Efficient Adsorptive Performance of Medical Stone Decorated by Carbon Dots

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ABSTRACT
Carbon dots could significantly change the property of a normal material and have received wide attention in the recent decade. In this research, glucose as a carbon source, carbon dots decorated medical stone (CD-MS) was successfully synthesized for efficient adsorptive removal of organic pollutants. Pyrolytic temperature and glucose concentration for the adsorbent preparation were proved to have a significant impact on the adsorptive performance. The optimal pyrolytic temperature and glucose concentration were found to be 300°C and 0.5 M, yielding the optimized adsorbent 0.5CD-MS-300 superior to other carbon dots decorated MS. Surface morphology analysis demonstrated that the carbon dots were successfully immobilized on the surface of MS while the atomic ratio of C increased from 2.6% of the raw MS to 11.25% of the 0.5CD-MS-300. Three organic pollutants including p-nitrophenol, orange II and methylene blue with different charge properties were employed to explore the adsorptive performance of the 0.5CD-MS-300. The results indicated that the surface of 0.5CD-MS-300 was negatively charged while carbon dots had significantly improved the adsorption capability of the raw MS. As such, the resulting adsorbent 0.5CD-MS-300 can be considered as a powerful adsorbent for the removal of some organic contaminants from wastewater.

INTRODUCTION
With the development of industry and agriculture, a large number of organic contaminants, such as dyes, endocrine-disruptors, and pharmaceutical and personal care products, have come into the natural environment along with the discharge of effluents (Westerhoff et al. 2005, Crini 2006). These organic pollutants existing in wastewater can cause a severe hazard to human and animal health due to the delivery of the food chain. To date, there are many treatment technologies, including coagulation, sonocatalytic degradation, photocatalytic degradation and adsorption, to be applied to the removal of the organic pollutants (Polubesova et al. 2006, Li et al. 2014, Mahata et al. 2007, Zhang et al. 2006). Among these technologies, the adsorption process is considered as a promising and reliable method due to its ease of operation, low energy consumption, low cost and high efficiency (Qu 2008).

Medical stone (MS) is one kind of natural silicate mineral without toxicity. Its main mineral components include Na(AlSi₃O₈), Na(AlO₂)(SiO₂)₃, K(AlSi₃O₈), K(AlO₂)(SiO₂)₃, Ca(Al₂Si₂O₈) and Mg(Al₂Si₂O₈) (Gao et al. 2011). MS has a tetrahedron structure of [SiO₄]³⁻ bonding to metal ions (such as K, Na, Ca, Mg, Cu and Al, etc.), contributing to the formation of a large internal surface area (Gao et al. 2012). Therefore, a series of excellent performances for MS are expected, such as biological activity, the ability of dissolving and adsorbing and adjusting pH capacity. Meanwhile, it is widely used for food preservation, water purification and medical care. In recent years, MS had been studied as an adsorbent material, which achieved remarkable advances on the adsorption of heavy metals in wastewater (Gao et al. 2011, Zhou et al. 2015). Even though, to our best knowledge, there have been few reports about the adsorption removal of organic contaminants.

Carbon dots are novel carbon-based nanomaterial. They contain abundant oxygen-containing functional groups (-OH, -COOH and -C=O) on the surfaces, with good solubilities and optical properties. As such, carbon dots are widely implemented for bioimaging, chemical sensors, and photovoltaic devices, etc. (Ding et al. 2014, Shen et al. 2012). In recent years, CDs have been considered an ideal candidate material for organic or inorganic pollutants removal due to their rich functional groups, ease of preparation and low toxicity (Hsu & Chang 2012). However, some significant disadvantages of CDs based adsorbent focused on the dissolution of CDs as well as difficulties of separation and regeneration (Liu et al. 2017). Consequently, it is a necessity of combining CDs with other constituents to synthesize CDs-functionalized adsorbents.
In this research, we prepared the carbon-dots decorated medical stone (CD-MS) for the efficient adsorption of organic pollutants. Glucose, with numerous hydroxy groups and formyl groups, was chosen as the precursor of CDs and can provide more active adsorption sites on the surfaces of the MS (Hsu & Chang 2012). The effects of pyrolytic temperature and glucose concentration for the adsorbent preparation were emphatically studied. At the same time, p-nitrophenol (PNP), orange II (ORII) and methylene blue (MB) with different charge properties in a neutral aqueous solution were selected as target organic pollutants to investigate the adsorptive performance and adsorption capacity of the CD-MS.

