graphite, which is usually related to sp² hybrid carbon atoms. In general, the degree of disorder of carbon material can be expressed as the ratio of the two integrated peak areas (*I*_D/*I*_G). By fitting calculations, it was found that *I*_D/*I*_G ratio of PDDA-G



Scheme 1 Schematic representation of assembly of (1) quartz/PDDA-G/(PSS/PDDA)₃/CDs and (2) quartz/PDDA/(PSS/PDDA)₃/CDs SAMs.



Fig. 1 (a) UV-Vis absorption spectra, (b) Raman spectra and (c) FTIR spectra of PDDA-G and GO.



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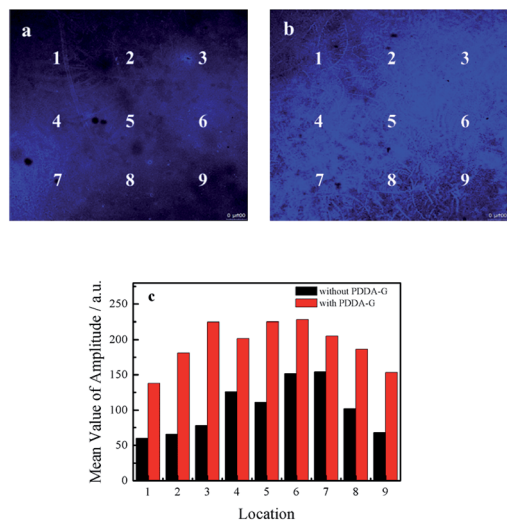


Fig. 5 Fluorescence microscopy images of (a) quartz/PDDA/(PSS/PDDA)₃/CDs SAMs and (b) quartz/PDDA-G/(PSS/PDDA)₃/CDs SAMs. (c) Confocal fluorescence values of quartz/PDDA/(PSS/PDDA)₃/CDs SAMs and quartz/PDDA-G/(PSS/PDDA)₃/CDs SAMs with different matching locations.

quartz/PDDA-G/(PSS/PDDA)₃/CDs SAMs was stronger than that of quartz/PDDA/(PSS/PDDA)₃/CDs SAMs due to the presence of graphene in the former, as shown in Fig. 5c.

To further investigate the enhancing effect of graphene, UV-Vis absorption spectra of quartz/PDDA-G/(PSS/PDDA)₃/CDs SAMs and quartz/PDDA/(PSS/PDDA)₃/CDs SAMs were recorded (Fig. S6†). Absorbance at 350 nm of CDs in these two SAMs was found to be unchanged, indicating that absorbed amounts of CDs on these two SAMs were consistent. Therefore, fluorescence intensity enhancement of CDs SAMs did not occur due to the increase in the amount of CDs in SAMs.

Optimization of PDDA-G concentration

To explore the effect of graphene concentration on fluorescence enhancement, PDDA-G assembled on quartz surface with concentrations of graphene varying from 0.001 mg mL⁻¹ to 0.1 mg mL⁻¹ was systematically studied. Results show that fluorescence intensity of CDs is PDDA-G concentration-dependent. Fluorescence intensity of CDs-based SAMs first increased and then decreased with the increase in PDDA-G concentration (Fig. S7†). When PDDA-G concentration was 0.005 mg mL⁻¹, fluorescence intensity of CDs was enhanced and thickness of the graphene layer at this PDDA-G concentration was *ca.* 4 nm (Fig. S8†). Since the transmittance decreased with the increase in PDDA-G concentration, fluorescence intensity of CDs SAMs was quenched.³⁶

Optimization of layers of polyelectrolyte

Fluorescence enhancement by graphene is dependent on the distance between CDs and PDDA-G surface. In this study, (PDDA/PSS)_n bilayers were employed to control the spacing distance between PDDA-G and CDs. The thickness of one bilayer of PSS/PDDA is *ca.* 3 nm,³⁷ and the fluorescence spectra of CDs with different numbers of spacer layers on quartz/PDDA-

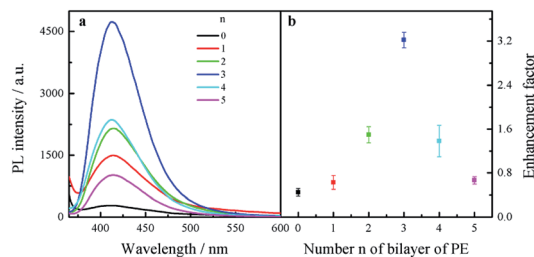


Fig. 6 (a) Fluorescence spectra of quartz/PDDA-G/(PSS/PDDA)_n/CDs SAMs with different layers of (PSS/PDDA)_n (*n* = 0–5). (b) Enhancement factor as a function of the number *n* of PSS/PDDA bilayers.

G/(PSS/PDDA)_n/CDs SAMs were examined (Fig. 6a). To further understand the enhanced efficiency of graphene, the enhancement factor (EF) was defined as the following equation:

$$EF = \frac{F_{Gr}}{F_{NGr}}$$

where F_{Gr} and F_{NGr} represent fluorescence intensity of SAMs with and without graphene, respectively.

