# **RSC Advances**



PAPER

View Article Online

View Journal | View Issue



Cite this: RSC Adv., 2018, 8, 21644

# Preparation of MoS<sub>2</sub>-based polydopaminemodified core—shell nanocomposites with elevated adsorption performances

New molybdenum disulfide (MoS<sub>2</sub>)-based core-shell nanocomposite materials were successfully prepared through the self-assembly of mussel-inspired chemistry. Characterization by Fourier transform infrared, thermogravimetric analysis, scanning electron microscope and transmission electron microscopy revealed that the surface of the flaked MoS<sub>2</sub> was homogeneously coated with a thin layer of polydopamine (PDA). Dye adsorption performances of the synthesized MoS<sub>2</sub>-PDA nanocomposites were investigated at different pH values and reaction times. Compared with pure MoS<sub>2</sub> nanosheets, the obtained core-shell nanocomposites showed elevated adsorption performances and high stability, indicating their potential applications in wastewater treatment and composite materials.

Received 6th April 2018 Accepted 4th June 2018

DOI: 10.1039/c8ra02964d

rsc.li/rsc-advances

## Introduction

Over the past several decades, graphene has been attracting a great deal of attention due to the unusual properties associated with its ultrathin structure.1-7 Among other twodimensional layer materials,8-11 molybdenum disulfide (MoS2) is one of the most attractive and has a similar structure to graphite. 12-16 MoS2 has a sandwich structure consisting of two layers of sulfur atoms and a middle layer of molybdenum atoms, which are bonded by van der Waals forces. 17-20 MoS2 nanosheets have many excellent physical and chemical properties, and demonstrate important applications in sensors,21-24 optoelectronic devices, 25,26 catalysis 27,28 and other fields.29 In addition, MoS2 nanocomposites have attracted wide study in the field of polymer nanocomposites. 30-33 For example, Li et al. investigated polyethylene glycol (PEG)-modified MoS2 surfaces,34 and the obtained products showed good removal efficiency toward some dyes.

In recent years, mussel-inspired chemistry has become a hot research topic in materials science, chemistry and other fields.<sup>35-37</sup> In the marine environment, mussels can secrete proteins through their feet, which have excellent adhesion and good biocompatibility. Dopamine, as an imitation mussel-protein material, has strong adhesion properties through

a complex self-assembly process to form a polydopamine (PDA)

coating with various functions.<sup>38</sup> The formed PDA layer can be used to modify the surface of inorganic and organic materials,

so it demonstrates wide prospects for application in the fields of

separation membranes, adsorbent materials, biomedical materials, biological binders and so on.<sup>39-42</sup> Now core-shell

materials have shown great potential applications in biology, electricity, catalysis and so on.<sup>43–46</sup> For example, Zhao *et al.* 

synthesized core-shell diamond-based nanocomposites,

exhibiting high activity and high catalytic performance.47

Although the catalytic performance of prepared composite materials may seem ideal, there are still some deficiencies, such

as high cost and harsh operating conditions. Thus, musselinspired chemistry has the advantages of mild reaction condi-

# 2. Materials and methods

### 2.1 Materials

Molybdenum disulfide (MoS<sub>2</sub>, 99.9%) nanosheets were obtained from Huajing Powdery Material Science and

Safranine T, and they showed enhanced adsorption ability.

tions, a wide range of applications and diverse functions, *etc.* On the other hand, core-shell nanocomposites have unique structural characteristics, integrating the properties of both internal and external materials.

In this work, we used PDA to modify the surface of MoS<sub>2</sub> nanosheets to synthesize core-shell nanocomposites. The asprepared composites were characterized by a series of morphological and spectral characterization techniques. The results showed that we had successfully prepared MoS<sub>2</sub>-PDA polymeric materials. The obtained MoS<sub>2</sub>-PDA composites were used as adsorbents for the removal of methylene blue and

<sup>&</sup>quot;State Key Laboratory of Metastable Materials Science and Technology, Yanshan University, Qinhuangdao 066004, China. E-mail: tfjiao@ysu.edu.cn

<sup>&</sup>lt;sup>b</sup>Hebei Key Laboratory of Applied Chemistry, School of Environmental and Chemical Engineering, Yanshan University, Qinhuangdao 066004, China. E-mail: zhanglexin@ysu.edu.cn

State Key Laboratory of Biochemical Engineering, Institute of Process Engineering, Chinese Academy of Sciences, Beijing 100190, P. R. China

**RSC Advances** Paper

Technological Co., Ltd. (Hunan, China). Hydroxyphenethylamine hydrochloride (dopamine, 98%) and tris(hydroxymethyl) aminomethane hydrochloride (Tris-HCl, 99%) were purchased from Aladdin Chemicals and Alfa Aesar Chemicals (Shanghai, China). Methylene blue (MB) and Safranine T (ST) were obtained from Tianjin KaiTong Chemical Reagent and Sinopharm Chemical Reagent Co., Ltd. without further purification. Deionized (DI) water was used to prepare aqueous solutions in all experiments.

