As common bisphosphonates drugs, pamidronate disodium and zoledronic acid have been widely investigated for bone metastases. In this paper, a new “turn-off” model based on carbon dots (CDs) from black tea was established to analyze the two kinds of bisphosphonate drugs, pamidronate disodium and zoledronic acid. Through the new sensor, both of drugs can be quantitative, respectively, with the limit of detection of $5 \times 10^{-9}$ mol·L$^{-1}$ and $6 \times 10^{-9}$ mol·L$^{-1}$. In addition, the fluorescence of newly prepared CDs can be quenched by two drugs with various degrees via photoinduced electron transfer, which can be perfectly used to distinguish them. Most importantly, this turn-off method has been employed to analyze the two drugs under the influence of foreign interference factors. This method provides a new view and guidance for the rapid analysis and recognition of drugs for bone metastases in vitro and in vivo.

1. Introduction

Due to their excellent photo stability, high luminous intensity, wide range of excitation spectrum, and narrow emission spectrum, carbon dots (CDs) have been applied as fluorescent sensor in many fields, such as biochemistry, food science, and medical analysis [1–5]. Recently, a kind of food-based carbon dots have attracted more and more attention because of its good biocompatibility, low toxicity, and easy preparation [6–8]. In the past, many reports of fluorescence detecting mode “turn-off” based on CDs have already been reported, such as the turn-off fluorescent sensor based on nitrogen-doped carbon dots applied to detect Hg$^{2+}$ ions by Zhang and Chen [9] and the turn-off fluorescence sensor for the detection of ferric ion in water using green synthesized N-doped carbon dots [10].

As a significant kind of drugs, bisphosphonates play an important role in the treatment of metabolic bone diseases such as bone metasis and have been expanded widely to inhibit bone resorption and management of skeletal disorder including osteoporosis and malignant hypercalcemia [11, 12]. Meanwhile, as two representative drugs among the bisphosphonates, pamidronate disodium and zoledronic acid have been studied by more and more researchers [13, 14]. However, despite their frequent use, the rapid analytical method of them whether in vitro or in vivo has largely remained very few, and the traditional chromatographic methods were mostly time consuming and expensive and, what’s more, have complicated pretreatments [15–17]. On the other hand, known as a fast, sensitive, and stable analytical technology, the fluorescent sensor based on quantum dots, especially carbon dots, have a good application prospect in the drug analysis [18, 19].

In this paper, a new turn-off sensor based on carbon dots from black tea was established to quantitatively characterize the two kinds of bisphosphonate drugs, pamidronate disodium and zoledronic acid. The morphology and chemical composition, stability, and fluorescence properties of the...
newly prepared CDs were analyzed in details. Through using the different binding ability between CDs and different drugs, this turn-off sensor can be successfully used to quantitatively analyze pamidronate disodium and zoledronic acid no matter under the influence of foreign interference factors or in the human plasma. Most importantly, the fluorescent sensor has been successfully used for the distinction of the two bisphosphonate drugs. This method illustrates its promising opportunities in the studies of other drugs in vivo or in vitro using the fluorescent sensor based on CDs.

2. Experimental

2.1. Experimental Materials and Reagents. The black tea was bought from Minghuang Natural Food Development Co., Ltd. Tris (hydroxymethyl) amino methane, HCl, KCl, NaCl, MgCl₂, ZnCl₂, CuCl₂, CaCl₂, and FeCl₃ were purchased from Sinopharm Chemical Reagent Co., Ltd. CDs from the black tea via the hydrothermal route were synthesized in our laboratory. Deionized distilled water prepared from a Mole water purification system was used.

2.2. Apparatus and Procedures. UV-Vis absorption spectra were examined by using a U-3900 UV-Vis spectrometer (Hitachi). The fluorescence spectra were acquired by an FL-7000 luminescence spectrometer (Hitachi). High-resolution transmission electron microscopy (HRTEM) was examined by using a Tecnai G2 F30 S-Twin microscope (Philips-FEI). Samples containing different concentrations of drugs and CDs were made up to 1 mL in 50 mmol·L⁻¹ Tris-HCl (pH = 6.0). The concentration of CDs was set at 8.0 × 10⁻⁵ g·mL⁻¹. The emission spectrum of the solution was then measured 5 min later. All optical measurements were performed at room temperature under ambient conditions, and the excitation wavelength (λₑₓ) was 360 nm.

