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Self-assembly fabrication, microstructures and antibacterial performance of layer-structured montmorillonite nanocomposites with cationic silica nanoparticles

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This paper reports the synthesis and antibacterial performance of a novel type of layer-structured montmorillonite (MMT) nanocomposite with cationic silica nanoparticles through an intercalating self-assembly method. The quaternary amine-functionalized SiO₂ nanoparticles were first prepared by grafting dimethyloctadecyl [3-(trimethoxysilyl) propyl] ammonium chloride (PQAC) onto the surfaces of silica nanoparticles, and their chemical composition and structure were characterized by Fourier-transform infrared spectroscopy, thermogravimetric analysis and X-ray photoelectron spectroscopy. Then, the reactant PQAC–SiO₂ nanoparticles were intercalated into the interlayers of MMT through electrostatic self-assembly to achieve the layer-structured MMT/PQAC–SiO₂ nanocomposites. The ordered and layered structural characteristics of the resulting nanocomposites were characterized using X-ray diffraction patterns and N₂ adsorption–desorption measurements and then confirmed by scanning electron microscopy and transmission electron microscopy. The optimum weight ratio of MMT/PQAC–SiO₂ nanoparticles was also achieved for the self-assembly fabrication of the nanocomposites on the basis of results mentioned above. The MMT/PQAC–SiO₂ nanoparticles gained an excellent thermal stability due to the intercalation of PQAC–SiO₂ nanocomposites into MMT and their onset degradation temperature was improved by 40 °C compared to that of PQAC–SiO₂ nanoparticles. Most of all, the nanocomposites presented a significant antibacterial effect against *Escherichia coli* and *Staphylococcus aureus* as the model microorganisms of Gram-negative and Gram-positive bacteria, respectively. This type of layered structural nanocomposite is expected to be applied for food packaging and containment of food materials, medical bandages for wound care, removal of environment pollutants like pesticides and phenol, etc.

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1. Introduction

Nanocomposites based on layer-structured silicate matrices and organic/inorganic compounds are currently one of the most interesting topics for the research and development of advanced functional materials and have received a great deal of attention from both scientific societies and industrial communities.¹ Generally, clay as a sort of layer-structured silicate can be broadly classified into two categories: natural and synthetic clays. Natural clays are basically composed of alternating sheets of “SiO₂” and “AlO₆” units with different mass ratios, which include kaolinite, halloysite, rectorite, chrysotile, smectite, vermiculite, pyragonite, mica, and chlorite.^{2,3} Among these natural clays, montmorillonite (MMT) clay is a type of smectite

with a structure of 2 : 1 phyllosilicates and has been attracting extensive interest in fundamental research and commercial development due to its high cationic exchange capacity, swelling capacity, high surface areas and consequential strong adsorption capacities as well as its broad applications in the preparation of nanocomposites.⁴ The natural MMT with exchangeable cations can be transformed into hydrophobic organic MMT by replacing the metal ions with inorganic ones like silica and organic ones like quaternary ammonium salts.^{5,6} Such modified materials have a larger basal spacing between two layers as well as lower surface energy and can be compatible with organic liquids or polymers. Moreover, the change in the surface property influences the use of MMT clay in different areas.⁷

There are numerous successful cases confirming that the hybrids of layered MMT clay and inorganic nanoparticles cannot only effectively enhance the intrinsic properties of inorganic nanoparticles but also improve the catalysis capability, catalysis capability, long-term thermal stability, high

