


Cite this: *RSC Adv.*, 2024, 14, 26103

# Preparation of hydroxyapatite and its elimination of excess fluoride from aqueous solution†

Yiming Zou,<sup>abcd</sup> Yaoting Wang,<sup>abcd</sup> Junfu Wang,<sup>abcd</sup> Song Wang,<sup>abcd</sup> Li Chen,<sup>abcd</sup> Yu Xi,<sup>abcd</sup> Ruliang Xie,<sup>\*abcd</sup> Jie Yang<sup>\*bc</sup> and Xin Xiao<sup>id</sup> <sup>\*cd</sup>

Excess fluoride in aqueous solutions can significantly affect dental and bone health. This study used two methods to prepare hydroxyapatite to remove fluoride ions from water. The experiments showed that the adsorption capacity and removal rate of hydroxyapatite (Xq-HAP) prepared by the novel method were higher than for the hydroxyapatite (Yt-HAP) prepared by the conventional method. The maximum fluoride ion trapping capacity of Xq-HAP could reach 29.04 mg g<sup>-1</sup> under the conditions of pH = 5 and an F ion concentration of 10 mg L<sup>-1</sup>. The materials were characterized by SEM, XRD, BET, XPS, and FTIR. An investigation was conducted to examine the impact of contact time, adsorbent dosage, fluoride concentration, solution pH, temperature, and several other parameters on the removal of fluoride. Adsorption equilibrium was reached in approximately 3 h at an initial fluoride concentration of 10 mg L<sup>-1</sup>. It can be seen that the adsorbent has a faster ability to trap fluoride ions. The adsorption kinetics and Langmuir isotherm indicated that fluoride ion adsorption is a monolayer chemisorption process. Further characterization and kinetic studies indicated that the removal mechanism involves ion exchange, electrostatic interactions, and complexation. After five adsorption cycles, the adsorption capacity reaches 23 mg g<sup>-1</sup>.

Received 21st March 2024

Accepted 6th July 2024

DOI: 10.1039/d4ra02147a

rsc.li/rsc-advances

## 1. Introduction

Heavy metals discharged into water bodies through industrial operations and human activities, even at trace levels, are considered a significant risk with long-term effects.<sup>1</sup> Fluorine is an element widely found in nature and is a trace element essential for human growth. The World Health Organization (WHO) has stipulated the maximum permissible concentration of fluoride as 1.0 mg L<sup>-1</sup> in the Hygienic Standard for Drinking Water for Domestic Use (GB 5749-2022).<sup>2</sup> Fluoride contamination of natural water has become a global problem because the semiconductor, electroplating, metallurgical, and ceramic industries discharge high concentrations of fluoride-containing wastewater. Some domestic factories involved in fluoride wastewater treatment have imperfect technology and equipment, resulting in the concentration of fluoride ions in the discharged wastewater being much higher than the national

emission standards.<sup>3</sup> Groundwater has been reported to be contaminated with high fluoride concentrations in some parts of the world, including Asia, Africa, Europe, and the Americas.<sup>4</sup> Fluoride-containing wastewater discharged more than the standard enters surface water through the scouring of surface runoff, and the long-term intake of excessive fluoride can cause the chronic disease fluorosis,<sup>5</sup> which has a significant impact on the structure and function of the human skeletal system, teeth, skeletal muscles, brain, and spinal cord,<sup>6</sup> and is hazardous to human health.

Therefore, rapid and effective technologies for treating excess fluoride in water are urgently needed.<sup>7</sup> Fluoride removal methods include the precipitation method,<sup>8</sup> ion exchange method,<sup>9</sup> membrane separation method<sup>10</sup>, and adsorption method.<sup>11</sup> The adsorption method has excellent potential for fluoride removal because of its advantages of low cost, simple operation, high removal rate, and reusable adsorbents.<sup>12</sup> From the classification of adsorbents, various adsorbents have been used in fluoride removal treatment, including apatite, activated carbon, activated alumina, biosorbent, nano-sorbent, and so on.<sup>13</sup> However, conventional adsorbents have disadvantages, such as few activation sites, poor treatment effect, weak affinity, low adsorption capacity, and their applications are limited.<sup>14</sup> Hydroxyapatite (Ca<sub>10</sub>(PO<sub>4</sub>)<sub>6</sub>(OH)<sub>2</sub>), which has low water solubility, low cost, high stability, easy accessibility, and good adsorption capacity, has now become a suitable material for the adsorption of fluoride ions in water.<sup>15</sup> The main adsorption

<sup>a</sup>Jiangsu Key Laboratory of Marine Bioresources and Environment, Jiangsu Ocean University, Lianyungang 222005, China. E-mail: xieruliang2007@163.com

<sup>b</sup>School of Ocean Food and Biological Engineering, Jiangsu Ocean University, Lianyungang 222005, China. E-mail: yangjie@jou.edu.cn

<sup>c</sup>Jiangsu Institute of Marine Resources Development, Lianyungang 222005, China. E-mail: xiaoxin@njst.edu.cn

<sup>d</sup>Jiangsu Key Laboratory of Function Control Technology for Advanced Materials, Jiangsu Ocean University, Lianyungang 222005, China

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ra02147a>



mechanisms of hydroxyapatite are electrostatic interactions, ion exchange, acid–base interactions, surface complexation, and dissolution<sup>16</sup>/precipitation.<sup>17</sup> Related experiments showed that the adsorption of fluoride ions decreases under acid–base conditions, and therefore, the study of solution pH is of great significance in the process of fluoride removal.<sup>18</sup> Furthermore, HAP has demonstrated exceptional efficacy in various catalytic applications, including environmental catalysis, energy catalysis, chemical synthesis catalysis, biocatalysis, and photocatalysis.<sup>19,20</sup> HAP is a catalyst that can fulfill these requirements. In organic synthesis, it can serve as a bifunctional acid–base catalyst for various organic reactions, such as esterification, amidation, and dehydration condensation. The acidity and basicity of its surface can regulate the reaction pathway, improving selectivity and yield.<sup>21</sup> A method was devised to gently remove the protective layer from alkylthio-gold nanoclusters (Au NCs) on hydroxyapatite (HAP) using hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) as the oxidizing agent. After oxidation, the thiol ligands on the supported Au NCs were removed, and the size and integrity of the supported Au were well preserved.<sup>22</sup> By pyrolyzing chicken bone waste at different temperatures (400 °C, 500 °C, 650 °C, and 800 °C), the influence of carbon residues in the bone marrow on the dispersion of Cu species introduced *via* impregnation with Cu(NO<sub>3</sub>)<sub>2</sub> solution was investigated. The results indicated that the hydroxyapatite formed after the pyrolysis of chicken bones was relatively stable, making the catalyst exhibit good activity and reusability.<sup>23</sup>

Most mussel shells are calcium carbonate with a content of up to 95% and contain small amounts of protein and polysaccharide substances.<sup>24</sup> The shell structure exhibits aragonite, calcite, spherical aragonite, and amorphous calcium carbonate,<sup>25</sup> which can be categorized in order of their stability from low to high: spherical aragonite, aragonite, and calcite.<sup>26</sup> Therefore, it can be used as an excellent green calcium source for the preparation of hydroxyapatite. Currently, the utilization of shellfish is limited to the edible portion, whereas the shell portion is seldom processed or utilized.<sup>27</sup> In 2012, the annual production of mussels in China amounted to  $5 \times 10^5$  tons, and the shells of mussels, which accounted for more than 1/3 of the weight of mussels, were discarded, and only one year could produce  $1.5 \times 10^5$  to  $1.8 \times 10^5$  tons.<sup>28</sup> The accumulation of large quantities of shells has caused severe pollution of land and air, and owing to the difficulty of natural decomposition, it has become a severe environmental problem in some areas of the southeastern coast of China. At the same time, the accumulated shells are a waste of resources. Owing to its robust ion exchange and biocompatibility,<sup>29</sup> HAP is regarded as a potential adsorbent. Therefore, using hydroxyapatite for wastewater treatment is essential for environmental protection and human health.

This study presents a process for preparing rod-shaped hollow hydroxyapatite using mussel shells as a template and adjusting the reaction time. This study aimed to synthesize HAP particles with fluoride ion adsorption performance. The effects of the contact time, adsorbent dosage, initial concentration, reaction temperature, and solution pH on the adsorption of fluoride ions by HAP were examined. The suitability of rod-shaped HAP as an adsorbent was investigated by determining

the adsorption isotherms and kinetic parameters. This study reveals that the adsorption of fluoride ions from aqueous solutions by rod hydroxyapatite particles paves the way for future research on more efficient and economical means of wastewater pollution purification.