2.3. Preparation of CDs from the Black Tea. The CDs were synthesized by using a facile, green, and low-cost
hydrothermal treatment based on the black tea as the carbon source as follows: 0.5 g the black tea and 40 mL water were added into 50 mL reaction kettle and then heated at 180°C for 3 h. The obtained CDs were centrifuged for 15 min at a rotating speed of 5000 r·min⁻¹ and then dialyzed in a dialysis bag (molecular weight cut off = 3500) for 12 h; finally, the product were freeze dried for further analysis.

2.4. Quantum Yield Measurement. The quantum yields (QYs) of the CDs were measured by comparing the integrated fluorescence intensity (320 nm excitation) and the absorbance value (320 nm) of CD and quinine sulfate was used as a reference. QY was calculated as follows:

\[ Q_X = Q_{ST} \cdot \left( \frac{I_X}{I_{ST}} \right) \cdot \left( \frac{A_{ST}}{A_X} \right) \cdot \left( \frac{\eta_X}{\eta_{ST}} \right), \]

where \( I \) is the integrated fluorescence intensity, \( A \) represents the absorbance value, and \( \eta \) means the refractive index of the solvent (both 1.33). The subscripts “X” and “ST” were CDs and quinine sulfate, respectively. The \( Q_{ST} \) was 54% while quinidine sulfate was in 0.1 mol·L⁻¹ H₂SO₄. In order to minimize the absorption effect, the absorbance value was set as 0.1 [9].

3. Results and Discussion

3.1. Fluorescence Properties and Characterization of CDs. As shown in Figure 1(a), it can be seen that the absorbance band of CDs was centered at around 270 nm in the UV spectrum which was from the \( n-\pi^* \) transition. The CDs emitted strong blue light under UV irradiation of 365 nm (Figure 1(b)), and while the \( \lambda_{ex} \) increased from 280 to 570 nm, the emission spectra of the CDs displayed a typical excitation wavelength-dependent characteristic, which had been suggested to be a result of varied fluorescence characteristics of CDs with various sizes or emissive sites existed on the surfaces of the CDs [2]. Chemical and structural information about the CDs using the FTIR spectroscopy was shown in Figure 1(c); the peak around
indicated the existence of –OH; the peaks around 2940 and 1240 cm\(^{-1}\) were assigned to the C–H; the peaks at 1660 cm\(^{-1}\) and 1620 cm\(^{-1}\) were attributed to the stretching vibration of C=O and C=C, respectively; the peaks at 1123 and 1007 cm\(^{-1}\) indicated C–OH and C–O, respectively. Figure 1(d) shows the high-resolution TEM (HRTEM) image of the CDs which reveals that the CDs had a size distribution around 2.2 nm. According to (1), the QY of the CDs based on the black tea was calculated to be 10.23% with quinine sulfate as a reference.

In order to further study the structure of CDs, the XPS analysis was conducted. Figure 2(a) shows the XPS spectrum of the CDs, and the three predominant peaks of C1s, N1s, and O1s at 284.61, 399.49, and 531.91 eV were corresponded to the binding energies. As can be seen from the result that the as-prepared CDs mainly contained C, O, and N, the contents of the three elements were calculated to be 73.8% (C), 21.7% (O), and 4.5% (N). The C1s spectrum (Figure 2(b)) can be deconvoluted into four peaks at around 284.6, 285.6, 286.6, and 287.9 eV, which were attributed to C–C, C–N, C–O, and C=O groups, respectively. In addition, the N1s spectrum (Figure 2(c)) can be deconvoluted into three peaks at 399.3, 400.1, and 400.7 eV which can be assigned to N–(C)\(_3\), N–H, and C–N–H groups, respectively. From the O1s spectrum shown in Figure 2(d), the three fitted peaks at 530.3, 531.3, and 532.7 eV were ascribed to C–OH/C–O–C, adsorbed oxygen, and –C=O groups, respectively. The result from XPS showed that the surface of the CDs from the black tea was functionalized by multiple oxygenated and nitrous groups which may lead the interactions with different drug molecules.