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durability, and regenerability of the resulting nanocomposites.^{8–10} Boutoumi *et al.*¹¹ synthesized a kind of MMT/TiO₂/polythiophene nanocomposites and found that it exhibited an enhanced degradation capability for the dye of rhodamine 6G. Mao *et al.*¹² reported the synthesis of mesostructured aluminosilicate intercalates and their application as an acidic catalyst for catalytic cracking. Ruiz-Hitzky *et al.*¹³ reported the preparation of a new class of hybrid materials based on the Na⁺-MMT/chitosan bionanocomposites and found that such bionanocomposites have an anion-exchange capability suitable to build potentiometric sensors. Meanwhile, some of studies were focused on the further modification of clays by antibiotic constituents. Lin *et al.*¹⁴ reported the immobilization of MMT with silver nanoparticles and found the resulting composites exhibited a mild inflammatory response and an outstanding antibacterial activity. Lavorgna *et al.*¹⁵ developed the active nanocomposite films consisting of layered MMT intercalated with chitosan and silver-nanoparticles through an ion exchange reaction, and they found that the films could release silver ions in a steady and prolonged manner. Although silver nanoparticles have a high antibacterial activity due to their large specific surface area, they can induce toxicity when used alone. This may result in a limitation to their application in broader areas.¹⁶ Zhou *et al.*¹⁷ reported a surface modification method for MMT by use of cationic gemini surfactants and found that these modified materials exhibit potential applications for removal of environment pollutants such as pesticides and for medical fields as antimicrobial materials. The MMT intercalated with quaternary ammonium compounds is well known as 'organo-clays' and has been widely used as disinfectants and antiseptics in food processing environments.¹⁸

As a class of traditional antimicrobial agents, quaternary ammonium salts have been widely applied for preoperative disinfection of unbroken skin and non-critical surfaces,¹⁹ mucous membranes,²⁰ and hand sanitizers.²¹ Although quaternary ammonium salts have a high antimicrobial efficiency and low antimicrobial resistance,²² they exhibit a non-controllable release behavior with a short-term antibacterial activity, and meanwhile, the unnecessary release of biocides can directly results in unpredictable long-term pollution.²³ Moreover, quaternary ammonium salts are easily decomposed at a temperature higher than 150 °C. The immobilization of quaternary ammonium salts onto the solid supporters like inorganic particles is considered as an effective and ecologically friendly method that can provide a long-term durability for this antimicrobial agent.²⁴ Nowadays, the nanostructure silica materials have been accepted as a potential candidate supporter due to their characteristics of good biocompatibility, high thermal stability and easy surface modification.^{25–27} Hebalkar *et al.*²⁸ reported the synthesis of bi-functional silica particles through a simple wet chemical method by attaching titania and silver nanoparticles onto the surface of silica particles in a controlled way to form a core-shell structure, and their investigations indicated that the silica surface exhibited the efficient photo-catalytically self-cleaning and antibacterial activities. A new strategy for the preparation of mesoporous silica materials with different shapes was developed by Hao

et al.,²⁹ and they found that the antibacterial activity of these mesoporous silica materials was dependent on their shapes. Qiu *et al.*³⁰ reported an electrostatic self-assembly method to prepare core-shell structural antimicrobial nanoparticles composed of silica and polyacrylamide and found that the resultant nanoparticles showed a stable and high antibacterial activity toward both *S. aureus* and *E. coli*.

In order to provide a long-term antibacterial capability and high thermal stability for quaternary ammonium salts, we attempted to employ MMT as a framework with a nanometric region and to use silica nanoparticles as a solid supporter to immobilize the polysiloxane quaternary ammonium chloride (PQAC) as an active ingredient in this work. It is highly expected that such the layer-structural immobilization can enhance the antimicrobial performance and thermal stability of PQAC due to the nanoscale effect in MMT interlayers. With this idea in mind, we first performed a chemical graft of PQAC as an antibacterial agent onto the surface of silica nanoparticles and then intercalated these PQAC-immobilized nanoparticles into MMT through an electrostatic self-assembly method.^{31,32} The microstructures of the resulting PQAC-silica/MMT nanocomposites were intensively studied, and their thermal and antibacterial properties were investigated. The aim of this study is to develop a novel type of MMT-based functional nanocomposites with an enhanced antibacterial activity and thermal stability.