## 2. Experimental

### 2.1 Materials

(NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub> (99%), HCl (99%), NaOH (99%), C<sub>3</sub>COOH (99%), and NaF (99%) were purchased from Shanghai McLean Co. All chemicals were of analytical grade and were used without further purification. Mussel shells were collected from the coastal edge of Lianyungang.

### 2.2 The synthesis of Xq-HAP and Yt-HAP

**2.2.1 Xq-HAP.** Mussel shells were pretreated in a muffle furnace at 220 °C for 24 h. Pretreated mussel shells (5 g) were mixed with 75 mL of 5% aqueous acetic acid solution in a 250 mL conical flask and shaken on a shaking table at 180 rpm for 24 h. The resulting colorless calcium acetate solution was poured out, and the remaining shells were rinsed three times with distilled water, dried, and mixed again with 40 mL of 5% aqueous acetic acid solution. This shaking step was repeated for 24 h. At the end of the process, the solution was poured out, and the remaining shells were rinsed and dried once again, at which point soft calcite (SC) was obtained. The obtained soft calcite was finely ground and 2 g of the powder was mixed with 1.58 g of diammonium hydrogen phosphate in a single-necked flask to obtain a Ca/P molar ratio of 1.67. Subsequently, 35 mL of deionized water was added, and after stirring until homogeneity was achieved, the pH of the solution was adjusted to 9.5. This mixture was then placed in an oil bath and incubated at 95 °C for 7 h. At the end of the reaction, the solution was washed to neutral pH using a vacuum filtration unit, and the solid obtained was dried in an oven at 120 °C for 4 h. The final white powder obtained is denoted as Xq-HAP, as shown in Fig. 1.

**2.2.2 Yt-HAP.** Mussel shells were first calcined in a muffle furnace at 220 °C for 24 h, and then uniformly broken and sieved through a 600 mesh using a cell breaker. 2 g sieved powder was mixed with 1.58 g of diammonium hydrogen

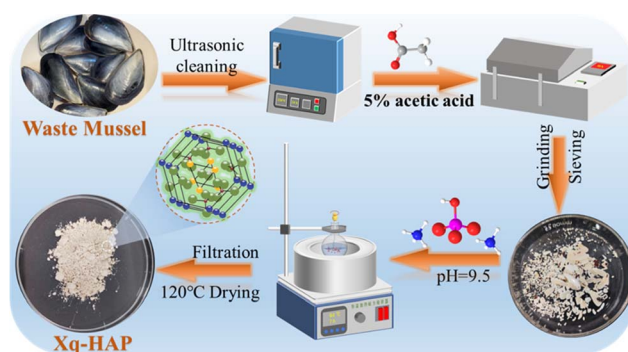


Fig. 1 Schematic illustration of the procedure for the synthesis of Xq-HAP.



phosphate in a single-necked flask, 35 mL of deionized water was added, and the pH was adjusted to 9.5. The mixture was incubated in an oil bath at 95 °C for 7 h. After the reaction, the solution was neutralized by washing with a vacuum filtration device, and then the solid was dried in an oven at 120 °C for 4 h. The final white powder is denoted as Yt-HAP. The resulting white powder was denoted as Yt-HAP.

### 2.3 Characterizations

The phase structure was determined *via* powder X-ray diffraction (XRD) using a RINT 2000 diffractometer (Rigaku) with Cu K $\alpha$  radiation. The morphology of the product was determined using a Hitachi S-4800 field-emission scanning electron microscope (SEM) and a JEOL JEM 2100F transmission electron microscope (TEM). The Brunauer–Emmett–Teller (BET) surface areas were determined using a Micromeritics ASAP 2460 instrument. X-ray photoelectron spectra (XPS) were recorded using a Thermo ESCALAB 250XI instrument. The specific surface areas and porosities of the samples were measured at 77 K using an N<sub>2</sub> adsorption analyzer (Micromeritics ASAP 2020 Plus). Fourier-transform infrared spectroscopy (FT-IR, nanoIR3-s) was used to analyze the characteristic functional groups on the HAP surface.

### 2.4 Adsorption experiments

All adsorption experiments were carried out in 100 mL conical flasks containing 50 mL of NaF solution and 50 mg of HAP adsorbent. The relationship between time and adsorption capacity was investigated using adsorption experiments at an initial fluorine concentration of 10 mg L<sup>-1</sup>. The fluoride ion concentration in the samples was measured using a PXSJ-226T ion meter and the adsorption capacity was calculated. The adsorption primary kinetics, secondary kinetics, and internal diffusion adsorption models were plotted to conduct a kinetic study. Adsorption isotherms were simulated at initial NaF concentrations of sodium fluoride ranging from 10 to 100 mg L<sup>-1</sup>. These conical flasks were placed in a shaker at temperatures (25 °C, 35 °C, and 45 °C) and a rotational speed of 150 rpm. Moreover, the residual fluoride ion concentration was measured after a certain period, and the adsorption capacity was calculated by fitting curves through Langmuir and Freundlich models. To investigate the effects of pH on the effectiveness of the adsorbent in removing fluorine, the pH was adjusted using 0.1 mol of HCl and NaOH solution. In the cyclic adsorption experiment, 0.5 mol L<sup>-1</sup> NaOH and 0.5 mol L<sup>-1</sup> NaCO<sub>3</sub> solution was selected as the desorption liquid, and its adsorption capacity was calculated after five experiments to judge its recycling capacity. The adsorption capacity ( $q_e$ , mg g<sup>-1</sup>) was calculated using the following equation:

$$q_e = \frac{(C_0 - C_e)V}{m}$$

where  $C_0$  (mg L<sup>-1</sup>) and  $C_e$  (mg L<sup>-1</sup>) are the initial and equilibrium concentrations of the fluoride solution, respectively;  $V$  (L) is the volume of the solution, and  $m$  (g) is the mass of the adsorbent.

## 3. Results and discussion

### 3.1 Characterization

Fig. 2 shows the XRD patterns of Xq-HAP and Yt-HAP. The prominent diffraction peaks of hydroxyapatite appear at  $2\theta = 22.8^\circ, 25.9^\circ, 28.9^\circ, 31.7^\circ, 32.3^\circ, 32.9^\circ, 34.04^\circ$ , and  $35.4^\circ$ , corresponding to the (111), (002), (210), (211), (112), (300), (202), and (301) crystal planes, respectively. All intensity peaks in the XRD spectrum of the hydroxyapatite powder are in perfect agreement with the structural data of hydroxyapatite, and the pattern is consistent with the hexagonal crystal structure of hydroxyapatite (JCPDS No. 09-0432).

The morphology of the Xq-HAP is shown in Fig. 3(A–D), a rock structure with a rough surface. A rough surface makes the adsorption performance more stable, and fluoride is less likely to come off. The diameters of these structures range from about 25–50 nm, with an average value of approximately 35 nm. As shown in Fig. 3(E and F), Yt-HAP exhibited an irregular clustered structure, and some clusters and agglomerates with rod-like structures were also observed. The EDS elemental surface sweep of Xq-HAP is shown in Fig. 3(G–J), and C, O, P, and Ca were uniformly distributed on the surface of Xq-HAP. The point-scan plot of the EDS elements of Xq-HAP is shown in Fig. 3(k), and C, O, P, and Ca were also distributed on the surface of Xq-HAP.