**MATERIALS AND METHODS**

**Chemicals:** Orange II (ORII) (mass fraction > 95%) was purchased from Beijing Chemical Reagents Company. Methylene blue (MB) (mass fraction > 98.5%, chemical pure) was purchased from Tianjin Chemical Reagent Research Institute. P-nitrophenol (PNP) used was of analytical grade and provided by Tianjin Guangfu Fine Chemical Research Institute, China. Other chemicals were of analytical grade and used without further purification. Deionized (DI) water was used throughout the research. The molecular structures of ORII, MB and PNP are shown in Fig. 1.

**Adsorbent preparation:** The natural medical stone (MS) was obtained from the Hunan Province of China. It was shattered with a multi-function pulveriser and subsequently sieved to a particle size between those of 80-mesh screens and 100-mesh screens. Firstly, 10 g of the powdered MS was immersed to 200 mL of 0.5 M or 0.05 M glucose solution and the mixture was shaken at a speed of 120 rpm for 12 h. Then, the resulting samples were rinsed with DI water for five times and oven-dried overnight at 80°C. The prepared carbon dots decorated medical stone (CD-MS) are denoted as 0.5CD-MS and 0.05CD-MS in the following studies.

Desired amounts of MS, 0.5CD-MS and 0.05CD-MS were tightly placed in a ceramic crucible with a lid and pyrolyzed in a muffle furnace under oxygen-limited conditions. The pyrolysis temperature was raised to the desired
values of 200°C, 300°C and 400°C at a heating rate of 10°C min⁻¹ and held constant for 2 h. After pyrolyzation, the samples were allowed to cool down to room temperature in the furnace. These pyrolyzed 0.05CD-MS were denoted as 0.05CD-MS-200, 0.05CD-MS-300 and 0.05CD-MS-400, respectively, while the pyrolyzed 0.5CD-MS are denoted as 0.5CD-MS-200, 0.5CD-MS-300 and 0.5CD-MS-400, respectively. Similarly, the adsorbent MS-200, MS-300 and MS-400 were also prepared.

**Batch adsorption studies:** Batch experiments were conducted in a series of 250-mL conical flasks. In the flasks, 20 mg of the adsorbent was added to 100 mL solution with an initial PNP, ORII or MB concentration of 20 mg/L. Then, the flasks were sealed and shaken at 298 K at a speed of 120 rpm for 24 h to achieve equilibrium. Solution pH was maintained at neutral unless otherwise stated. To examine the effect of glucose concentration, 5 mg adsorbent (0.5CD-MS or 0.05CD-MS) was used to adsorb PNP with an initial concentration of 5 mg/L.

**Characterization:** The morphology and superficial element compositions of MS and 0.5CD-MS-300 were characterized by a Hitachi S-4800 field scanning electron microscope (FESEM) coupled with an energy dispersive spectrometer (EDS).

**Analysis methods:** All the samples were collected and filtered by microporous membranes (0.45μm) before analysis. The concentration of PNP, ORII and MB was measured using UVmini-1240 spectrophotometer (Shimadzu, Japan) at the wavelength of maximum adsorption of 317 nm, 483 nm, and 664 nm, respectively (Cheng et al. 2019, Mi et al. 2016). The adsorption capacity was calculated using the following equation:

$$q_e = \frac{(C_0 - C_e)V}{M} \quad \ldots(1)$$

Where, $q_e$ (mg/g) is the adsorption capacity at equilibrium; $C_0$ and $C_e$ (mg/L) are the initial and equilibrium concentrations of PNP, ORII, and MB, respectively; $V$ (L) is the solution volume, and $M$ (g) is the weight of adsorbents.