2. Experimental section

2.1. Materials

Na-MMT clay with a cationic exchange capacity of 145 mequiv./100 g was commercially obtained from Nanocor Inc., USA. The colloidal silica nanoparticles (Ludox-SM) with a particle size of 7 nm were purchased from Sigma-Aldrich, USA, and they were suspended in water with a weight fraction of 30 wt%. A solution of PQAC [(CH₃O)₃Si(CH₃)₃N⁺(CH₃)₂(C₁₀H₂₁)] in methanol with a volume fraction of 60 vol% was commercially supplied by Acros Organics, Belgium. The other chemicals were purchased from Beijing Chemical Reagent Co., Ltd., China and used as received.

2.2. Preparation methods

A quaternization reaction was first performed for the colloidal silica nanoparticles. In a typical reaction process, 7 ml of colloidal silica sol was diluted in 40 ml of deionized water, and then the resulting suspension was added dropwise into 10 ml of PQAC solution under magnetic agitation at room temperature. With the formation of a white precipitate, and the suspension was aged at room temperature with mild stirring for 24 h. Sequentially, the supernatant liquid was discarded, and the precipitate was washed thoroughly with deionized water and methanol. The resultant PQAC-modified silica nanoparticles (marked as PQAC-SiO₂) were redispersed in deionized water with a weight fraction of 1.2 wt% for further use.³³

A series of layer-structured nanocomposites with different mass ratios of MMT/PQAC-SiO₂ were prepared through an intercalating self-assembly method. In a typical process, 1.6 g of



MMT powders were dispersed in 80 ml of deionized water, and then the resulting slurry was heated to reflux at 80 °C for 6 h to make MMT powders fully swollen. The obtained suspension was centrifuged for 30 minutes, and then the suspension in the upper layer was collected with for the further intercalating self-assembly with PQAC-SiO₂ nanoparticles. The suspension containing PQAC-SiO₂ nanoparticles was added to the swollen MMT suspension and then was stirred at 70 °C for 8 h. Ultimately, the resultant MMT/PQAC-SiO₂ nanocomposites were washed several times with deionized water and then dried at 70 °C for 24 h.

2.3. Characterization and measuring method

Fourier-transform infrared (FTIR) spectroscopy was conducted on a Bruker Alpha-T FTIR spectrometer at a scanning number of 32 using a KBr sampling sheet. The thermal stability of nanocomposites was characterized by thermogravimetric analysis (TGA) on a TA Instruments SDT2960 thermogravimetric analyzer at a heating rate of 5 °C min⁻¹ under a nitrogen atmosphere. The zeta potentials of the MMT and PQAC-SiO₂ suspensions were measured by a Malvern Zetasizer Nano-ZS ZEN 3600 zeta potential analyzer. X-ray photoelectron spectroscopy (XPS) analysis was performed with a VG Scientific ESCALAB 250 X-ray photoelectron spectrometer with a focused monochromatized Al K α radiation. The obtained XPS spectra were fitted for the Casa XPS software, in which a Shirley background was assumed. X-ray diffraction (XRD) powder patterns were obtained from a Rigaku D/Max 2500 VBZ/PC X-ray diffractometer using Cu-K α radiation and operated at 40 kV and 50 mA with a scan rate of 1° min⁻¹. The morphologies of nanocomposites were observed by scanning electron microscopy (SEM) using a JEOL JSM-7800 scanning electron microscope and their microstructures were characterized by transmission electron microscopy (TEM) on a FEI Tecnai G2 20 transmission electron microscope operating at an acceleration voltage of 200 kV. The N₂ adsorption-desorption isotherms were measured at 77 K using a Quantachrome Autosorb-1 absorption analyzer with prior degassing under vacuum at 120 °C for 12 h. The data of specific surface area was calculated from the adsorption branch of the isotherm by using a Barrett-Joyner-Halenda (BJH) method.

