Introduction

With the discovery of antibiotics, such as penicillin, for human beings to fight infectious diseases, there is a common phenomenon of abusing antibiotics to cope with various illnesses. However, antibiotics are difficult to metabolize in a short period of time in animals, plants and humans. Simultaneously, emissions from pharmaceutical companies have also caused enormous environmental pollution. Thus, a large amount of used antibiotics went to the soils, rivers, animals, etc. The accumulation of these residual antibiotics not only pollutes the environment, but also causes harm to human health. With the emergence of super bacteria, there is serious concern that society will lack the required antibiotics in future. The warnings of scientists have attracted the attention of the public and governments. Therefore, searching for effective and non-toxic methods to remove antibiotic pollutants has become a hot research topic.

There are a number of methods which aim to decrease the amount of antibiotics in the environment, including adsorption, photocatalysis, electrochemical oxidation, advanced oxidation and biodegradation. Among these methods, adsorption is a simple and cheap way to solve the problem of antibiotic pollution and can be divided into two categories: chemisorption and physisorption. Chemisorption requires high activation energy and usually higher temperatures, while physisorption is usually caused by molecular force and can occur at lower temperature. Furthermore, the chemisorption reaction may be undetectable at low temperature or pressure. As characteristic of physisorption, for gases at temperatures above their critical temperature, adsorption is confined to a monolayer (unimolecular). Therefore, the adsorption capacity is related to the specific surface area and structures of the adsorbents. The effects of the specific surface area and structure of different types of single adsorbent materials and composite adsorbent materials on the adsorption capacity have been studied in depth. Promising materials, such as zeolites for adsorption of tetracycline, attapulgite/carbon composites for adsorption of tetracycline, and oil shale powders and functional MOFs for adsorption of norfloxacin, were reported. A large number of experimental results show that the specific surface area and the structure of composites are the key factors that make adsorption more effective than with a single material. Therefore, there is an increasing focus on developing composite adsorbent materials with an optimized structure and high specific surface area via simple preparation processes amenable to mass production.

In this work, ultralong TiO$_2$ nanotubes were prepared using a simple hydrothermal treatment. In order to sustain the ultralong structure and increase the specific surface area, the ultralong TiO$_2$ nanotubes were covered with an ultrathin carbon layer by carbonizing ethanol, and thus a core–shell TiO$_2$@C ultralong nanotube composite was formed. Three common antibiotics, i.e., tetracycline (TC), ofloxacin (OFO) and norfloxacin (NFO), were adsorbed by the TiO$_2$@C composite, and the adsorption capacities were measured to be 240 mg g$^{-1}$ (TC), 232 mg g$^{-1}$ (OFO), and 190 mg g$^{-1}$ (NFO), respectively. The adsorption mechanism is consistent with a Langmuir pseudo-first-order kinetic model.
study sheds light on the mechanism of metal oxide/carbon hybrids for adsorbing antibiotics and identifies routes to improve performance.

**Experimental section**

**Preparation of ultralong titanium dioxide nanotubes**

Firstly, 0.5 g P25 (a commercial titanium dioxide which is an anatase and rutile two-phase mixture nanopowder with an average particle diameter of 25 nm) was added into 40 mL of 10 mol L⁻¹ NaOH solution and dispersed in a 50 mL reaction kettle, which was ultrasonicated for 5 min to obtain a homogeneous solution. The reaction kettle was sealed and put into an oil bath at 120 °C for 48 h with magnetic stirring (Fig. 1a). When the reaction finished, the reactor was taken out and cooled to room temperature. The obtained white products were washed with 0.1 mol L⁻¹ HNO₃ and centrifuged several times until the pH reached 7. The final products were washed with deionized water and placed in a vacuum drying oven at 60 °C for 8 h.

**Preparation of TiO₂@C core–shell structures**

The as-prepared products were placed in the middle of a tube furnace, N₂ gas was purged through anhydrous ethanol and then into the tube furnace, and the reaction temperature was set at 400 °C for 10 h under a nitrogen atmosphere. Black products (TiO₂@C composite) were obtained as the reaction finished. The experimental schematic diagram is shown in Fig. 1b.