**RESULTS AND DISCUSSION**

**Effect of Pyrolytic Temperature and Glucose Concentration on PNP Adsorption**

Both the pyrolytic temperature and glucose concentration were important for adsorbent preparation. It was observed from Fig. 2 that the CD-MS pyrolyzed at 300°C had a comparatively higher adsorption capacity for PNP removal. The concentration of glucose for the adsorbent preparation was expected to have a significant impact on the uptake of contaminants as well. As illustrated, it was noted that the adsorption capacity of PNP by 0.5CD-MS was higher than that by 0.05CD-MS under the same pyrolytic temperature. Further, at the pyrolytic temperature of 300°C, the adsorption capacities by the 0.5CD-MS and 0.05CD-MS were 4.16 mg/g and 1.88 mg/g, respectively. This might be attributed to the increasing bonding sites on the CD-MS surfaces resulted from the increase of glucose concentration. Similar experimental results were also observed by Xue et al. (2013). As such, 0.5CD-MS-300 was used for the following experiments.

On the other hand, as the structure of pore canals and surface area of internal pores of CD-MS varied as a conse-

![Fig. 2: Effect of glucose concentration and pyrolytic temperature on adsorption of PNP. PNP 5.0 mg/L, neutral solution pH.](image-url)
quence of the different pyrolytic temperatures, the adsorption capacity of PNP by the resulting adsorbents was expected to vary accordingly. The relationship between pyrolytic temperature and adsorption capacity of PNP is presented in Fig. 2. It shows that the adsorption capacity of PNP by 0.5CD-MS-300 reached the maximum of 4.16 mg/g, which might be attributed to the removal of crystal water in the pore canals and the increase of the surface area of internal pores as the pyrolytic temperatures rose. However, continuous increase of pyrolytic temperature was not helpful for the uptake of PNP and the adsorption capacity declined, which might be attributed to the collapse of pore canals.

**Characterization of the Carbon Dots Decorated Medical Stone**

As stated above, the 0.5CD-MS-300 demonstrated the highest adsorption capacity for PNP in comparison to other adsorbents. The surface morphology of the 0.5CD-MS-300 and the raw MS was characterized by SEM and presented herein. From Fig. 3a and c, it was observed that the raw MS presented a natural crystal structure, although it was difficult to find some pore canals from the MS surfaces. Differently, the surface structure of the 0.5CD-MS-300 became loose after pyrolysis, and some sponge pores were scattered onto the surfaces of the adsorbent, which might facilitate the adsorption of contaminants (Yu et al. 2013). This demonstrated pyrolysis treatment can change the surface structure of MS. From Fig. 3b and d, many fine particles aggregated on the surfaces of MS, which were deduced to be carbon dots immobilized on the surfaces of MS.

To further explore whether carbon dots had decorated on the surfaces of MS, EDS analysis was conducted. As illustrated in Fig. 4a and b, comparing to the atomic ratio of C increased from 2.6% of the raw MS to 11.25% of the 0.5CD-MS-300, which was reasonably attributed to the introduction of CDs on the surfaces of MS. Accordingly, the atomic ratio of O decreased from 57.97% of the raw MS to 53.70% of the 0.5CD-MS-300. Pyrolysis under different temperatures and immobilization of carbon dots had significantly altered the surface properties of the raw MS.

**Effect of Adsorbent Dosage on PNP Adsorption**

The effect of adsorbent (0.5CD-MS-300) dosage from 100 mg/L to 800 mg/L was investigated and presented in Fig. 5. The plot showed that uptake of PNP decreased gradually with an increase of adsorbent dosage. One reason was that the unsaturated adsorption sites increased with increasing adsorbent dosage at a fixed concentration and volume of adsorbent dosage.
PNP. The other was attributed to the decrease of surface area and the increase of diffusion path length of adsorbent due to the particle aggregation (Shirmardi et al. 2016, Kumar et al. 2010).