The antimicrobial performance of MMT/PQAC-SiO₂ nanocomposites was measured according to a standard plate count method as described by the ref. 34–36 *Escherichia coli* (*E. coli*) as a Gram-positive bacterium and *Staphylococcus aureus* (*S. aureus*) as a Gram-negative bacterium were commercially obtained from National Center for Medical Culture Collections and selected to antibacterial tests, and both of them contained 10 (ref. 6 and 7) colony forming units (CFU) per each milliliter. For a typical measuring process, 50 mg ml⁻¹ of suspension containing MMT/PQAC-SiO₂ nanocomposites was mixed with 25 ml of normal saline to evaluate the antibacterial activity. Prior to the formal measurement, these bacteria were cultivated in the Luria-Bertani medium for 24 h. The cultivated bacteria were diluted and then added into the suspension containing MMT/PQAC-SiO₂ nanocomposites to achieve a viable cell concentration of

about 10³ CFU ml⁻¹. The bacteria solutions without nanocomposites were also prepared as controls. Aliquots (100 μ l) were withdrawn every 1 h to homogeneously cast on a series of Petri dishes coated with the Luria-Bertani agar medium. The plates were moved to the culturing chamber and incubated for 48 h at 37 °C. The bacterial colonies presented on the plate were imaged and counted. Each of nanocomposites was measured for each incubation time and each testing was performed in triplicate. Then, the average number of surviving bacterial colonies was obtained from the above test. The survival rates (*R*) for the adherent *E. coli* and *S. aureus* in the medium were calculated by the following formula:

$$R = \frac{C_t}{C_0} \times 100\% \quad (1)$$

where *C_t* and *C₀* present the average numbers of surviving bacterial colonies at the incubation time of 0 and *t*, respectively.

3. Results and discussion

3.1. Structural characterizations of PQAC-SiO₂ nanoparticles

PQAC-SiO₂ nanoparticles were synthesized through a polycondensation reaction between the silanol groups of silica nanoparticles and PQAC, and their chemical composition and structure were characterized by FTIR spectroscopy, and the resulting infrared spectra were illustrated in Fig. 1, where the spectra of pristine silica nanoparticles and PQAC were also presented as references. Pristine silica nanoparticles are found to exhibit two absorption peaks at 1109 and 475 cm⁻¹ in their infrared spectrum, which are attributed to the Si-O asymmetric and symmetric vibrations, respectively.³⁰ Furthermore, an absorption band corresponding to the Si-O bending vibration is also found at 799 cm⁻¹. Compared to pristine silica nanoparticles, it can be observed that the infrared spectra of PQAC-SiO₂ nanoparticles show two new absorption bands at 1038 and 1135 cm⁻¹ for the network structure formed by the Si-O-Si

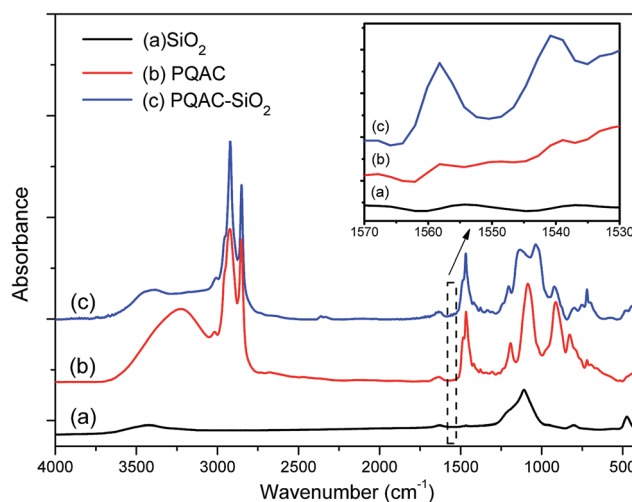


Fig. 1 FTIR spectra of pristine silica nanoparticles, PQAC and PQAC-SiO₂ nanoparticles.



linkage, implicating that the hydrolyzed PQAC has condensed with the silanol groups on the surfaces of silica nanoparticles. Moreover, there are two characteristic absorption bands observed at 1558 and 1539 cm^{-1} due to the C–N deformation vibration,³⁷ and meanwhile, the other characteristic peaks also appears at 2920, 2850, 1470, and 720 cm^{-1} in the infrared spectrum of PQAC–SiO₂ nanoparticles as observed in Fig. 1. These absorption bands are attributed to the stretching and scissoring deformation vibrations of methyl and methylene groups in PQAC, indicating that PQAC has been covalently grafted onto the silica nanoparticles. In addition, the appearance of a broad band from 3200 cm^{-1} to 3600 cm^{-1} suggests that both PQAC and PQAC–SiO₂ nanoparticles contain an abundant of hydroxyl groups.