Infrared spectroscopy is an essential way to study the changes in functional groups before and after adsorption. As shown in Fig. 4, the phosphate PO<sub>4</sub><sup>3-</sup> vibrational modes typically exhibit absorption peaks at 900–1200 cm<sup>-1</sup>. The stretching vibration of the P–O bond may cause the peak to appear at approximately 1040 cm<sup>-1</sup>. There is a possibility of ion exchange between the phosphate ion PO<sub>4</sub><sup>3-</sup> in Xq-HAP and fluoride ions in the solution. Since fluoride ions have a smaller ionic radius and a higher charge density, they are more likely to bind to Ca<sup>2+</sup> on the surface of hydroxyapatite, thereby replacing some of the phosphate ions and generating a CaF<sub>2</sub> precipitate. The broad peak at 3570.4 cm<sup>-1</sup> is the absorption peak of the stretching vibration of the O–H bond in Xq-HAP, and the characteristic peak of the hydroxyl group almost disappeared after adsorption,

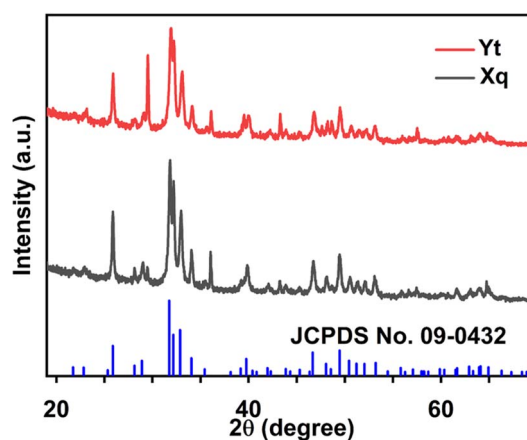


Fig. 2 XRD pattern of Xq-HAP and Yt-HAP.





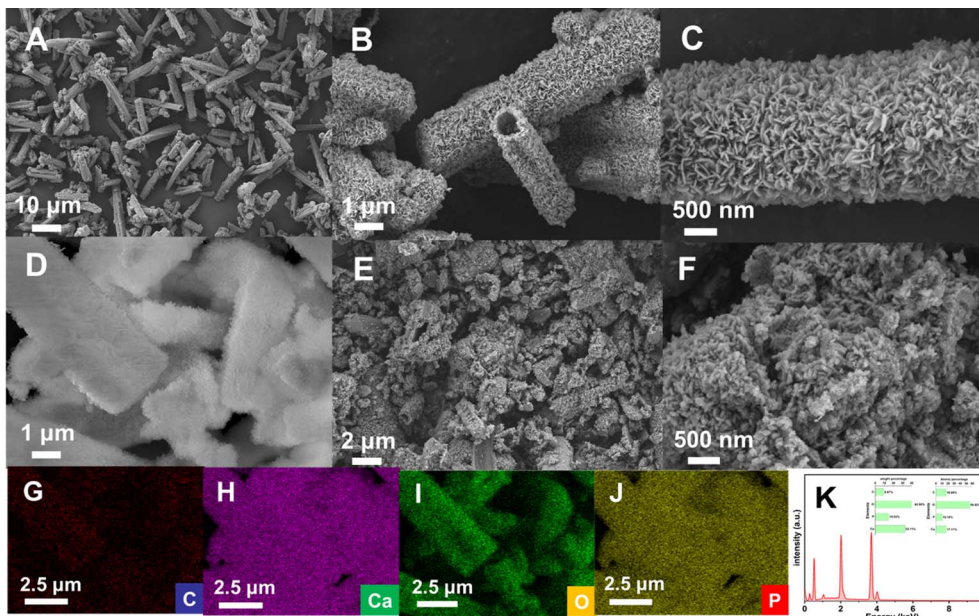


Fig. 3 (A–D) Xq-HAP SEM images, (E and F) Yt-HAP SEM images, (G–J) EDS elemental mapping for C, Ca, O and P respectively. (K) EDS spectrum of Xq-HAP.

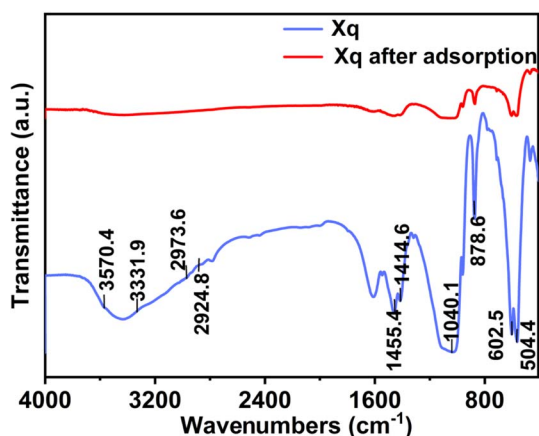


Fig. 4 FT-IR spectrum of Xq-HAP before and after adsorption.

which suggests that  $F^-$  was ion-exchanged with  $OH^-$ . During the adsorption process, the hydroxyl groups on the surface of hydroxyapatite may react chemically with fluoride ions to form new chemical bonds, leading to changes in the characteristic peaks of the hydroxyl groups and the appearance of new peaks. Through the analysis of infrared spectrograms, the characteristic peaks of phosphate ions and hydroxyl groups changed before and after the adsorption of Xq-HAP, indicating that the adsorption process was a chemical reaction dominated by ion exchange, revealing the mechanism of the fluoride ions adsorbed by hydroxyapatite.

XPS characterization was used to analyze the chemical states of three elements, Ca, O, and P, on the surface of Xq-HAP. The XPS spectrum of Xq-HAP is shown in Fig. S1.† The two peaks of elemental carbon correspond to 284.97 eV and 289.52 eV, respectively, and the binding energies correspond to the

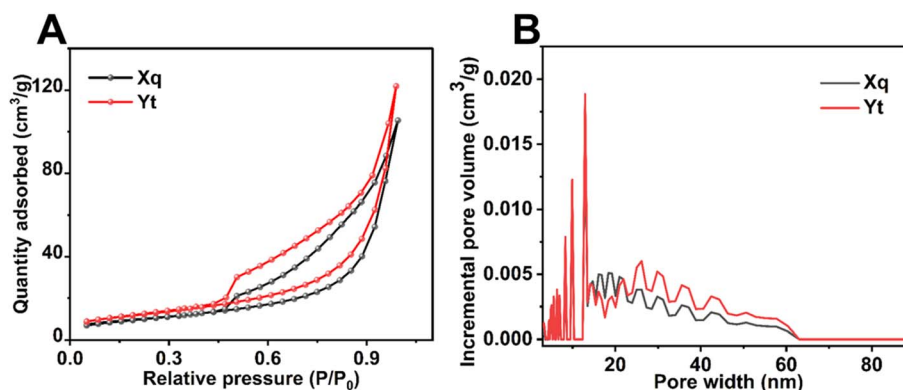


Fig. 5 (A) Nitrogen adsorption–desorption isotherms and (B) the BJH pore-size distribution curve of Xq-HAP and Yt-HAP at different reaction time.



chemical bonding of C–C or C=C or C=O, respectively. The two characteristic peaks are located at 347.39 eV and 350.98 eV, attributed to the  $2p_{3/2}$  and  $2p_{1/2}$  orbitals of Ca, respectively, which coincide with the typical binding energies of the Ca element. The single peak of P corresponds to a binding energy of 133.45 eV, which is the characteristic binding energy of the P element in Xq-HAP. In addition, the binding energy of O 1s is 531.38 eV, which is attributed to  $\text{OH}^-$ , and the ratio of the peak areas of Ca and P elements is approximately 1.67, indicating that the prepared Xq-HAP material is pure, well-structured, and hard.

Fig. 5(A and B) shows the  $\text{N}_2$  adsorption and desorption curves of the two adsorbent materials, and it can be seen that the adsorption curves of the two materials conform to the IV-type curves, which indicates that the pore columns in the materials are micropores. As shown in Table 1, the specific surface area of Yt is  $42.6 \text{ m}^2 \text{ g}^{-1}$ , which is higher than that of Xq ( $35.3 \text{ m}^2 \text{ g}^{-1}$  and the pore volume of Yt ( $0.122 \text{ m}^3 \text{ g}^{-1}$ ) was also higher than that of Xq ( $\text{m}^3 \text{ g}^{-1}$ ). However, Xq possesses a larger average pore size, indicating that Xq mediates more pore structure than Yt material. Fig. 5(B) shows the pore-size distributions of the two materials. Both exhibit multistage pore

structures, and mediator pore structures play a significant role in adsorption. These multistage pore structures facilitate the adsorption of pollutant molecules.

### 3.2 Adsorption property

**3.2.1 Effect of contact time.** In a conical flask, (0.02 g, 0.04 g, 0.05 g, and 0.1 g) were weighed and poured into 50 mL of a fluoride ion solution at an initial concentration of  $10 \text{ mg L}^{-1}$  at  $25^\circ\text{C}$  for the experiment. Fig. S2(A)† shows the adsorption efficiency of Xq-HAP for fluoride ions over time. Fig. S2(B)† shows a graph of the adsorption capacity of Xq-HAP for fluoride ions over time. Fig. S2(C)† shows a graph of the adsorption efficiency of Yt-HAP for fluoride ions over time. Fig. S2(D)† shows the fluoride ion desorption capacity of Yt-HAP over time. As can be seen from the figure, the removal effect hardly changed after the experiment was carried out for 180 min, and the adsorption equilibrium time for this experiment was 180 min.