**Adsorptive Performance of Carbon Dots Decorated Medical Stone**

The surface charge property of an adsorbent always plays an important role during the adsorptive removal of organic contaminants. Three organic pollutants with different charge properties including PNP, ORII and MB were selected to explore the adsorptive performance of the 0.5CD-MS-300. The initial concentration of the three contaminants was fixed at 20 mg/L. As illustrated in Fig. 6, it was noted that the adsorption capacity of PNP and MB by 0.5CD-MS-300 was much higher than that of ORII. The adsorption capacity of PNP, MB, and ORII was 27.87 mg/g, 15.37 mg/g, and 0 mg/g, respectively. This indicated that the 0.5CD-MS-MS had a high adsorption selectivity for organic pollutants with different charge properties.

To our knowledge, PNP ($pK_a = 7.15$) in water solution exists in two species of molecule and anion, and molecular

![Fig. 4: EDS analysis results of the raw MS(a) and the 0.5CD-MS-300 (b).](image)
Fig. 5: Effect of adsorbent dosage on PNP adsorption. PNP 5.0 mg/L, neutral solution pH.

Fig. 6: Adsorption of PNP (a), ORII (b) and MB (c) onto the raw MS and the 0.5CD-MS-300. Contaminant concentration 20.0 mg/L, neutral solution pH.
PNP can convert to anions at pH > 7.15 due to dissociation (Sarkar et al. 2010). Meanwhile, MB is moderately alkaline in water and produces cation (C⁺ or CH⁺) (Gobi et al. 2011). By contrast, ORII (pKₐ= 10.6 and pKₐ= 1) has three different forms depending on the pH of the aqueous solution, noted H₂L which is doubly protonated and dominant at pH < 1, HL⁻ which is mono protonated and dominant at pH 1-10.6, L²⁻ which is non-protonated and dominant at pH > 10.6 (Abramian & El-Rassy 2009). Therefore, the three pollutants have different electrical characteristics in a neutral aqueous solution and the predominant species of PNP, MB and ORII are molecule, cation (C⁺ or CH⁺), and anion (HL⁻), respectively. Accordingly, from the results mentioned above, it could be deduced that the surface of 0.5CD-MS-300 was negatively charged, which was beneficial to MB adsorption due to an electrostatic attraction. Meanwhile, the results indicated that electrostatic attraction/repulsion was an important adsorption mechanism for 0.5CD-MS-300, which also facilitated the high adsorption selectivity for these organic pollutants. However, an interesting phenomenon was that the adsorption capacity of PNP is excellent. As mentioned, most of PNP exists as molecules in a neutral solution, which indicated that the effect of electrostatic force was not dominant. Therefore, considering the abundant O-containing functional groups of carbon dots on the 0.5CD-MS-300, the higher adsorption capacity of PNP might be attributed to H-bond interaction between PNP molecules and the surface functional groups of the adsorbent, π-π interaction, and π-stacking forces (Zheng et al. 2017).

In addition, the 0.5CD-MS-300 demonstrated a higher adsorption capacity towards PNP and ORII than the raw MS in general, indicating that the introduction of carbon dots was beneficial to the adsorption capability for PNP and ORII. By contrast, the introduction of carbon dots declined the adsorption of MB. Therefore, the surface of the 0.5CD-MS-300 was predominantly negatively charged.

CONCLUSION
Carbon dots decorated medical stone (CD-MS) was creatively prepared by immobilizing carbon dots on the surfaces of medical stone (MS). Compared to the raw MS, the 0.5CD-MS-300 demonstrated an excellent adsorption capability for PNP. Each experiment for the adsorption of the three pollutants including PNP, ORII and MB with different charge properties, indicated that the surface of 0.5CD-MS-300 was negatively charged. Furthermore, 0.5CD-MS-300 had a high adsorption selectivity for different organic pollutants, which was attributed to the electrostatic attraction/repulsion as one of the fundamental adsorption mechanisms. In a word, carbon dots had successfully decorated the raw medical stone and the resulting adsorbent 0.5CD-MS-300 can be considered as a powerful adsorbent for the removal of organic pollutants from wastewater.

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