The surface elemental compositions of PQAC–SiO₂ nanoparticles as well as the pristine silica ones were analyzed by XPS spectroscopy, and the obtained spectra were given in Fig. 2. As observe from Fig. 2, both pristine silica and PQAC–SiO₂ nanoparticles show the O 1s, C 1s and Si 2p signals at binding energy of 532.4, 284.8 and 102.9 eV in their survey XPS spectra of these two samples, respectively. It is noteworthy that the content of C atom for PQAC–SiO₂ nanoparticles increases significantly to 34.7% from 6.56% of pristine silica nanoparticles according to the signal intensity, whereas their content of O atom declines from 57.96% to 41.82%. Moreover, the N 1s signal appears at 402.45 eV in the XPS spectrum of PQAC–SiO₂ nanoparticles, which is due to the NH_4^+ group derived from PQAC.³⁸ Such a change in surface elemental composition confirms the existence of PQAC moieties on the surface of silica nanoparticles.

The thermal stabilities of PQAC–SiO₂ nanoparticles and PQAC were characterized by TGA. Fig. 3 shows the TGA and derivative TG thermograms of these two samples. PQAC is found to exhibit a typical two-step thermal degradation in the temperature ranges of 150–300 °C and 370–510 °C due to the decomposition of quaternary ammonium ions and the pyrolysis of alkane chains,³⁹ respectively, and it also shows a char yield of 18.2 wt% at the end of decomposition. It seems that PQAC–SiO₂

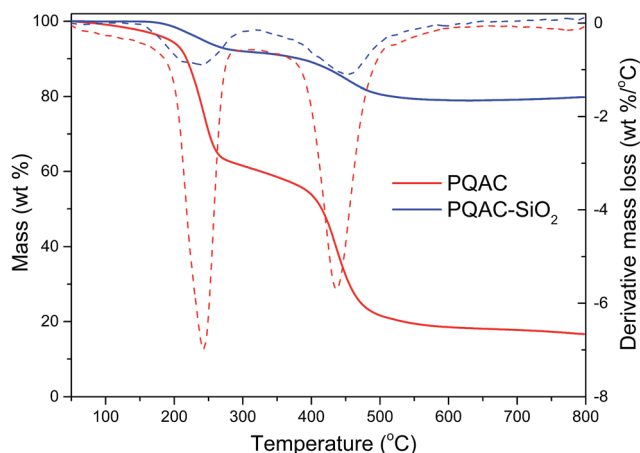


Fig. 3 TGA and DTG thermograms of silica nanoparticles, PQAC and PQAC–SiO₂ nanoparticles.

nanoparticles present a similar thermal degradation behavior with PQAC and, however, show a mass loss of 21.4 wt% at the end of thermal degradation. Such a mass loss can be attributed to the grafting ratio of PQAC on the silica nanoparticles.

3.2. Preparation and structural characterizations of MMT/PQAC–SiO₂ composites

The MMT/PQAC–SiO₂ composites were prepared through an intercalating electrostatic self-assembly method. This self-assembly mechanism is schematically illustrated in Fig. 4. It is expected that PQAC–SiO₂ nanoparticles are endowed with a positive charge on their surfaces through the functionalization with PAQC. On the other hand, MMT as a layer-structured material carries a permanent negative charge on the laminates, which has been balanced by the cationic counterions within the interlayer space. In this case, positively charged PQAC–SiO₂ nanoparticles are introduced into the negatively charged MMT laminates, a rapid heterocoagulation takes place definitely due to multiple charge complex formation between two materials resulting from electrostatic interaction.⁴⁰ Fig. 5 shows the zeta potentials of pristine silica nanoparticles, PQAC–SiO₂ nanoparticles and MMT with variation of pH values of reactant