**3.2.2 Effect of the amount of adsorbent.** The effects of the dosages of the two adsorbent materials on the removal of fluoride ions were further investigated. The fluoride ion solution chosen for the experiment was 50 mL and the initial concentration was  $10 \text{ mg L}^{-1}$ . The dosages of Xq-HAP and Yt-HAP adsorbent materials were 20 mg, 40 mg, 50 mg, and 100 mg, respectively, and the experiments were carried out at  $25^\circ\text{C}$ , with adsorption equilibrium reached within 180 min. The adsorption results are shown in Fig. 6, and the removal rate of fluoride ions gradually increased when the dosage of the Xq-HAP adsorbent material was increased from 20 mg to 40 mg. The fluoride ion removal rate was 55.74% at a dosage of 50 mg, and the adsorption capacity for fluoride ions reached  $27.84 \text{ mg}$

Table 1 Specific surface area and pore parameters of adsorbents for Xq and Yt

Sample	$S_{\text{BET}} (\text{m}^2 \text{ g}^{-1})$	$S_{\text{micr}} (\text{m}^2 \text{ g}^{-1})$	$V_{\text{total}} (\text{m}^3 \text{ g}^{-1})$	$D_{\text{average}} (\text{nm})$
Yt	42.6	1.8	0.122	13.5
Xq	35.3	1.6	0.115	14.8

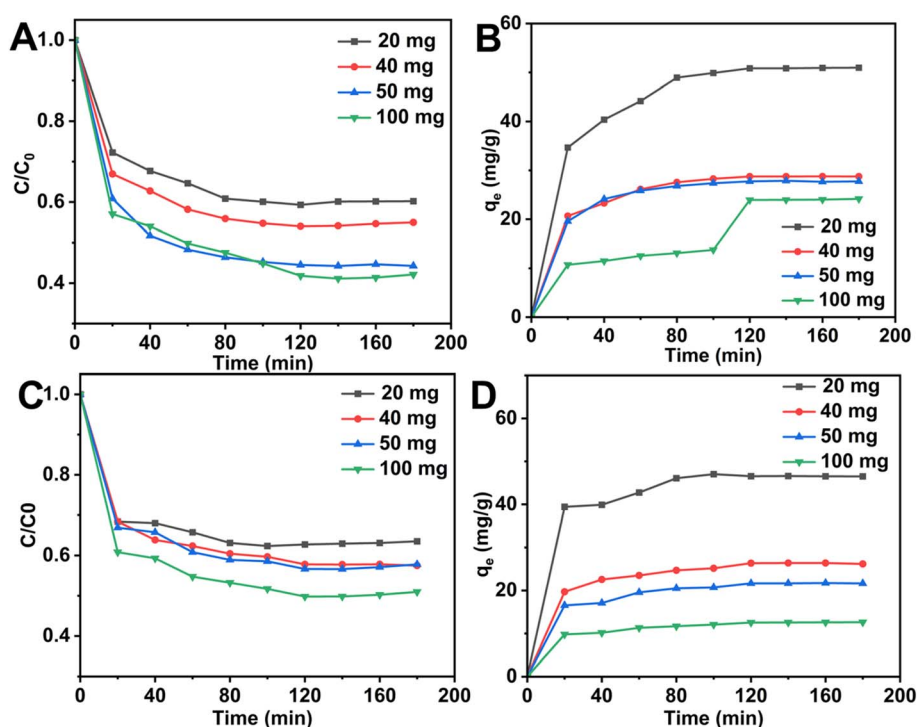


Fig. 6 The fluoride removal rate and adsorption capacity of Xq-HAP (A and B) and Yt-HAP (C and D) varied with the dosage.



Table 2 Comparison of adsorption capacities of different materials

Sample	Adsorption capacity (mg g <sup>-1</sup> )	Adsorption condition	References
Carbonaceous materials	0.8–3.5	20 mg L <sup>-1</sup> , 20 mL, 200 mg, shake, 3 h	30
Chitosan beads	1.8	10 mg L <sup>-1</sup> , 50 mL, 250 mg, shake, 1 h	31
Shell@fluorapatite nanoarray	15	10 mg L <sup>-1</sup> , 50 mL, 10 mg, shake, 20 min	32
CeO <sub>2</sub> @SiO <sub>2</sub> microspheres	32.24	50 mg L <sup>-1</sup> , 100 mL, 150 mg, shake, 2 h	33
Highly activated alumina media	7.47	10 mg L <sup>-1</sup> , 10 mL, 10 mg, shake, 24 h	34
Xq-HAP	29.04	10 mg L <sup>-1</sup> , 50 mL, 50 mg, shake, 2 h	This work

g<sup>-1</sup>. The fluorine removal rate was 43.38%, and the adsorption capacity reached 21.75 mg g<sup>-1</sup> at a Yt-HAP dosage of 50 mg. Considering the high efficiency, low cost, and other factors, the optimum dosage of the Xq-HAP adsorbent material was selected as 50 mg for a solution of fluoride ions of 10 mg L<sup>-1</sup>.

**3.2.3 Comparison of the adsorption effect of different materials.** As shown in Table 2, Xq-HAP can be compared with other adsorption materials. It can be seen that Xq-HAP has outstanding adsorption capacity, is simple to prepare, and the cost of the preparation material is low and cost-effective. Compared with previous work, Xq-HAP is more favorable for industrial production. In the application of water treatment fluoride removers, hydroxyapatite not only can effectively control the precipitation process, but also does not release secondary pollutants, showing good environmental performance.

**3.2.4 Effect of adsorption kinetics.** The adsorption kinetic curves for Xq-HAP and Yt-HAP were obtained by measuring the

amount of fluoride ions adsorbed by the two materials at different time points. The experiments revealed the adsorption behavior of these two materials over time. The proposed primary kinetics, secondary kinetics, and intraparticle diffusion modeling equations are as follows:

$$\ln(q_e - q_t) = \ln q_e - k_1 t$$

$$\frac{t}{q_t} = \frac{1}{k_2 q_e} + t/q_e$$

$$q_t = k_3 t^{0.5} + c$$

where  $q_t$  (mg g<sup>-1</sup>) is the adsorption capacity of fluoride ions at any moment  $t$ ,  $q_e$  (mg g<sup>-1</sup>) is the adsorption capacity of fluoride ions when equilibrium is reached;  $k_1$  (min<sup>-1</sup>) is the proposed

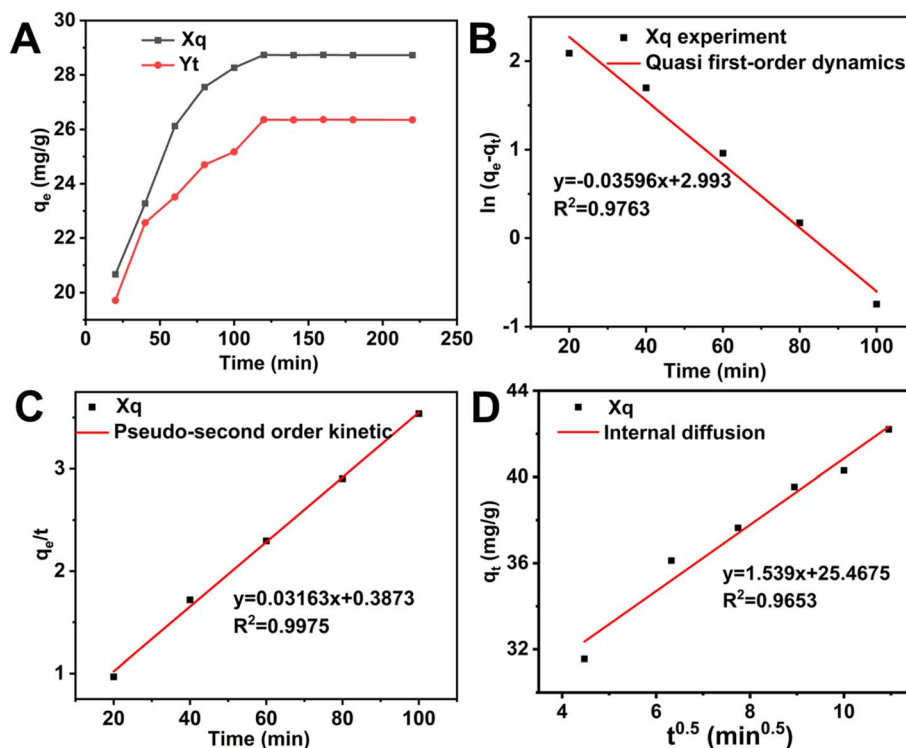


Fig. 7 (A) Adsorption capacity curve with time, (B) the quasi first-order dynamics, (C) the pseudo-second order kinetic, and (D) internal diffusion model.



primary reaction rate constant;  $k_2$  ( $\text{g} (\text{mg}^{-1} \text{min}^{-1})$ ) is the proposed secondary reaction rate constant, and  $k_3$  ( $\text{mg g}^{-1} \text{min}^{-0.5}$ ) is the intraparticle diffusion rate.