### 2.2 Fabrication of the MoS<sub>2</sub>-PDA nanocomposites

The synthesis of core-shell MoS2-PDA nanocomposite was undertaken according to the method given in previous literature.48 Briefly, 300 mg of MoS2 nanosheets were first dispersed in 100 mL of Tris buffer (10 mM, pH = 8.5) with sonication for 10 minutes, and then 200 mg of dopamine was added to the above solution with stirring. After stirring at room temperature in the dark for 12 h, 24 h, and 48 h, the obtained composite products were washed several times with water and ethanol, respectively, then centrifuged at 6000 rpm for 10 min. Finally, the products were freeze-dried for 2-3 days for the next experiments.

### 2.3 Adsorption performance test

In order to investigate the dye adsorption performances of MoS<sub>2</sub> nanosheets and MoS2-PDA nanocomposites, we selected MB and ST as model dyes to complete the experiment. 10 mg of MoS<sub>2</sub> nanosheets and 10 mg of the prepared MoS<sub>2</sub>-PDA nanocomposites were added to 100 mL of MB solution (8 mg L<sup>-1</sup>) and ST solution (30 mg L<sup>-1</sup>), respectively. After different adsorption time intervals, the samples were centrifuged and the supernatant analyzed by a UV-Vis spectrophotometer (664 nm for MB; 530 nm for ST). In addition, we also studied the effect of MoS2 and MoS2-PDA on MB adsorption with different initial solution pH values. The pH of the initial solution was adjusted to values of 2-11 by adding diluted aqueous HCl solution or NaOH solution.20 All experiments were carried out at room temperature under dark conditions. The amount of adsorbed dye per unit mass of adsorbent  $q_t$  (mg g<sup>-1</sup>) at time t (min) is calculated from the following formula:

$$q_t = \frac{(C_0 - C_t)V}{m} \tag{1}$$

where  $C_0$  is the initial concentration of the adsorption solution  $(\text{mg L}^{-1})$ ,  $C_t$  is the concentration of the adsorption solution at time  $t \text{ (mg L}^{-1})$ , m is the total MoS<sub>2</sub> sample added (g), and V is the volume of the adsorption solution (L).

## Characterization

The microstructures of the samples were obtained using a fieldemission scanning electron microscope (SEM) (S-4800II, Hitachi, Japan) equipped with an energy dispersive X-ray spectroscope (EDS) and a transmission electron microscope (TEM) (HT7700, Hitachi High-Technologies Corporation, Japan) with an accelerating voltage of 20 kV. Thermogravimetric analysis (TGA) was performed by a NETZSCH STA 409 PC Luxx

simultaneous thermal analyzer (Netzsch Instruments Manufacturing Co, Ltd, Germany). BET measurements (NOVA 4200-P, US) were taken to characterize the specific surface areas and pore diameter distribution. Fourier transform infrared (FTIR) spectra were accomplished on a Thermo Nexus 470 FT-IR spectrometer (KBr disk). X-ray photoelectron spectroscopy (XPS) was performed using a Bragg diffraction setup (SMART LAB, Rigaku, Japan) with an Al Ka X-ray source. The adsorption experiments were monitored using a Shimadzu UV2550 spectrophotometer. All experimental processes were undertaken in a beaker under dark conditions. The synthesized composite materials were completely dewatered using an FD-1C-50 freeze dryer (Beijing Boli Experimental Instruments Co., Ltd., China). All aqueous solutions were prepared with water purified in a double-stage Millipore Milli-Q Plus purification system.

#### 3. Results and discussion

### 3.1 Characterization of nanocomposites

Firstly, Fig. 1 shows the entire experimental process, including the preparation of MoS<sub>2</sub>-PDA and the adsorption experiments of the obtained nanocomposites. First, MoS2-PDA nanocomposites were obtained by modification of MoS2 nanosheets with a PDA layer in Tris buffer solution and dried by centrifugation. Then two model dyes (MB and ST) were used to evaluate the adsorption capacity of the prepared composite materials.

The nanostructures of the obtained materials were investigated by SEM and TEM, as shown in Fig. 2. Obviously, compared with the original MoS2 (Fig. 2a), with an increase in modification time, the layer thickness of PDA coated on the MoS<sub>2</sub> surface increased and showed a tendency to aggregate together, as shown in Fig. 2b-d. In addition, Fig. 2e represents the TEM image of the MoS<sub>2</sub>-PDA composite with a modification time of 48 h, which was named MoS2-PDA-48. In the TEM image, it was obvious that the outer edge of the MoS2 nanosheets demonstrated some ultra-thin layer structures, which indicated that the PDA layer had been successfully anchored on the surface of the MoS<sub>2</sub> nanosheets. Moreover, the typical XRD pattern of MoS<sub>2</sub> (the inset image of Fig. 2a) and the EDX pattern of MoS<sub>2</sub>-PDA-48 (Fig. 2f) also provided obvious characteristics and suggested the above speculations.