**Characterization**

The prepared TiO₂@C composites were characterized by scanning electron microscopy (SEM, S-4800, Japan) and transmission electron microscopy (TEM, JEM-2100F, Japan). The crystal structure and components of the material were determined using an X-ray diffractometer (XRD, Rigaku, Cu-Kα radiation, Japan). The specific surface area and the pore size distribution of the material were determined from the Brunauer–Emmett–Teller (BET) test (ASAP 2020, Micromeritics, America). The surface chemical structure was determined by Fourier transform infrared spectroscopy (FT-IR) (6700, Nicolet, USA) and Raman spectroscopy (Lab RAM, HR800) with 532 nm laser light. The absorbance of the TiO₂@C composite materials was measured using a UV spectrophotometer (G-9, Rangqi, China). The functional groups and the elemental states on the surface of the adsorbents were examined by X-ray photoelectron spectroscopy (XPS) analysis, which was conducted on a Kratos Axis Ultra DLD spectrometer, using monochromated Al Kα X-rays at a base pressure of 1 × 10⁻⁹ Torr.

**Results and discussion**

The result of XRD analysis is shown in Fig. 2; the main diffraction peaks could be indexed to the (101), (103), (200), (105), (213), (116) and (107) reflections, corresponding to anatase phase TiO₂ with the JCPDS card no. 01-0562. The result also confirms that C exists in the TiO₂ products (marked with red asterisks). The C species were further tested by Raman spectroscopy and the result is shown in Fig. S1.† The peak intensity ratio between the G band and D band (IG/ID) is 1.1796, indicating that the proportion of carbon defect structures in the carbon structure is relatively small and the dominant component in the carbon structures is the sp².

The low- and high-magnification SEM images of the TiO₂@C products are shown in Fig. 3a and b. From the SEM images, it

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Fig. 1 Schematic of the preparation process for TiO₂@C.

Fig. 2 XRD pattern of the obtained products.

Fig. 3 Microstructure characterization of the core–shell TiO₂@C nanotubes: (a and b) low and high-magnification SEM images, and (c) TEM and (d and e) HRTEM images.
can be seen that the length of the TiO$_2$@C nanotubes can be up to several micrometers, representing an ‘ultralong’ structure. From the TEM image of Fig. 3c, it can be confirmed that the composite material is a tubular nanostructure. More detailed information on the material can be obtained from Fig. 3d; the outer diameter of the nanotubes is ~35 nm, and the inner diameter is ~15 nm, and the thickness of the carbon layer is 5–6 nm (Fig. 3e). This shows that the carbon layer was successfully coated on the outside of the TiO$_2$ nanotubes, and the interplanar spacing of 0.35 nm indicates the (101) lattice plane of the TiO$_2$ crystal.

The detailed specific surface area of the TiO$_2$@C core–shell nanotubes can be obtained from the N$_2$ adsorption–desorption curve (Fig. 4a). The specific surface area of TiO$_2$@C can be calculated from the N$_2$ adsorption–desorption isotherm, and the BET value is 346.26 m$^2$ g$^{-1}$, which is higher than that of the single TiO$_2$ (Fig. S2 in the ESI). Fig. 4b shows the pore size distribution of the TiO$_2$@C nanotubes, with a main diameter of around 15 nm, which corresponds to the diameter of the nanotubes. The diameter of ~60 nm can be attributed to the gaps between the nanotubes.

The adsorption of the three kinds of antibiotics is based on the classical monolayer Langmuir model and the multilayer Freundlich adsorption model. The Langmuir equation depicts the process of randomly adsorbing species onto the surface of the adsorbent. The linearization equation of the Langmuir adsorption model is as follows:

$$
\frac{c_e}{q_e} = \frac{1}{K_L q_m} + \frac{c_e}{q_m}
$$

where $c_e$ is the balance concentration (mg L$^{-1}$), $K_L$ is the adsorption constant (L mg$^{-1}$), $q_m$ is the maximum adsorption capacity (mg g$^{-1}$), and $q_e$ represents the quantity (mg g$^{-1}$) of antibiotics adsorbed at equilibrium.