As shown in Fig. 7(A), during the first 60 min, the adsorbent material had sufficient and available active sites, the adsorption drive was high, and the adsorption rate was high. The adsorption rate slowed slightly from 60 to 120 min, which is attributed to the diffusion of fluoride ions on the surface of the adsorbent into the pores, resulting in a large number of active sites exposed on the adsorbent surface being occupied by fluoride. After 180 min, the adsorption capacity gradually reached equilibrium and the maximum adsorption capacity reached  $28.74 \text{ mg g}^{-1}$ . As shown in Fig. 7B, C and Table 3, Xq-HAP was more in line with the model of the proposed second-order dynamics, where the model correlation coefficient of the proposed second-order dynamics ( $R^2$ ) was 0.9975. The correlation coefficients were all higher than the corresponding value of 0.9763 for the proposed one-level kinetic model, and it can be inferred that the adsorption of fluorine by Xq-HAP is mainly a chemisorption effect. As shown in Fig. 7(D), the effect of the internal diffusion model fitting indicates that adsorption on the adsorbent surface of Xq-HAP and the diffusion process into the pores jointly affect the adsorption efficiency during the

adsorption process. This finding agrees with the results reported in numerous studies.

**3.2.5 Effect of initial concentration.** In order to investigate the impact of two adsorption materials on the elimination of fluoride ions, the study specifically examined the influence of the initial concentration of fluoride ions. In the experiment, the fluoride ion solution was 50 mL, the initial concentration of the fluoride ion solution was  $5 \text{ mg L}^{-1}$ ,  $10 \text{ mg L}^{-1}$ , and  $20 \text{ mg L}^{-1}$ , and the dosage of adsorbent material was 50 mg. The experiment was conducted at  $25^\circ\text{C}$ , and adsorption equilibrium was reached in 180 min. As shown in Fig. 8, while the initial concentration of the fluoride ion solution varied from small to large, the removal rate of fluoride ions by Xq-HAP and Yt-HAP decreased. The fluoride ion removal by Xq at  $5 \text{ mg L}^{-1}$ ,  $10 \text{ mg L}^{-1}$ , and  $20 \text{ mg L}^{-1}$  was 52%, 46%, and 22%, respectively. The adsorption capacities were  $14.41 \text{ mg g}^{-1}$ ,  $28.78 \text{ mg g}^{-1}$ , and  $19.18 \text{ mg g}^{-1}$ , respectively, which was attributed to the insufficient adsorption drive at low fluoride ion concentration. The higher the fluoride ion concentration, the less effective the sites for adsorption of Xq, resulting in a decreased adsorption capacity. The fluoride ion removal rates by Yt-HAP were 42%, 42%, and 18%, and the adsorption capacities were  $11.86 \text{ mg g}^{-1}$ ,  $26.41 \text{ mg g}^{-1}$ , and  $16.24 \text{ mg g}^{-1}$ , respectively. These were taken together, so the optimum initial concentration of the fluoride ion solution was selected as  $10 \text{ mg L}^{-1}$ .

**3.2.6 Effect of adsorption temperature.** In order to investigate the effect of two adsorbent materials on the removal of fluoride ions at different temperatures, the concentration of fluoride ions in the experiment was  $10 \text{ mg L}^{-1}$ , each conical flask was taken as 50 mL, and the dosage of adsorbent materials was 50 mg. The experimental temperatures were controlled to

Table 3 Adsorption kinetic parameters of the pseudo-first-order and pseudo-second-order kinetic models

Adsorbent	Quasi first-order dynamics			Pseudo-second order kinetic		
	$K_1$	$Q_e (\text{mg g}^{-1})$	$R^2$	$K_2$	$Q_e (\text{mg g}^{-1})$	$R^2$
Xq	0.036	19.95	0.9763	0.0316	31.65	0.9975

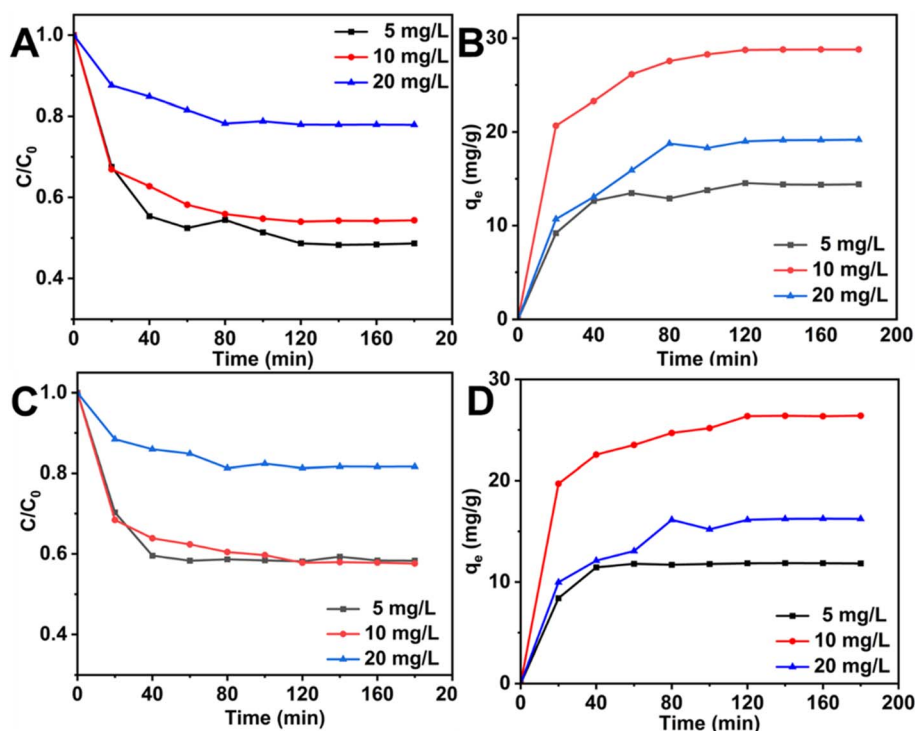


Fig. 8 The fluoride removal rate and adsorption capacity of Xq-HAP (A and B) and Yt-HAP (C and D) vary with initial concentration.





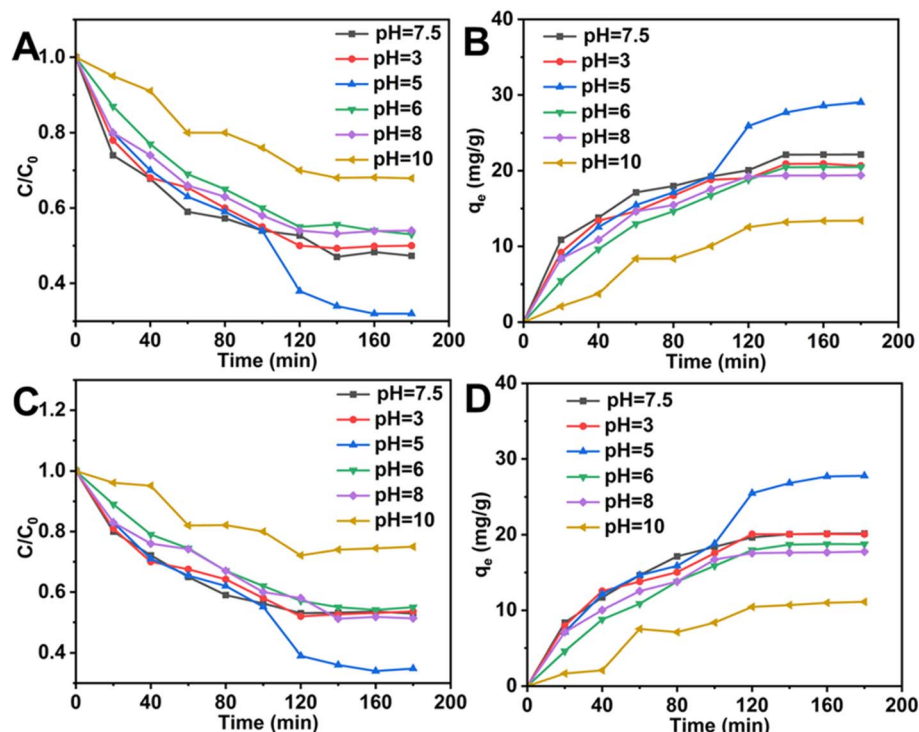


Fig. 9 The fluoride removal rate and adsorption capacity of Xq-HAP (A and B) and Yt-HAP (C and D) vary with pH.