In addition, the thermal stability of MoS2 and the obtained MoS2-PDA composites were characterized and are shown in Fig. 3. According to the TG curves, MoS<sub>2</sub> showed a low mass loss

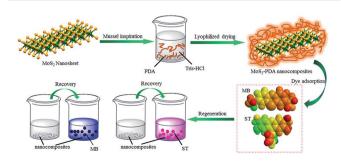


Fig. 1 Schematic illustration of the fabrication of MoS<sub>2</sub>-PDA composites by mussel-inspired chemistry.

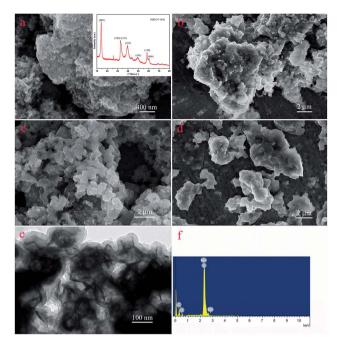


Fig. 2 SEM and TEM images of as-prepared materials: (a) SEM image and inserted XRD data of  $MoS_2$ ; (b–d) SEM images of core–shell  $MoS_2$ –PDA composites after different modification times (12, 24, and 48 h); (e) and (f) TEM image and EDX spectra of  $MoS_2$ –PDA-48.

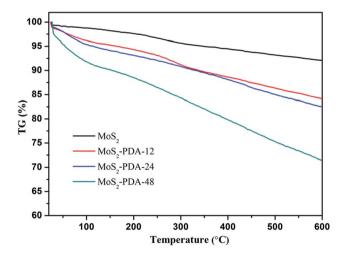


Fig. 3  $\,$  TG curves of MoS $_2$  nanosheets and the obtained MoS $_2$ -PDA nanocomposites.

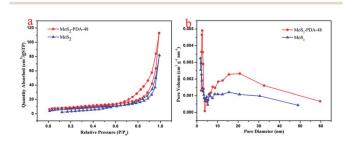


Fig. 4 (a) Adsorption–desorption isotherms and (b) pore size distribution of MoS<sub>2</sub> and the MoS<sub>2</sub>–PDA-48 nanocomposite.

Table 1 Physical data of MoS<sub>2</sub> and the MoS<sub>2</sub>-PDA-48 nanocomposite

Sample	Specific surface area (m² g <sup>-1</sup> )	Average pore diameter (nm)	Pore volume (cm <sup>3</sup> g <sup>-1</sup> )
MoS <sub>2</sub>	25.83	19.62	0.127
MoS <sub>2</sub> -PDA-48	30.57	22.84	0.175

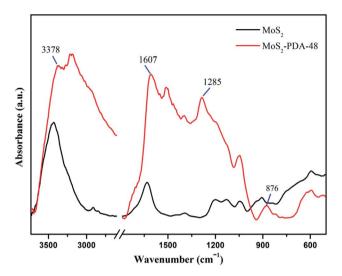


Fig. 5 IR spectra of MoS<sub>2</sub> and the MoS<sub>2</sub>-PDA-48 composite.

(about 7.9%) from room temperature to  $600\,^{\circ}$ C, which indicated good thermal stability. The mass losses of the  $MoS_2$ -PDA composites took the values of 15.8%, 17.4%, and 28.6% with different modification times (12 h, 24 h, and 48 h, respectively). The obtained results indicated that the PDA layer had been successfully coated onto the  $MoS_2$  surface and the amount of PDA component in the composites increased with an increase in modification time.

In order to study the microstructure characteristics of  $MoS_2$  and  $MoS_2$ –PDA-48, Fig. 4a shows the BET measurements using  $N_2$  adsorption–desorption isotherms. The adsorption–desorption isotherms of both  $MoS_2$  and  $MoS_2$ –PDA-48 belonged to type IV isotherms. The hysteresis loop of  $MoS_2$  has an abrupt increase in the amount of adsorption at  $P/P_0 = 0.6$ –0.95, which

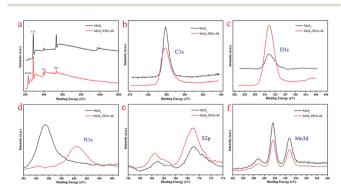


Fig. 6 (a) XPS survey spectra of  $MoS_2$  and  $MoS_2$ -PDA-48; high resolution scans of different elements: (b) C 1s; (c) O 1s; (d) N 1s; (e) S 2p; (f) Mo 3d.