The Freundlich adsorption model is an empirical equation that is simple and widely used. It can simulate a multi-layer adsorption process, which means that the adsorption sites on the adsorption surface are not uniform. The linearized form of the Freundlich adsorption model equation is as follows:

$$
\ln q_e = \ln K_F + \left(\frac{1}{n}\right) \ln C_e
$$

where $K_F$ and $n$ represent the Freundlich constant [(mg g$^{-1}$) (L mg$^{-1}$)$^{1/n}$] and adsorption intensity of the adsorbents, respectively. The mass of the TiO$_2$@C materials used in this study was 5 mg, for adsorption of 5–40 mg L$^{-1}$ tetracycline (TC), 5–40 mg L$^{-1}$ ofloxacin (OFO) and 2–20 mg L$^{-1}$ norfloxacin (NFO). Adsorption equilibrium was achieved after magnetic stirring at room temperature. Fig. 4 shows the experimental data of the TiO$_2$@C nanotubes adsorbing the three kinds of antibiotics: OFO, TC and NFO. Fig. 5a and b are the Langmuir model fitting charts, and Fig. 5c and d show the corresponding Freundlich model fitting charts. The fitting data are shown in Table 1. From the correlation coefficient ($R^2$) value of the two models, it can be concluded that the three kinds of antibiotics adsorbed by the material are more consistent with the Langmuir model. The maximum adsorption capacities of OFO, TC and NFO, calculated using the Langmuir model, are 232 mg g$^{-1}$, 240 mg g$^{-1}$, and 190 mg g$^{-1}$, respectively. The experimental results show that the adsorption amount of this material is greatly improved compared with other inorganic materials, as shown in Table 2.

### Adsorption kinetic model

The adsorption kinetics of the three kinds of antibiotics were studied. The pseudo-first-order model was based on the assumption that the adsorption was controlled by diffusion. The pseudo-first-order dynamic model is as follows:

$$
\ln(q_e - q_t) = \ln q_e - K_1 t
$$
The pseudo-second-order kinetic model assumes that the adsorption rate is determined by the square value of the number of adsorbed vacancies on the surface of the adsorbent. The pseudo-second-order dynamic model is as follows:\(^{(4)}\)

\[
\frac{t}{q_t} = \frac{1}{K_2q_e^2} + \frac{t}{q_e}
\]

where \(q_e\) is the concentration of adsorption at equilibrium (mg g\(^{-1}\)), \(q_t\) is the amount of adsorption (mg g\(^{-1}\)) at time \(t\), \(K_1\) is the pseudo-first-order adsorption rate constant, and \(K_2\) is the pseudo-second-order adsorption rate constant.\(^{(17)}\) The fitting data of the dynamics of the three kinds of antibiotics are given in Table 3.

Fig. 6 shows the pseudo-first-order and pseudo-second-order kinetic models of the three antibiotics, and the intra-particle-diffusion model. As Table 3 shows, the pseudo-first-order model best describes the adsorption mechanism. Thus, the adsorption process of TiO\(_2@C\) adsorbing OFO, TC and NFO is mainly controlled by diffusion.\(^{(22)}\) In order to obtain more information on the adsorption process, data are fitted to the intra-particle-diffusion model, as follows:\(^{(34)}\)

\[
q_t = K_{id}t^{1/2} + C
\]

Adsorption can be divided into two processes: adsorbent surface adsorption and slow pore diffusion. The particle diffusion model is the most suitable for describing the dynamics of particles in the particle diffusion process.\(^{(34)}\) The linear fitting of \(q_t\) and \(t^{1/2}\) gives a straight line that passes through the origin and shows that particle diffusion is the rate determining step.\(^{(15)}\) From Fig. 6c, it can be seen that the TiO\(_2@C\) particle diffusion model is divided into three stages; the first phase is the diffusion of the boundary layer, which is mainly the external mass transfer of the adsorbent. The second phase refers to the internal diffusion of particles, which is mainly the transport of microstructures of adsorbents, such as holes. These two phases make up the intra-particle diffusion model. Because the contact angle between the material itself and the solution is different, the time needed for different materials to reach adsorption equilibrium is different. The third phase is usually not considered as a rate-controlling step, generally referring to the procession of adsorption being close to adsorption equilibrium, which means that the active sites of the internal and external surfaces are mostly occupied. Compared with single titanium dioxide nanotube materials, the contact angle of the composites changed due to the synergistic effect of titanium dioxide nanotubes and the carbon layer, which greatly improved the adsorption performance of the material\(^{(16)}\) (Fig. S3\(^\ddagger\)).