be 25 °C, 35 °C and 45 °C, and the adsorption equilibrium was reached in 180 min. Fig. S3(A)† shows a graph of the fluoride ion adsorption efficiency of Xq-HAP over time. Fig. S3(B)† shows the graph of the adsorption capacity over time. Fig. S3(C)† shows the fluoride ion adsorption efficiency of Yt-HAP over time. Fig. S3(D)† represents the relationship between adsorption capacity and time. As the temperature of the reaction system increased from 25 °C to 35 °C and then to 45 °C, the adsorption efficiency of Xq-HAP for fluoride ions increased from the initial 34% to 45% and then to 52%, and the adsorption capacity also increased from 21.13 mg g<sup>-1</sup> to 28.86 mg g<sup>-1</sup> and then to 32.64 mg g<sup>-1</sup>. The adsorption efficiency of Yt-HAP for fluoride ions increased from the initial 29% to 42% to 46%, and the adsorption capacity also increased from 18.40 mg g<sup>-1</sup> to 26.43 mg g<sup>-1</sup> to 29.12 mg g<sup>-1</sup>. It can be seen that the adsorption reaction is a heat-absorbing reaction. Increasing the temperature during the experiment is favorable for the adsorption reaction, which indicates that Xq is heat-resistant and can adapt to harsh environments. The final temperature of the reaction was maintained at 35 °C.

**3.2.7 Effect of initial pH.** To determine the effect of the two adsorption materials on the efficiency of fluoride ion removal under different initial pH conditions, 50 mL of a fluoride ion solution with an initial concentration of 10 mg L<sup>-1</sup> was used in the experiment. The initial pH values of the fluoride solutions were 3, 5, 6, 8, 10, and 7.5. The experiment was conducted at 35 °C, and the adsorption process reached the adsorption equilibrium at 180 min. As shown in Fig. 9(A–D), Xq-HAP was the most effective in removing fluoride ions when the pH was 5. Xq-HAP removed 68% of the fluoride ions, and the adsorption capacity

was 29.04 mg g<sup>-1</sup>. It can be seen that the adsorption of fluoride ions is very poor under the condition that the initial solution pH value is strong acid or strong basic. A possible reason is that in an acidic solution, the solution contains a large number of hydrogen ions, which produces a weak electrolyte hydrogen fluoride with fluoride ions, thus affecting the adsorption effects and making the removal of fluoride ions less effective. When the solution becomes alkaline, the higher concentration of OH<sup>-</sup> ions causes competition with F<sup>-</sup> ions for adsorption. As a result, the materials' ability to adsorb fluoride ions is greatly diminished, leading to a drop in the effectiveness of fluoride ion removal.

**3.2.8 Adsorption isotherm.** The relationship between the fluoride concentration and adsorption capacity was investigated by fitting different adsorption isotherm models.

Langmuir adsorption isotherms:

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

Freundlich adsorption isotherm:

$$\ln q_e = \ln k_F + \frac{\ln c_e}{n}$$

Temkin adsorption isotherm:

$$q_e = b \ln a + b \ln c_e$$

where  $q_e$  denotes the mass of fluoride ions adsorbed per unit mass of adsorbent at adsorption equilibrium in mg g<sup>-1</sup>,  $c_e$





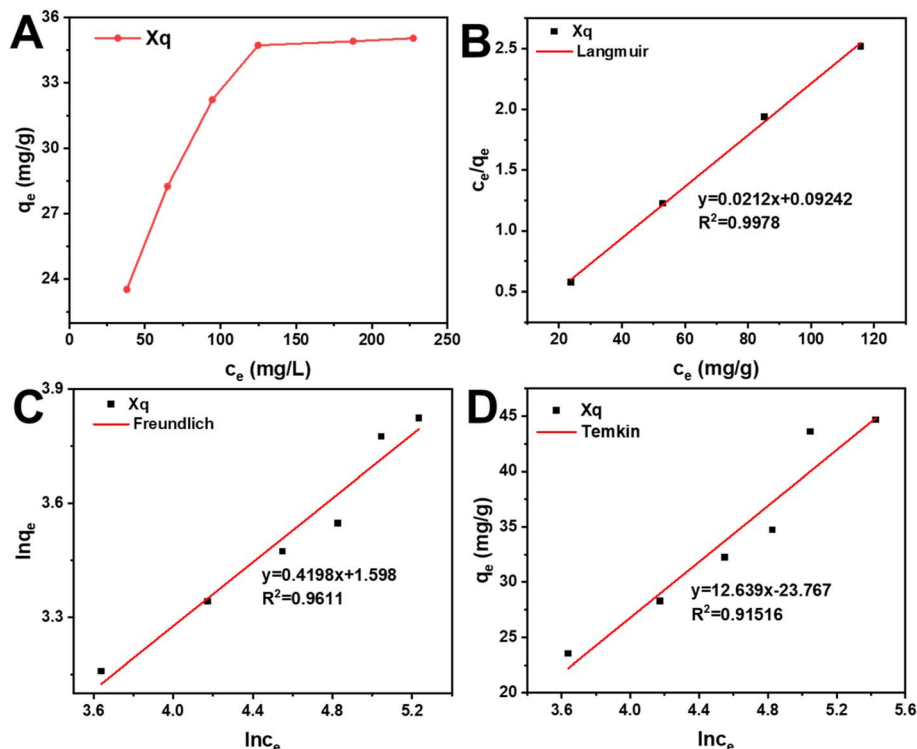


Fig. 10 (A) Adsorption isotherms of fluoride on Xq-HAP samples (dosage = 1 g L<sup>-1</sup>, *T* = 25 °C, contact time = 24 h, and pH = 7), (B) Langmuir isotherm model, (C) Freundlich isotherm model, and (D) Temkin isotherm model.

Table 4 Langmuir and Freundlich isotherm model parameters for fluoride adsorption

Adsorbent	Langmuir model			Freundlich model		
	$q_{\max}$ (mg g <sup>-1</sup> )	$K_L$ (L mg <sup>-1</sup> )	$R^2$	$K_F$ (mg <sup>1-n</sup> L <sup>n</sup> g <sup>-1</sup> )	$1/n$	$R^2$
Xq-HAP	47.16	0.2294	0.9978	0.133	0.331	0.997

denotes the equilibrium concentration of fluoride ions in mg L<sup>-1</sup>,  $q_m$  denotes the maximum mass of fluoride ions adsorbed per unit mass of adsorbent in mg g<sup>-1</sup>,  $k_L$  denotes the Langmuir constant associated with the adsorption capacity in L mg<sup>-1</sup>,  $k_F$  denotes the Freundlich constant associated with the adsorption capacity of the adsorbent, and  $1/n$  is the adsorption strength.