 $\textbf{Table 2}\quad \text{Elemental contents (\%) of MoS}_2 \, \text{and MoS}_2 - \text{PDA} - 48 \, \text{based on XPS analysis}$ 

Samples	Mo [at%]	S [at%]	C [at%]	O [at%]	N [at%]
MoS <sub>2</sub>	13.99	27.71	28.94	26.10	3.26
MoS <sub>2</sub> -PDA-48	3.51	9.11	52.56	27.24	7.58

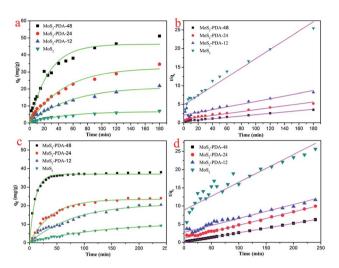


Fig. 7 Adsorption kinetics curves of MB (a and b) and ST (c and d) onto  $MoS_2$  nanosheets and prepared  $MoS_2$ -PDA nanocomposites with different reaction times.

is characteristic of a mesoporous material. For the  $MoS_2$ -PDA-48 nanocomposites, a hysteresis loop could be observed at pressure  $P/P_0=0.8$ -0.95 that indicated a mesoporous structure. The pore size distribution curves were obtained according to the BET method and are shown in Fig. 4b. The pore size distribution of  $MoS_2$ -PDA-48 showed a larger mesoporous size of 22.84 nm, compared to 19.62 nm for  $MoS_2$ . The data for BET surface area and total pore volume are listed in Table 1. The specific surface area of  $MoS_2$ -PDA-48 was calculated to be 30.57

 $\rm m^2~g^{-1}$ , which was much larger than that of MoS<sub>2</sub> (25.83  $\rm m^2~g^{-1}$ ). The higher specific surface area can increase the number of active sites on the surface of the MoS<sub>2</sub>-PDA composite and enhance the chances of dye molecules anchoring at active sites, thus making adsorbents with better adsorption properties.

FT-IR spectra of MoS<sub>2</sub> and the MoS<sub>2</sub>-PDA-48 composite are demonstrated in Fig. 5. The curve of MoS2-PDA-48 shows a characteristic band at 3378 cm<sup>-1</sup>, which could be ascribed to the stretch vibrations of -NH<sub>2</sub>, -NH-, and -OH.<sup>49-55</sup> The characteristic peak at 1607 cm<sup>-1</sup> could be deduced as a C=C stretching vibration in the benzene ring. At the same time, the weak peaks around 1285 and 876 cm<sup>-1</sup> could be assigned to C-OH stretching vibration from the catechol groups and C-O stretching, respectively, which indicate that the surface of MoS<sub>2</sub> was successfully modified by a PDA layer. 56-59 In addition, the XPS analysis (Fig. 6) demonstrates the composition of MoS<sub>2</sub> and MoS<sub>2</sub>-PDA-48. Fig. 6a demonstrates the characteristic peaks in the curve of the MoS<sub>2</sub>-PDA-48 composite, such as Mo (3d), O (1s), C (1s), and N (1s). Additionally, the high resolution XPS spectra are shown in Fig. 6b-f. Compared with pure MoS<sub>2</sub>, the relative intensity of C 1s of MoS2-PDA-48 decreased and the O 1s peak increased significantly (Fig. 6b and c). In the meantime, it was clear that the relative intensity of the N 1s of MoS<sub>2</sub>-PDA-48 at 400 eV increased obviously in Fig. 6d. In addition, as shown in Fig. 6e and f, the relative intensity of Mo 3d and S 2p in MoS<sub>2</sub>-PDA-48 was significantly reduced compared with the MoS<sub>2</sub> nanosheets, which also indicates that the PDA layer was successfully coated onto the surface of the MoS2 nanosheets. On the other hand, the relative elemental analysis data based on XPS analysis is shown in Table 2. The atom percentages of Mo and S in the table fell from 13.99% and 27.71% to 3.51% and 9.11%, respectively. At the same time, the atomic percentage of N in MoS<sub>2</sub>-PDA-48 accounted for 7.58%, indicating successful PDA modification on the surface of the MoS<sub>2</sub> nanosheets.