In order to understand the change that has occurred on the surface of the TiO\(_2\) nanotubes, FTIR and XPS tests were carried out and TC was selected as an example. Fig. 7a shows the FTIR comparison, before and after the adsorption of TC. The

| Antibiotics     | Adsorbents       | \(q_m\) (mg g\(^{-1}\)) | Conditions | Ref. |
|-----------------|------------------|--------------------------|------------|-----|
| Tetracycline    | Zeolite          | 130                      | pH = 7     | 14  |
|                 | GSO              | 137                      | pH = 7     | 23  |
|                 | PS/GO            | 197.9                    | pH = 6     | 24  |
|                 | PC@CMCS          | 136.9                    | pH = 7     | 25  |
|                 | Ultralong hydrogen titanate nanobelts | 151.51       | pH = 7     | 26  |
|                 | TiO\(_2@C\)      | 240                      | pH = 7     | This work |
| Norfloxacin     | AMPH             | 1.42                     | pH = 7     | 27  |
|                 | MIP              | 29.35                    | pH = 8     | 28  |
|                 | Ultralong hydrogen titanate nanobelts | 111.73       | pH = 7     | 26  |
|                 | TiO\(_2@C\)      | 190                      | pH = 7     | This work |
| Ofloxacin       | Oil shale powders| 15.4                     | pH = 7.5   | 16  |
|                 | Ultralong hydrogen titanate nanobelts | 148.14       | pH = 7     | 26  |
|                 | TiO\(_2@C\)      | 232                      | pH = 7     | This work |

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**Table 2** The adsorption capacity comparison of the adsorbent materials

| Antibiotics | Adsorbents        | \(q_m\) (mg g\(^{-1}\)) | Conditions | Ref. |
|-------------|-------------------|--------------------------|------------|-----|
| OFO         |                   |                          |            |     |
| TC          |                   |                          |            |     |
| NFO         |                   |                          |            |     |

**Table 3** OFO, TC and NFO dynamic correlation fitting data

| Model | \(C_0\) | \(K_1\) | \(q_{e,cal.}\) | \(R^2\) | \(K_2\) | \(q_{e,cal.}\) | \(R^2\) |
|-------|--------|--------|----------------|--------|--------|----------------|--------|
| OFO   | 40     | 0.00647| 230.2          | 0.95   | 7.32 \times 10^{-6} | 769.2 | 0.52 |
| TC    | 40     | 0.00909| 252.1          | 0.97   | 2.84 \times 10^{-6} | 454.5 | 0.50 |
| NFO   | 20     | 0.00874| 191.2          | 0.97   | 2.99 \times 10^{-5} | 248.7 | 0.47 |
conversion of the amide group. A hydrogen ion was introduced into the soaking TiO2 nanotubes from dilute nitric acid, combined with a small amount of carbon into the C\(\text{-}\text{H}\) bond. The absorption peak at 1384 cm\(^{-1}\) is the result of the N-O bond introduced by TiO2 nanotubes with dilute nitric acid.\(^3\) The smaller absorption peaks at 1066 cm\(^{-1}\) and 903 cm\(^{-1}\) are due to the telescopic vibration of the C-O bond. The absorption peak of TiO2@C at 1609 cm\(^{-1}\) is the characteristic peak of the amide group. After adsorption of TC, the peak shifted to 1630 cm\(^{-1}\), indicating that TC was successfully adsorbed on TiO2@C. The sharp peak at 468 cm\(^{-1}\) is the result of the vibration of the Ti-O bond of titanium dioxide nanotubes. In addition, the carbonyl group is formed by the combination of some oxygen elements, which makes the Ti-O bond peak of the TiO2@C composite blunt and the peak shifts to 497 cm\(^{-1}\). The analysis of FT-IR results showed that the adsorption of TiO2@C on TC was mainly physical adsorption, which was in accordance with the pseudo-first-order kinetic model. To further prove that TC has been adsorbed by TiO2@C, XPS spectra (Fig. S4\(^\dagger\)) of the C 1s peaks of adsorbents before and after adsorption are compared. After deconvolution, the C 1s signal can be separated into a C=C\(\text{-}\text{C}\) peak at 284.5 eV, a C-O peak at 285 eV and a C=O peak at 286.2 eV. Simultaneously, the content of C=C\(\text{-}\text{C}\) increased sharply. In contrast, the content of C=O is the lowest among the three kinds of chemical bonds. In summary, the increase of C=C\(\text{-}\text{C}\) is due to the benzene ring of TC. When the TC antibiotic was adsorbed by TiO2@C, the C=O would be replaced by C=C\(\text{-}\text{C}\). Thus, it is noted from the XPS spectra that the TC antibiotic can be adsorbed by TiO2@C effectively.