![Figure 3: Fluorescence quenching behavior of CDs after the addition of various concentrations (5 × 10^{-9} mol·L^{-1} to 4 × 10^{-7} mol·L^{-1}) of pamidronate disodium (A-I (a)) and the addition of various concentrations (6 × 10^{-9} mol·L^{-1} to 8 × 10^{-7} mol·L^{-1}) of zoledronic acid (A-G (c)). The linear plot of relative fluorescence intensity (F_0/F_1) of CDs as a function of pamidronate disodium (5 × 10^{-9} mol·L^{-1} to 4 × 10^{-7} mol·L^{-1} (b) and zoledronic acid (6 × 10^{-9} mol·L^{-1} to 8 × 10^{-7} mol·L^{-1} (d)). Every data point was the mean of three measurements. The error bars were the standard deviation.](image)
3.2. Effective Quenching of Fluorescence of CDs by Pamidronate Disodium and Zoledronic Acid: Turn-Off Process. As shown in Figure 3(a), the increased addition of pamidronate disodium led to the gradual decreased fluorescence of CDs. And the relationship between the concentration of pamidronate disodium and the fluorescence intensities of CDs follows the “Stern-Volmer” equation:

\[ \frac{F_0}{F_1} = 1 + K_{sv}[M], \]  

where \( F_0 \) and \( F_1 \) were the fluorescence intensities of CDs in the absence and presence of the drugs, respectively; \([M]\) was the concentration of pamidronate disodium, and the quenching constant \( K_{sv} \) defines the quenching efficiency of pamidronate disodium. The equation shows that \( F_0/F_1 \) was directly proportional to the concentration of pamidronate disodium. As shown in the Stern-Volmer plot in the inset of Figure 1, the \( K_{sv} \) of pamidronate disodium to the fluorescence of CDs was \( 4.8 \times 10^6 \text{ L} \cdot \text{mol}^{-1} \), with linear range from \( 5 \times 10^{-9} \text{ mol} \cdot \text{L}^{-1} \) to \( 4 \times 10^{-7} \text{ mol} \cdot \text{L}^{-1} \) and a correlation coefficient \( R = 0.996 \). On the other hand, as shown in Figure 3(c), when the concentration of zoledronic acid was increasing from \( 6 \times 10^{-9} \text{ mol} \cdot \text{L}^{-1} \) to \( 8 \times 10^{-7} \text{ mol} \cdot \text{L}^{-1} \), the fluorescent intensity of the CDs at around 430 nm also decreases gradually, which indicating that zoledronic acid can effectively quench the fluorescence of CDs. Meanwhile, through the Stern-Volmer equation (Figure 3(d)), the \( K_{sv} \) of zoledronic acid to the fluorescence of CDs was \( 2.8 \times 10^6 \text{ L} \cdot \text{mol}^{-1} \) with the linear range from \( 6 \times 10^{-9} \text{ mol} \cdot \text{L}^{-1} \) to \( 8 \times 10^{-7} \text{ mol} \cdot \text{L}^{-1} \) and a correlation coefficient \( R = 0.991 \). The detect limit of pamidronate disodium and zoledronic acid was, respectively, \( 5 \times 10^{-9} \text{ mol} \cdot \text{L}^{-1} \) and \( 6 \times 10^{-9} \text{ mol} \cdot \text{L}^{-1} \). Obviously, pamidronate disodium exhibits better quenching ability to the fluorescence of CDs than zoledronic acid with the value of \( K_{sv} \) almost twice higher than that of zoledronic acid and seen in Figure 4, the addition of pamidronate disodium shows a significantly weaker fluorescence quenching phenomenon than the same concentration of zoledronic acid, which may be caused by the different intermolecular binding capacities. As shown in Scheme 1, compared with the imidazolyl group contained in zoledronic acid, the amino group contained in pamidronate disodium has the smaller space obstruction and a better affinity to the polycarboxyl group modified on the surface of the CDs, leading to much stronger quenching phenomenon to the fluorescence via photoinduced electron transfer and exhibiting excellent selectivity in detecting the two drugs with similar structures. To explain the sensing mechanism in depth, lifetimes of CDs before and after adding bisphosphonates were investigated and the...
result was shown in Figure 5. The lifetime of CDs after adding different bisphosphonates was reduced by different degrees which can act as a proof for the photoinduced electron transfer mechanism. And what’s more, the reduction fluorescence lifetime of CDs caused by pamidronate disodium was more than that caused by zoledronic acid. This also demonstrated the different quenching capabilities of bisphosphonates towards the carbon dots.