### 3.2 Adsorption performances

In order to study the adsorption properties of the prepared MoS<sub>2</sub>-PDA nanocomposites, two model dyes (MB and ST) were

Table 3 Kinetic parameters of the  $MoS_2$  and  $MoS_2$ -PDA composites for MB and ST adsorption

МВ	Pseudo-first-order model			Pseudo-second-order model		
	$q_{\mathrm{e}}  (\mathrm{mg \ g^{-1}})$	$R^2$	$k_1  (\mathrm{min}^{-1})$	$q_{ m e}~({ m mg~g^{-1}})$	$R^2$	$k_2 ({\rm g \ mg^{-1} \ min^{-1}})$
$MoS_2$	6.7309	0.9772	0.0208	8.0567	0.9340	0.0032
MoS <sub>2</sub> -PDA-12	21.1223	0.9763	0.0181	27.2183	0.8321	$6.46 \times 10^{-4}$
MoS <sub>2</sub> -PDA-24	32.5736	0.9753	0.0212	38.2409	0.9372	$7.33 \times 10^{-4}$
MoS <sub>2</sub> -PDA-48	46.2684	0.9665	0.0389	54.9451	0.9801	$8.84 \times 10^{-4}$
	Pseudo-first-orde	er model		Pseudo-second-o	rder model	
Safranine T	$q_{ m e}~({ m mg~g}^{-1})$	$R^2$	$k_1  (\mathrm{min}^{-1})$	$q_{\rm e}~({ m mg~g}^{-1})$	$R^2$	$k_2 (\mathrm{g \ mg^{-1} \ min^{-1}})$
$MoS_2$	10.9608	0.9865	0.0077	13.1268	0.8101	$6.52\times10^{-4}$
MoS <sub>2</sub> -PDA-12	23.4885	0.9878	0.0251	26.2192	0.9231	$5.18\times10^{-4}$
MoS <sub>2</sub> -PDA-24	21.0367	0.9714	0.0127	27.7239	0.9845	0.0011
MoS <sub>2</sub> -PDA-48	36.9899	0.9929	0.0959	38.610	0.9996	0.0063

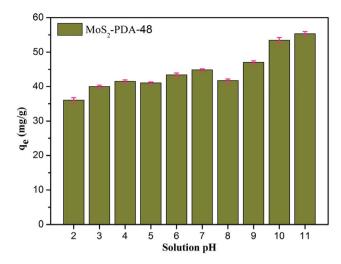


Fig. 8 The effect of solution pH on MB adsorption by  $MoS_2$ -PDA-48 nanocomposites.

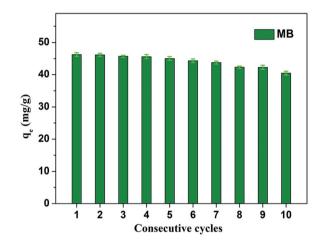


Fig. 9 Relative adsorption capacity of MoS<sub>2</sub>-PDA-48 nanocomposite towards MB for different consecutive cycles.

used in this research work. The concentrations of residual dyes at different reaction time intervals were monitored at different absorbance wavelengths (664 nm for MB; 530 nm for ST). Fig. 7 shows the results of the adsorption kinetics of  $MoS_2$  nanosheets and  $MoS_2$ –PDA nanocomposites for the removal of MB and ST. It can obviously be observed from Fig. 7a and c that the dye adsorption capacities of the  $MoS_2$ –PDA nanocomposites

increased markedly compared to pure MoS<sub>2</sub> in the equilibrium state, and the adsorption effect of MoS<sub>2</sub>-PDA-48 was the best. The detailed kinetic parameters are shown in Table 3. It should be noted that PDA is a kind of hydrophilic and adhesive material. After PDA anchored onto the surface of the MoS<sub>2</sub> nanosheets, the core-shell nanostructures formed demonstrated a porous surface with a number of enhanced active sites to improve the adsorption performance of the composites.

In addition, the adsorption kinetic process can be described by classical kinetic models as follows:

The pseudo-first-order model can be represented by eqn (2):

$$\log(q_{\rm e} - q_{\rm t}) = \log q_{\rm e} - \frac{k}{2.303}t\tag{2}$$

The pseudo-second-order model can be represented by eqn (3):

$$\frac{t}{q_t} = \frac{1}{kq_e^2} + \frac{t}{q_e} \tag{3}$$

where  $q_{\rm e}$  is the equilibrium adsorption capacity, mg g<sup>-1</sup>;  $q_t$  is the adsorption capacity at time t, mg g<sup>-1</sup>; the  $k_1$  and  $k_2$  values are the kinetic rate constants. Kinetic results (Table 3) could be better characterized by a pseudo-first-order model with a correlation coefficient ( $R^2 > 0.9665$ ) or a pseudo-second-order model with a correlation coefficient ( $R^2 > 0.8101$ ). Comparing the above adsorption data, it could easily be observed that MoS<sub>2</sub>–PDA-48 showed the best adsorption capacity, which could be related to the rich functional groups on the surface of PDA. And the ability to remove MB seemed higher than that for ST. In addition, the gradual increase in the maximum adsorption indicated that the experimental method used in this study is feasible.