As shown in Fig. 10 and Table 4, the Langmuir model fit of Xq-HAP had correlation coefficients as high as 0.9978, and the maximum adsorption capacity was calculated to be 47.16 mg g<sup>-1</sup> according to the theoretical. The Langmuir model is closer to the actual adsorption process than the Freundlich model, which has a correlation coefficient of 0.9611, and  $1/n < 1$  is considered as a measure of favorable or unfavorable adsorption in the Freundlich model. Therefore, it can be concluded that the Xq-HAP material exhibits significant advantages in fluoride adsorption, and its adsorption process exhibits homogeneous surface properties. Careful analysis revealed that the Xq-HAP material exhibited monolayer adsorption when fluoride was adsorbed. The Temkin adsorption isotherm model was used to study the effects of high fluoride concentrations, and the

Xq-HAP correlation coefficient was 0.9152, suggesting that the vast majority of the process was chemisorption, with electrostatic adsorption in the physical adsorption serving as a supporting effect. Due to their comparable ionic radii, F<sup>-</sup> can

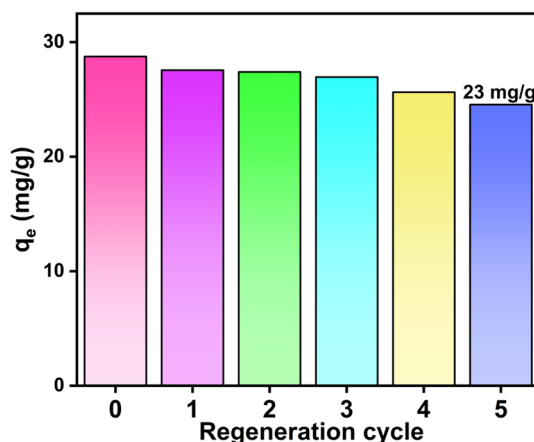


Fig. 11 Recyclability studies of Xq-HAP.



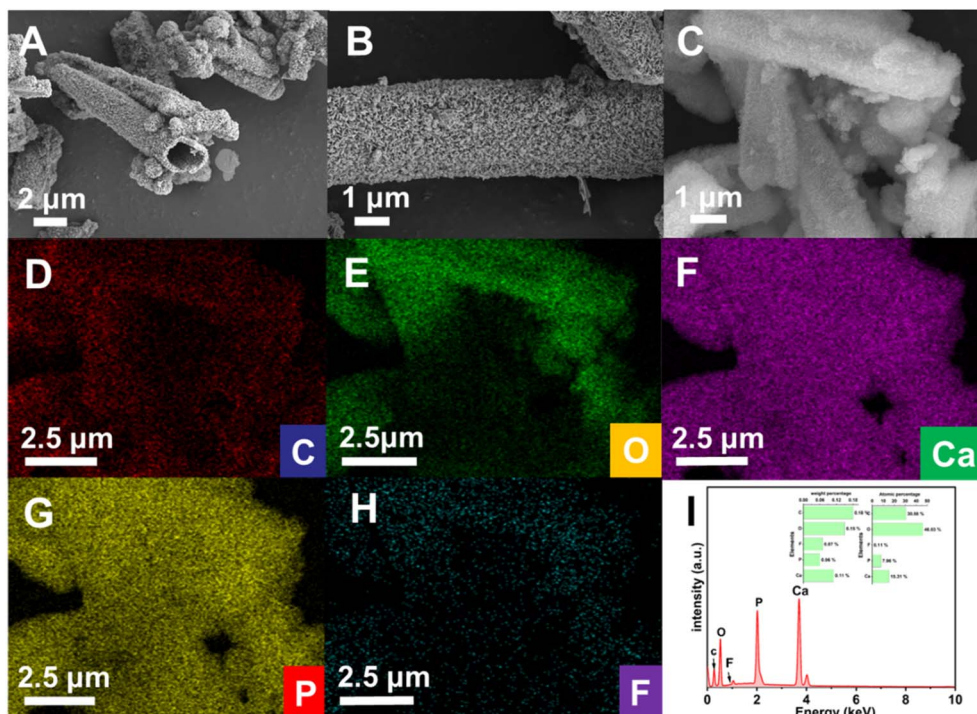


Fig. 12 (A–C) SEM images, (D–H) EDS elemental mapping for C, O, Ca, P and F respectively, (I) EDS spectrum of Xq-HAP after adsorption.

engage in ion exchange with  $\text{OH}^-$  and  $\text{Ca}^{2+}$ . This results in the formation of a complex fluorine-based apatite.

**3.2.9 Desorption experiments.** The recycling of the material and the regeneration ability of the adsorbent are the key to

judge its performance. The  $0.5 \text{ mol L}^{-1}$  NaOH and  $5 \text{ mol L}^{-1}$   $\text{NaCO}_3$  solutions were selected as desorbents. The material was rinsed with deionized water, centrifuged, and then dried. The desorption-treated materials were then subjected to repeated

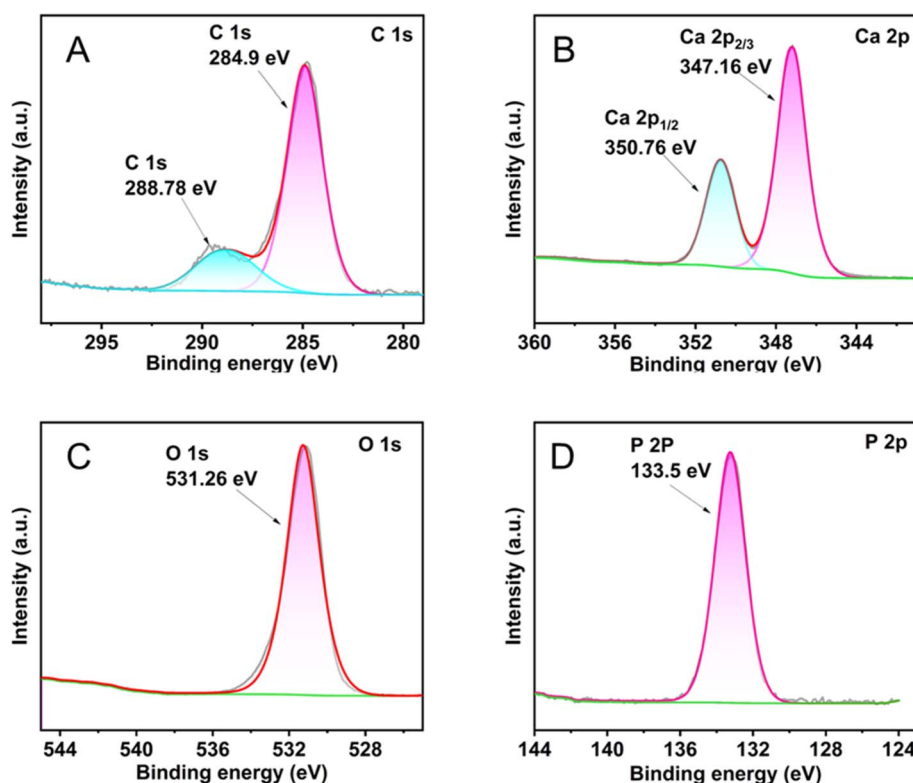


Fig. 13 XPS spectra of the Xq-HAP after fluoride adsorption: (A) C 1s, (B) Ca 2p, (C) O 1s, and (D) P 2p.



adsorption experiments. The effect of adsorption is shown in Fig. 11. The results showed that the adsorption performance of the adsorbent decreased slightly after five cycles but still maintained a high adsorption capacity of 23 mg g<sup>-1</sup> for fluoride ions.

### 3.3 Stability of materials

Fig. 12(A–C) reveals that the material maintained its rod-like shape even after adsorption, suggesting a high level of stability. Fig. 12(D–H) indicates the presence of elemental fluorine, together with Ca, P, and O elements, confirming the successful capture of fluoride ions.

X-ray photoelectron spectroscopy (XPS) analysis can be used to assess the chemical valence changes of the materials before and after the reaction with fluoride ions following Xq-HAP adsorption. The existence of C 1s, O 1s, Ca 2p, and P 2p peaks in Fig. 13(A–D) shows the presence of Xq-HAP including C, O, Ca, and P components. A comparison of the spectra of Xq-HAP after the adsorption of fluoride ions with those before adsorption revealed some changes in the peaks, confirming the successful adsorption of fluoride ions by Xq-HAP. In X combined with the spectrum before adsorption, the analysis shows that the Ca 2p spectrum can be decomposed into the spectra of two pairs of spin-orbitals, the Ca 2p<sub>3/2</sub> located at 347.39 eV and the located at 350.98 eV of Ca 2p<sub>1/2</sub>. The peak area of Ca decreased dramatically after adsorption of fluoride ions by Xq-HAP. The percentage of Ca<sup>2+</sup> decreased from 16.12% to 15.27%, which was hypothesized that it might be due to the fact that Ca<sup>2+</sup> was released into the solution or replaced by fluoride ions. Whereas, in combination with the O 1s exhibited before adsorption, it can be seen that the O 1s is located near the peak at 531.38 eV belonging to the OH<sup>-</sup> of Xq-HAP before the adsorption reaction. After the adsorption reaction, the oxygen fraction peak of adsorbed water was observed at 531.26 eV. The material maintained an extremely high stability after the adsorption of fluoride ions, and the doping of fluoride ions did not change its surface chemical state, which provided a strong guarantee for its recycling performance. Essentially, the primary processes responsible for the attachment of fluoride ions to Xq-HAP involve electrostatic interactions and surface complexation. Moreover, the resulting surface complexes formed following the attachment exhibit remarkable stability.