Enhancement factor was plotted against the number of bilayers, as shown in Fig. 6b. The optimal layer number of polyelectrolytes was found to be 3 layers and thickness was *ca.* 9 nm; corresponding EF was calculated to be *ca.* 3.2.

Effect of degree of graphene reduction on fluorescence enhancement

To investigate the effect of the change in graphene structure on fluorescence enhancement, graphene with different degrees of reduction was synthesized under various reduction conditions, namely, room temperature, photochemical reduction and hydrothermal reaction. On conducting the synthesis reactions, the strongest peak in the UV-Vis absorption spectra of graphene was found to gradually shift from 230 nm to 270 nm,³⁸ as shown in Fig. 7a, indicating that the degree of graphene reduction gradually increased. Moreover, Raman spectra of four different graphene samples were recorded. The increase in I_D/I_G ratio indicated that the degree of graphitization gradually increased³⁹ (Fig. 7b). These experimental results showed that the degree of graphene reduction of PDDA-G was the highest. Degree of graphene reduction was found to have a significant effect on fluorescence enhancement efficiency (Fig. 8). As the degree of graphene reduction increased, the fluorescence enhancement efficiency continuously increased. However,

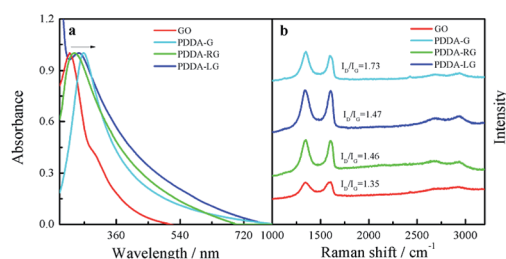


Fig. 7 (a) UV-Vis absorption spectra and (b) Raman spectra of GO, PDDA-G, PDDA-RG and PDDA-LG.



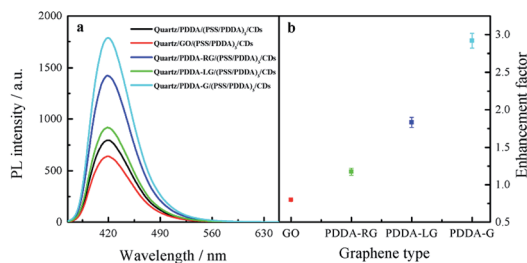


Fig. 8 (a) Fluorescence spectra of CDs SAMs and (b) enhancement factor of graphene with different degrees of graphene reduction.

fluorescence spectra of quartz/GO/(PSS/PDDA)₃/CDs SAMs show much weaker intensity than those of other platforms, probably due to the enhanced energy transfer from CDs to GO.²¹

Sensing for Tc based on inner filter effect

Since graphene can enhance fluorescence of SAMs, detection sensitivity should be improved. Therefore, CDs-based SAMs with graphene were expected to develop a sensing device with high sensitivity. Our experiments revealed that fluorescence intensity of CDs could be effectively quenched by introduced Tc. In general, quenching mechanisms involve energy transfer, electron transfer, inner filter effects, *etc.* In this study, it was found that fluorescence lifetime of CDs did not change in the presence of Tc (Fig. S9b†). To further probe the mechanism of fluorescence quenching by Tc, the reduction potential of CDs was detected as -0.475 V (Fig. S10a†). Then, the lowest unoccupied molecular orbital (LUMO) and highest occupied molecular orbital (HOMO) energy levels of CDs were calculated as -3.93 eV and -7.47 eV, respectively. The E_{LUMO} and E_{HOMO} of Tc have been obtained as -3.87 eV and -6.20 eV, respectively.⁴⁰ Therefore, it was considered that electrons in the HUMO of CDs could not be transferred to the HUMO of Tc, as shown in Fig. S10b.† Furthermore, emission spectra of CDs effectively overlap with absorption spectra of Tc (Fig. S9a†). Therefore, it can be deduced that CDs' fluorescence was quenched by Tc probably due to the inner filter effect (IFE) rather than energy transfer or electron transfer.

After optimization of experimental conditions, Tc at different concentrations was added into SAMs system. It was found that fluorescence intensity of SAMs in 10 mM PB buffer solution of pH 6.8 gradually decreased (Fig. 9a). The relationship between F_0/F of quartz/PDDA-G/(PSS/PDDA)₃/CDs SAMs and Tc concentration was established (Fig. 9b). It was found that when Tc concentration was varied in the range of 4.762–90.91 nM, a good linear relationship between the ratio F_0/F and Tc concentration could be obtained as $F_0/F = 6.432 \times 10^6 C + 1.003$, with $R^2 = 0.9924$, where C is the Tc concentration with M as the unit and F and F_0 represent the fluorescence intensity of SAMs in the presence or absence of Tc, respectively. LOD of CDs SAMs with graphene was measured and calculated to be 0.9284 nM, which is lower than 96 nM of CDs SAMs without graphene and lower by 1–3 orders of magnitude than those of other reported methods (Table S1†).

The effect of antibacterial drugs and other related substances on quartz/PDDA-G/(PSS/PDDA)₃/CDs SAMs were further studied. When Tc concentration was set to 56.60 nM,



plasmatic materials for surface-enhanced fluorescence, is expected to be a substitute for traditional precious metal materials.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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