Fig. 8 shows the effect of solution pH on the adsorption of MB to  $MoS_2$ –PDA-48. It can be seen that as the pH of the solution increased from 2 to 11, the adsorption capacity showed an increasing trend from 36.02 mg g<sup>-1</sup> to 55.32 mg g<sup>-1</sup>. However, this process varied with different pH values. This difference can be presumed to be a result of the protonation of the functional groups on the nanocomposite surface as well as the  $\pi$ – $\pi$  stacking and electrostatic interactions with the MB molecules. In addition, in order to demonstrate the reuse of the obtained composites, 10 consecutive cycles were repeated using the same  $MoS_2$ –PDA-48 nanocomposite and fresh MB solution (Fig. 9). This showed that the adsorption capacity of the  $MoS_2$ –PDA-48 nanocomposite still retained a removal rate of 87.4% towards

 Table 4
 Comparison of MB adsorption capacities in the reported literature

Materials	$Q_{\rm max} \left({\rm mg~g^{-1}}\right)$	Characteristics	Ref.	
Carbon nanotubes	35	Simple process, environmentally friendly	61	
GO/Fe <sub>3</sub> O <sub>4</sub> nanohybrids	32	Poor stability	62	
PDA microspheres	90.7	High-efficiency, large adsorption capacity	63	
e <sub>3</sub> O <sub>4</sub> @MnO <sub>2</sub> core-shell nanocomposite	9.71	Weak adsorption capacity	64	
VDF/PDA membranes	172.3	Good regeneration ability	65	
e <sub>3</sub> O <sub>4</sub> @graphene	45.27	Weak adsorption capacity	66	
$IoS_2$ -PDA	46.3	Eco-friendly preparation, good stability	This w	

**RSC Advances** Paper

MB and reached 40.45 mg  $g^{-1}$ . Moreover, Table 4 lists the MB adsorptions of the relevant materials reported in the literature. 61-66 In contrast, the present prepared MoS<sub>2</sub>-PDA composite materials showed a larger adsorption capacity and eco-friendly preparation process, demonstrating wide application in wastewater treatment and self-assembled core-shell composite materials.

#### 4. Conclusions

In summary, we have synthesized core-shell MoS<sub>2</sub>-PDA nanocomposites simply by using mussel-inspired chemistry. A series of characterization techniques demonstrated that the PDA was successfully coated onto the surface of the MoS2 nanosheets. The prepared MoS<sub>2</sub>-PDA composites showed effective removal capacities towards two model dyes. The adsorption process was illustrated by pseudo-first-order and pseudo-second-order kinetic models. The present research work is expected to show potential applications in wastewater treatment and selfassembled core-shell composite materials.

# Conflicts of interest

There are no conflicts to declare.

# Acknowledgements

We greatly appreciate the financial supports of National Natural Science Foundation of China (No. 21473153 and 51771162), Support Program for the Top Young Talents of Hebei Province, China Postdoctoral Science Foundation (No. 2015M580214), Research Program of the College Science & Technology of Hebei Province (No. ZD2018091), and Scientific and Technological Research and Development Program of Qinhuangdao City (No. 201701B004).

# Notes and references

- 1 A. K. Geim and K. S. Novoselov, Nat. Mater., 2007, 6, 183.
- 2 K. S. Novoselov, A. K. Geim, S. V. Morozov, D. Jiang, M. I. Katsnelson and I. V. Grigorieva, Nature, 2005, 438, 197.
- 3 L. Yan, Y. B. Zheng, F. Zhao, S. Li, X. Gao, B. Xu, P. S. Weiss and Y. Zhao, Chem. Soc. Rev., 2012, 41, 97-114.
- 4 B. K. Kim, eXPRESS Polym. Lett., 2012, 6, 772.
- 5 B. H. Nguyen and V. H. Nguyen, Adv. Nat. Sci.: Nanosci. Nanotechnol., 2016, 7, 023002.
- 6 S. P. Lonkar, Y. S. Deshmukh and A. A. Abdala, Nano Res., 2015, 8, 1039-1074.
- 7 F. Perreault, D. F. A. Fonfeca and M. Elimelech, Chem. Soc. Rev., 2015, 44, 5861-5896.
- 8 J. H. Yu, H. R. Lee, S. S. Hong, D. Kong, H. W. Lee, H. Wang, F. Xiong, S. Wang and Y. Cui, Nano Lett., 2015, 15, 1031-
- 9 V. W. Brar, A. R. Koltonow and J. Huang, ACS Photonics, 2017,
- 10 M. K. Jana and C. N. R. Rao, Philos. Trans. R. Soc., A, 2016, 374, 20150318.