## 4. Conclusion

The study involved the preparation of Xq-HAP adsorbent material using discarded mussel shells from the coastal area of Lianyungang. This material was used to remove fluoride ions from aqueous solution. Compared to hydroxyapatite (Yt-HAP) prepared using traditional methods, Xq-HAP demonstrated a higher adsorption capacity and removal rate. XRD analysis showed that Xq-HAP had a well-defined hexagonal structure and cubic phase. The fluoride removal effect of Xq-HAP as an adsorbent was investigated, and the results showed that the optimum pH for the adsorption of fluoride ions by Xq-HAP was 5, and equilibrium was reached after 180 min of contact. The

kinetic analysis also showed that the adsorption efficiency of Xq-HAP was higher than that of Yt-HAP, and the proposed second-order equation was the best way to describe the adsorption of fluoride ions by Xq-HAP, suggesting that adsorption is a surface phenomenon as well as a rate-limiting process. SEM revealed that Xq-HAP had a rod-like structure and there was no substantial difference in the morphology of the material before and after adsorption. Overall, the adsorbent has excellent potential for applications in fluoride removal.

## Data availability

The authors confirm that the data supporting the findings of this study are available within the article.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by Lianyungang Postdoctoral Research Foundation (LYG20230017) the Research Program of “521 Project” of Lianyungang City of China.

## References

- 1 D. B. Adamu, L. T. Tufa, J. Lee, E. Zereffa, T. A. Segne and M. H. Razali, *J. Environ. Chem. Eng.*, 2023, **11**, 111196.
- 2 A. Angelin, M. Kalpana, K. Govindan and S. Kavitha, *Environ. Sci. Pollut. Res.*, 2021, **30**, 71614–71627.
- 3 V. E. Badillo-Almaraz, J. Armando Flores, H. Arriola, F. A. López and L. Ruiz-Ramirez, *J. Radioanal. Nucl. Chem.*, 2007, **271**, 741–744.
- 4 L. Cheng, Z. Guan, L. Si, C. Mang, X. Weng, Q. Zhang and Z. Ma, *J. Water Reuse Desalin.*, 2018, **8**, 479–489.
- 5 G. Choudhary, M. Yadav, B. Saini, P. R. Sharma, S. Pandey, V. Kant and R. K. Sharma, *Desalination*, 2024, **574**, 133355.
- 6 L. L. Dai, F. Nudelman, C. H. Chu, E. C. M. Lo and M. L. Mei, *J. Dent.*, 2021, **105**, 103581.
- 7 M. Gao, W. Wang, H. Yang and B.-C. Ye, *Microporous Mesoporous Mater.*, 2019, **289**, 109620.
- 8 J. J. García-Sánchez, V. Martínez-Miranda and M. Solache-Ríos, *J. Fluorine Chem.*, 2013, **145**, 136–140.
- 9 J. J. García-Sánchez, M. Solache-Ríos, J. M. Martínez-Gutiérrez, N. V. Arteaga-Larios, M. C. Ojeda-Escamilla and I. Rodríguez-Torres, *J. Fluorine Chem.*, 2016, **186**, 115–124.
- 10 A. Jeyaseelan, I. Aswin Kumar, M. Naushad and N. Viswanathan, *J. Mol. Liq.*, 2022, **354**, 118830.
- 11 A. Jeyaseelan, N. Viswanathan, I. A. Kumar and M. Naushad, *J. Mol. Liq.*, 2023, **383**, 122125.
- 12 M. Jiménez-Reyes and M. Solache-Ríos, *J. Hazard. Mater.*, 2010, **180**, 297–302.
- 13 M. Jiménez-Reyes and M. Solache-Ríos, *Water, Air, Soil Pollut.*, 2013, **224**, 1499.
- 14 P. Liang, R. An, R. Li and D. Wang, *Int. J. Biol. Macromol.*, 2018, **111**, 255–263.



- 15 W. Liang, L. Zhan, L. Piao and C. Rüssel, *Mater. Res. Bull.*, 2011, **46**, 205–209.
- 16 X.-R. Ma, X.-Y. Wei, R. Dang, W. Guo, Y.-H. Kang, X. Li, Y. Gao, J.-J. Bai, Y. Zhang, Z.-F. Zhang, Y.-J. Ma and Z.-M. Zong, *Appl. Clay Sci.*, 2021, **211**, 106191.
- 17 E. E. Merodio-Morales, H. E. Reynel-Ávila, D. I. Mendoza-Castillo, C. J. Duran-Valle and A. Bonilla-Petriciolet, *Int. J. Environ. Sci. Technol.*, 2019, **17**, 115–128.
- 18 S. Mohan, D. K. Singh, V. Kumar and S. H. Hasan, *J. Fluorine Chem.*, 2017, **194**, 40–50.
- 19 N. Sangiorgi, R. Bondoni, A. Sangiorgi, L. Aversa, R. Tatti, R. Verucchi, A. Adamiano, M. Sandri, A. Tampieri and A. Sanson, *Ceram. Int.*, 2021, **47**, 9701–9710.
- 20 V. Russo, A. D'Angelo, C. Salvi, R. Paparo, M. E. Fortunato, E. M. Cepollaro, O. Tarallo, M. Trifuoggi, M. Di Serio and R. Tesser, *J. Environ. Chem. Eng.*, 2024, **12**, 111973.
- 21 X. Song, F. Zhou, H. Ma, Y. Liu and G. Wu, *Mol. Catal.*, 2023, **542**, 113105.
- 22 B. Zhang, J. Fang, J. Li, J. J. Lau, D. Mattia, Z. Zhong, J. Xie and N. Yan, *Chem.-Asian J.*, 2016, **11**, 532–539.
- 23 K. Zhang, Y. Shao, Y. Jiang, L. Zhang, S. Zhang, Y. Wang, S. Hu, J. Xiang and X. Hu, *Mol. Catal.*, 2024, **555**, 113887.
- 24 S. Samatya, H. Mizuki, Y. Ito, H. Kawakita and K. Uezu, *React. Funct. Polym.*, 2010, **70**, 63–68.
- 25 T. Sani, M. Adem, G. Fetter, P. Bosch and I. Diaz, *Water, Air, Soil Pollut.*, 2016, **227**, 90.
- 26 M. Yi, K. Wang, H. Wei, D. Wei, X. Wei, B. Wei, L. Shao, T. Fujita and X. Cui, *J. Hazard. Mater.*, 2023, **442**, 130027.
- 27 S. Zhang, Y. Lyu, X. Su, Y. Bian, B. Yu and Y. Zhang, *Environ. Earth Sci.*, 2016, **75**, 401.
- 28 X. Zhang, Y. Qi, Z. Chen, N. Song, X. Li, D. Ren and S. Zhang, *Appl. Surf. Sci.*, 2021, **543**, 148727.
- 29 M. Zhou, H. Yang, Z. Wang, J. Ren, R. Wang and Y. He, *Environ. Sci. Pollut. Res.*, 2022, **30**, 32641–32654.
- 30 I. Abe, S. Iwasaki, T. Tokimoto, N. Kawasaki, T. Nakamura and S. Tanada, *J. Colloid Interface Sci.*, 2004, **275**, 35–39.
- 31 N. Viswanathan, C. Sairam Sundaram and S. Meenakshi, *J. Hazard. Mater.*, 2009, **167**, 325–331.
- 32 Y. Xia, X. Huang, W. Li, Y. Zhang and Z. Li, *J. Hazard. Mater.*, 2019, **361**, 321–328.
- 33 F. Wang, K. Wang, Y. Muhammad, Y. Wei, L. Shao and X. Wang, *ACS Sustain. Chem. Eng.*, 2019, **7**, 14716–14726.
- 34 A. Z. Haddad, C. D. Pilgrim, A. M. Sawvel, J. N. Hohman and A. J. Gadgil, *Adv. Sustainable Syst.*, 2019, **3**, 1900005.