The D-R isotherm model can be applied to identify the type of adsorption, i.e., physical or chemical adsorption process. The linear D-R isotherm\(^27\) models are as follow and the results are shown in Fig. 8:

\[
\ln q_e = \ln q_m - B_D e \tag{6}
\]

\[
\epsilon = RT \ln(1 + 1/C_e) \tag{7}
\]

where \(q_e\) and \(q_m\) (mmol g\(^{-1}\)) are the equilibrium antibiotics solution concentration on the adsorbent and maximum adsorption capacity, respectively, \(B_D\) is a constant related to the adsorption energy (mol\(^2\) J\(^{-2}\)), \(\epsilon\) is the Polanyi potential, \(C_e\) is the equilibrium concentration (mol L\(^{-1}\)), \(R\) is the gas constant and \(T\) is the absolute temperature (K).

The adsorption free energy \((E)\) can be calculated according to Polanyi potential from the following equation:

\[
E = 1/\sqrt{2} \times B_D \tag{8}
\]

Compared with the value of \(R^2\), it is obvious that the Langmuir model is more fit for the experimental data. As for the D-R equations, if the value of \(E\) is less than 8 kJ mol\(^{-1}\), physical sorption is considered as the main mechanism, while chemical sorption will be expected if the value of \(E\) is in the range of 8-16 kJ mol\(^{-1}\). As shown in Table 4, the three values of \(E\) are all less than 8 kJ mol\(^{-1}\). This proves that physisorption is the main sorption mechanism. The results of this analysis yielded consistent pseudo-first-order adsorption kinetics.

The thermodynamics parameters regarding antibiotic adsorption onto TiO@C, such as Gibbs free energy \((\Delta G)\), and the solid and liquid phases at equilibrium \((K_d)\), can be calculated using the following equations:\(^{28}\)

\[
K_d = q_d/C_e \tag{9}
\]

\[
\Delta G = -R \ln(K_d) \tag{10}
\]

Generally, if \(-20 < \Delta G < 0\) kJ mol\(^{-1}\), physical adsorption is dominant and it changes to more negative values than
The TiO$_2@$C exhibits a mesoporous structure and large specific surface area. The adsorption experiment proves that the TiO$_2@$C composite adsorbent can adsorb three kinds of typical antibiotics (TC, OFO, and NFO) effectively. Moreover, the main determinant of the adsorption process is proved to be physical adsorption rather than chemical adsorption, and the adsorption process fits the Langmuir and pseudo-first-order models. Therefore, the work provides a practical case for the future development of more composite adsorbents and lays a foundation for the industrialization of composite materials.

Conclusions

Core–shell TiO$_2@$C ultralong nanotubes were prepared using a solvothermal process followed by anhydrous ethanol carbonization. The TiO$_2@$C exhibits a mesoporous structure and large specific surface area. The adsorption experiment proves that the TiO$_2@$C composite adsorbent can adsorb three kinds of typical antibiotics (TC, OFO, and NFO) effectively. Moreover, the main determinant of the adsorption process is proved to be physical adsorption rather than chemical adsorption, and the adsorption process fits the Langmuir and pseudo-first-order models. Therefore, the work provides a practical case for the future development of more composite adsorbents and lays a foundation for the industrialization of composite materials.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The work has been funded by the National Natural Science Foundation of China (51602193, 21601122, and 51701022), the Shanghai “Chen Guang” Project (16CG63), the Fundamental Research Funds for the Central Universities (WD1817002), and the Talent Program of Shanghai University of Engineering Science, Science and Technology Facilities Council for STFC/MDC Futures Early Career Award.

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