- 11 M. Xu, T. Liang, M. Shi and H. Chen, Chem. Rev., 2013, 113, 3766-3798.
- 12 M. A. Lukowski, A. S. Daniel, F. Meng, A. Forticaux, L. Li and S. Jin, J. Am. Chem. Soc., 2013, 135, 10274-10277.
- 13 K. Kaasbjerg, K. S. Thygesen and K. W. Jacobsen, Phys. Rev. B, 2012, **85**, 115317.
- 14 G. Eda, T. Fujita, H. Yamaguchi, D. Voiry, M. Chen and M. Chhowalla, ACS Nano, 2012, 6, 7311-7317.
- 15 D. Lembke, S. Bertolazzi and A. Kis, Acc. Chem. Res., 2015, 48, 100-110.
- 16 L. Zhou, B. He, Y. Yang and Y. He, RSC Adv., 2014, 4, 32570-
- 17 G. Eda, H. Yamaguchi, D. Voiry, T. Fujita, M. Chen and M. Chhowalla, Nano Lett., 2011, 11, 5111-5116.
- 18 H. Li, O. Zhang, C. C. R. Yap, B. K. Tay, T. H. T. Edwin and A. Olivier, Adv. Funct. Mater., 2012, 22, 1385-1390.
- 19 W. Qiao, S. Yan, X. Song, X. Zhang, X. He and W. Zhong, Appl. Surf. Sci., 2015, 359, 130-136.
- 20 Q. Huang, M. Liu, J. Chen, Q. Wan, J. Tian and L. Huang, Appl. Surf. Sci., 2017, 419, 35-44.
- 21 J. Z. Ou, A. F. Chrimes, Y. Wang, S. Tang, M. S. Strano and K. Kalantarzadeh, Nano Lett., 2014, 14, 857-863.
- 22 F. K. Perkins, A. L. Friedman, E. Cobas, P. M. Campbell, G. G. Jernigan and B. T. Jonker, Nano Lett., 2013, 13, 668-673.
- 23 C. Zhu, Z. Zeng, H. Li, F. Li, C. Fan and H. Zhang, J. Am. Chem. Soc., 2013, 135, 5998-6001.
- 24 B. Radisavljevic, A. Radenovic, J. Brivio, V. Giacometti and A. Kis, Nat. Nanotechnol., 2011, 6, 147.
- 25 A. Splendiani, L. Sun, Y. Zhang, T. Li, J. Kim and C. Y. Chim, Nano Lett., 2010, 10, 1271-1275.
- 26 D. Lembke, S. Bertolazzi and A. Kis, Acc. Chem. Res., 2015, 48, 100-110.
- 27 Z. Zhou, Y. Lin, P. Zhang, E. Ashalley, M. Shafa and H. Li, Mater. Lett., 2014, 131, 122-124.
- 28 X. Man, L. Yu, J. Sun and S. Li, Funct. Mater. Lett., 2016, 9, 1650065.
- 29 K. Maslana, Przem. Chem., 2017, 96, 689-691.
- 30 S. K. Kumar and R. Krishnamoorti, Annu. Rev. Chem. Biomol. Eng., 2010, 1, 37-58.
- 31 R. H. Gonçalves, R. Fiel, M. R. Soares, W. H. Schreiner, C. M. Silva and E. R. Leite, Chemistry, 2015, 21, 15583-15588.
- 32 K. Pramoda, M. Kaur, U. Gupta and C. N. Rao, Dalton Trans., 2016, 45, 13810-13816.
- 33 X. Wang, W. Xing, X. Feng, L. Song and Y. Hu, Polym. Rev., 2017, 57, 440-466.
- 34 X. Qiao, F. Hu, D. Hou and D. Li, Mater. Lett., 2016, 169, 241-
- 35 H. Lee, S. M. Dellatore, W. M. Miller and P. B. Messersmith, Science, 2007, 318, 426-430.
- 36 H. C. Yang, J. Luo, Y. Lv, P. Shen and Z. K. Xu, J. Membr. Sci., 2015, 483, 42-59.
- 37 X. Zhang, M. Liu, Y. Zhang, B. Yang, Y. Ji and L. Feng, RSC *Adv.*, 2012, **2**, 12153–12155.
- 38 H. Lee, N. F. Scherer and P. B. Messersmith, Proc. Natl. Acad. Sci. U. S. A., 2006, 103, 12999-13003.