3.3. Selectivity of the Sensor for Pamidronate Disodium and Zoledronic Acid Detection. As to evaluate the selectivity of the CDs sensor, 10 different biologically relevant metal ions including Na⁺, K⁺, Mg²⁺, Ca²⁺, Fe³⁺, Zn²⁺, Cu²⁺, Bovine serum albumin (BSA), human serum albumin (HSA), and the mixture of all the above were selected as foreign interference with a concentration at 1×10⁻⁷ mol·L⁻¹. The comparison is shown in Figure 6 which can indicate that the interference effect of all the 10 interferences were slight. In addition, even in the spiked samples of the human plasma and urine, a recovery of 101.3% and 99.4% for pamidronate disodium (4×10⁻⁷ mol·L⁻¹) and a recovery of 102.1% and 100.8% for zoledronic acid (8×10⁻⁷ mol·L⁻¹) were obtained. What’s more, other kinds of bisphosphonates such as nерidronate sodium and alendronate sodium also failed to have a significant impact on the fluorescence of CDs. It was obvious that a special effect appeared between CDs and pamidronate disodium or zoledronic acid, so the as-prepared CDs had a high selectivity for pamidronate disodium and zoledronic acid analysis.

4. Conclusion

In this study, a new turn-off model based on newly prepared carbon dots (CDs) from the black tea was established to analyze pamidronate disodium and zoledronic acid. By using the new sensor, both of drugs can be quantitative, respectively, with high sensitivity and stability. Meanwhile, the fluorescence of the CDs can be quenched by two drugs with various degrees via photoinduced electron transfer, which provides the sensor a high selectivity to the two structurally similar drugs. What’s more, this turn-off method shows good stability and has been employed to analyze the two drugs under the influence of common foreign interference factors which indicating its good application prospects for drugs analysis in vitro and in vivo.

Conflicts of Interest

The authors declare that they have no conflicts of interest regarding the publication of this paper.

Acknowledgments

This work was financially supported by the Tianjin Natural Science Foundation (no. 12JCYBJC16400) and the Tianjin Municipal Health Bureau Fund Project (nos. 15KG124, 12KG121).

References

[1] S. Zhu, Q. Meng, L. Wang et al., “Highly photoluminescent carbon dots for multicolor patterning, sensors, and bioimaging,” Angewandte Chemie, vol. 125, no. 14, pp. 4045–4049, 2013.
[2] S. Zhu, Y. Song, X. Zhao, J. Shao, J. Zhang, and B. Yang, “The photoluminescence mechanism in carbon dots (graphene quantum dots, carbon nanodots, and polymer dots): current state and future perspective,” Nano Research, vol. 8, no. 2, pp. 355–381, 2015.
[3] Y. Dong, H. Pang, H. B. Yang et al., “Carbon-based dots co-doped with nitrogen and sulfur for high quantum yield and excitation-independent emission,” Angewandte Chemie International Edition, vol. 52, no. 30, pp. 7800–7804, 2013.
[4] U. Barua, S. Gogoi, A. Konwar et al., “Carbon dot based sensing of dopamine and ascorbic acid,” Journal of Nanoparticles, vol. 2014, Article ID 178518, 8 pages, 2014.
[5] K. Yang, Y. Wang, W. Wang, H. Miao, and X. Yang, “Dual-channel sensor of carbon dots cooperating with gold nano-clusters employed for assaying multiple targets,” Biosensors and Bioelectronics, vol. 91, no. 15, pp. 566–573, 2017.
[6] J. Shen, S. Shang, X. Chen, D. Wang, and Y. Cai, “Facile synthesis of fluorescence carbon dots from sweet potato for
Fe	extsuperscript{3+} sensing and cell imaging,” *Materials Science and Engineering: C*, vol. 76, pp. 856–864, 2017.