- 39 M. Liu, G. Zeng, K. Wang, Q. Wan, L. Tao and X. Zhang, Nanoscale, 2016, 8, 16819–16840.
- 40 Y. Liu, K. Ai and L. Lu, Chem. Rev., 2014, 114, 5057-5115.
- 41 D. E. Fullenkamp, J. G. Rivera, Y. K. Gong, K. H. Lau, L. He and R. Varshney, *Biomaterials*, 2012, 3, 3783–3791.
- 42 C. Cheng, S. Nie, S. Li, H. Peng, H. Yang and L. Ma, *J. Mater. Chem. B*, 2013, 1, 265–275.
- 43 D. Hou, H. Tao, X. Zhu and M. Li, *Appl. Surf. Sci.*, 2017, **419**, 580–585.
- 44 M. L. Wu, Y. Y. Li, R. Yue, X. D. Zhang and Y. M. Huang, Sci. Rep., 2017, 7, 42773.
- 45 M. Huang, X. L. Zhao, F. Li, W. Li, B. Zhang and Y. X. Zhang, J. Mater. Chem. A, 2015, 3, 12852–12857.
- 46 R. Guo, T. F. Jiao, R. R. Xing, Y. Chen, W. C. Guo, J. X. Zhou, L. X. Zhang and Q. M. Peng, *Nanomaterials*, 2017, 7, 317.
- 47 X. N. Zhao, T. F. Jiao, R. R. Xing, H. Huang, J. Hu, Y. Qu, J. X. Zhou, L. X. Zhang and Q. M. Peng, RSC Adv., 2017, 7, 49923–49930.
- 48 G. J. Zeng, M. Y. Liu, X. H. Liu, Q. Huang, D. Z. Xu, L. C. Mao, H. Y. Huang, F. J. Deng, X. Y. Zhang and Y. Wei, *Appl. Surf. Sci.*, 2016, 387, 399–405.
- 49 R. Guo, T. Jiao, R. Li, Y. Chen, W. Guo, L. Zhang, J. Zhou, Q. Zhang and Q. Peng, ACS Sustainable Chem. Eng., 2018, 6, 1279–1288.
- 50 J. Zhou, Y. Liu, T. Jiao, R. Xing, Z. Yang, J. Fan, J. Liu, B. Li and Q. Peng, *Colloids Surf.*, *A*, 2018, **538**, 7–13.
- 51 J. Song, R. Xing, T. Jiao, Q. Peng, C. Yuan, H. Möhwald and X. Yan, ACS Appl. Mater. Interfaces, 2018, 10, 2368–2376.
- 52 Y. Liu, C. Hou, T. Jiao, J. Song, X. Zhang, R. Xing, J. Zhou, L. Zhang and Q. Peng, *Nanomaterials*, 2018, 8, 35.

- 53 S. Huo, P. Duan, T. Jiao, Q. Peng and M. Liu, *Angew. Chem.*, *Int. Ed.*, 2017, **56**, 12174–12178.
- 54 J. Zhou, F. Gao, T. Jiao, R. Xing, L. Zhang, Q. Zhang and Q. Peng, *Colloids Surf.*, *A*, 2018, 545, 60–67.
- 55 X. Luo, K. Ma, T. Jiao, R. Xing, L. Zhang, J. Zhou and B. Li, *Nanoscale Res. Lett.*, 2017, **12**, 99.
- 56 K. Li, T. Jiao, R. Xing, G. Zou, J. Zhou, L. Zhang and Q. Peng, Sci. China Mater., 2018, 61, 728–736.
- 57 R. Xing, T. Jiao, Y. Liu, K. Ma, Q. Zou, G. Ma and X. Yan, Polymers, 2016, 8, 181.
- 58 R. Xing, K. Liu, T. Jiao, N. Zhang, K. Ma, R. Zhang, Q. Zou, G. Ma and X. Yan, Adv. Mater., 2016, 28, 3669–3676.
- 59 Y. Liu, K. Ma, T. Jiao, R. Xing, G. Shen and X. Yan, *Sci. Rep.*, 2017, 7, 42978.
- 60 R. R. Xing, W. Wang, T. F. Jiao, K. Ma, Q. R. Zhang, W. Hong, H. Qiu, J. X Zhou, L. X. Zhang and Q. M. Peng, ACS Sustainable Chem. Eng., 2017, 5, 4948–4956.
- 61 Y. J. Yao, F. F. Xu, M. Chen, Z. X. Xu and Z. W. Zhu, *Bioresour. Technol.*, 2010, **101**, 3040–3046.
- 62 T. F. Jiao, Y. Z. Liu, Y. T. Wu, Q. R. Zhang, X. H. Yan, F. M. Gao, A. J. P. Bauer, J. Z. Liu, T. Y. Zeng and B. B. Li, Sci. Rep., 2015, 5, 12451.
- 63 J. Fu, Z. Chen, M. Wang, S. Liu, J. Zhang and J. Zhang, *Chem. Eng. J.*, 2015, **259**, 53–61.
- 64 W. Wang, T. F. Jiao, Q. R. Zhang, X. N. Luo, J. Hu, Y. Chen, Q. M. Peng, X. Yan and B. B. Li, RSC Adv., 2015, 5, 56279– 56285.
- 65 F. Ma, N. Zhang, X. Wei, J. Yang, Y. Wang and Z. Zhou, *J. Mater. Chem. A*, 2017, 5, 14430–14443.
- 66 Y. Yao, S. Miao, S. Liu, L. P. Ma, H. Sun and S. Wang, Chem. Eng. J., 2012, 184, 326–332.