[7] L. Wang and H. S. Zhou, “Green synthesis of luminescent nitrogen-doped carbon dots from milk and its imaging application,” *Analytical Chemistry*, vol. 86, no. 18, pp. 8902–8905, 2014.

[8] S. Zhao, M. Lan, X. Zhu et al., “Green synthesis of bifunctional fluorescent carbon dots from garlic for cellular imaging and free radical scavenging,” *ACS Applied Materials & Interfaces*, vol. 7, no. 31, pp. 17054–17060, 2015.

[9] R. Zhang and W. Chen, “Nitrogen-doped carbon quantum dots: facile synthesis and application as a “turn-off” fluorescent sensor for detection of Hg	extsuperscript{2+} ions,” *Biosensors and Bioelectronics*, vol. 55, pp. 83–90, 2014.

[10] T. N. J. I. Edison, R. Atchudan, J. J. Shim, S. Kalimuthu, B.-C. Ahn, and Y. R. Lee, “Turn-off fluorescence sensor for the detection of ferric ion in water using green synthesized N-doped carbon dots and its bio-imaging,” *Journal of Photochemistry and Photobiology B: Biology*, vol. 158, pp. 235–242, 2016.

[11] Y. Sakamoto, “Bilateral typical femoral fractures in a patient with metastatic breast cancer on long-term bisphosphonate therapy: a case report,” *Journal of Osteoporosis and Physical Activity*, vol. 2, no. 1, pp. 1–4, 2014.

[12] V. Viereck, G. Emons, V. Lauck et al., “Bisphosphonates pamidronate and zoledronic acid stimulate osteoprotegerin production by primary human osteoblasts,” *Biochemical and Biophysical Research Communications*, vol. 291, no. 3, pp. 680–686, 2002.

[13] L. Yang and S. Du, “Efficacy and safety of zoledronic acid and pamidronate disodium in the treatment of malignant skeletal metastasis: a meta-analysis,” *Medicine*, vol. 94, no. 42, p. e1822, 2015.

[14] Y. Orita, I. Sugitani, S. Takao, K. Toda, J. Manabe, and S. Miyata, “Prospective evaluation of zoledronic acid in the treatment of bone metastases from differentiated thyroid carcinoma,” *Annals of Surgical Oncology*, vol. 22, no. 12, pp. 4008–4013, 2015.

[15] R. Karthikeyan, O. S. A. I. Koushik, and P.S. Babu, “Stability indicating method development and validation of zoledronic acid in injectables by RP–HPLC,” *Journal of International Research in Medical and Pharmaceutical Sciences*, vol. 7, no. 4, pp. 159–174, 2016.

[16] Ł. Matuszewski, A. Matuszewska, T. Mazurkiewicz, and S. Ohga, “Determination of bisphosphonates by ion-pair HPLC,” *Journal of the Faculty of Agriculture Kyushu University*, vol. 56, no. 2, pp. 213–216, 2011.

[17] E. Ban, J. Y. Park, H. T. Kim, and C.-K. Kim, “Determination of alendronate in low volumes of plasma by column-switching high-performance liquid chromatography method and its application to pharmacokinetic studies in human plasma,” *Archives of Pharmacal Research*, vol. 34, no. 12, pp. 2079–2086, 2011.

[18] T. Feng, X. Ai, G. An, P. Yang, and Y. Zhao, “Correction to charge-convertible carbon dots for imaging-guided drug delivery with enhanced in vivo cancer therapeutic efficiency,” *ACS Nano*, vol. 10, no. 5, p. 5587, 2016.

[19] Z. Yang, M. Xu, Y. Liu et al., “Nitrogen-doped, carbon-rich, highly photoluminescent carbon dots from ammonium citrate,” *Nanoscale*, vol. 6, no. 3, pp. 1890–1895, 2014.