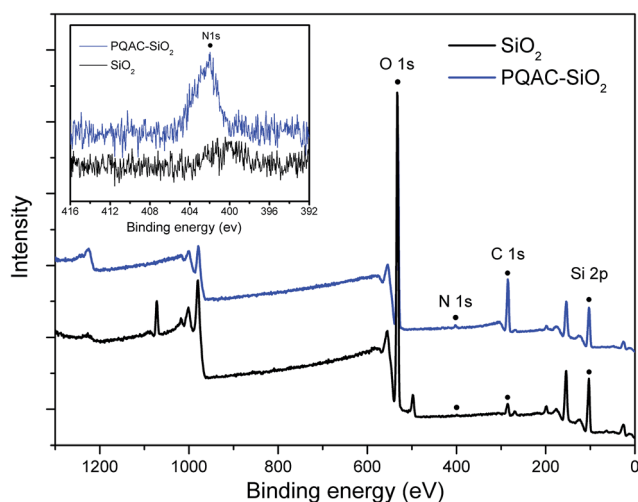


Fig. 2 XPS spectra of pristine silica nanoparticles and PQAC–SiO₂ nanoparticles.

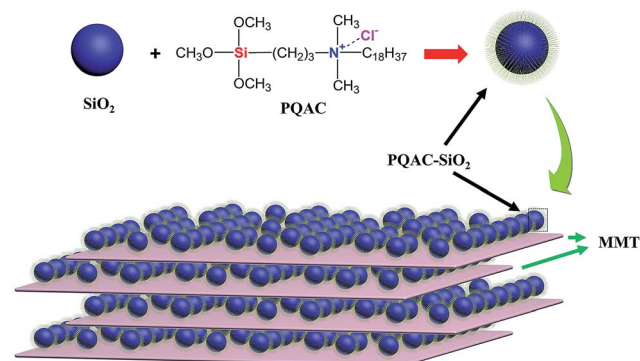


Fig. 4 Schematic formation process of MMT/PQAC–SiO₂ nanocomposites.



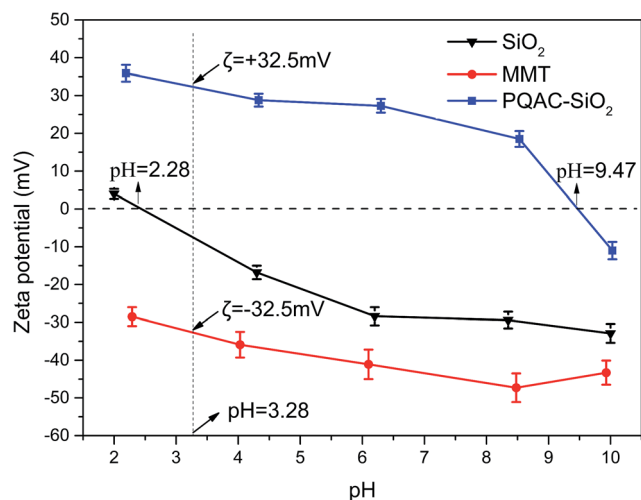


Fig. 5 Plots of zeta potential as a function of pH value for silica nanoparticles, MMT and PQAC-SiO₂ nanoparticles in water.

suspension. It is observed in Fig. 5 that pristine silica nanoparticles shows a gradually decreasing negative zeta potential with an increase of pH value and also presents an isoelectric point at pH 2.28. However, the zeta potential of PQAC-SiO₂ nanoparticles exhibit a significant variation trend ranging from +35.9 to -10.8 mV with increasing the pH value of reactant suspension, and their isoelectric point was found at pH 9.47. On the other hand, MMT presents a negative zeta potential variation in the range of -28.5 to -47.3 mV. In this case, it is easily determined that PQAC-SiO₂ nanoparticles and MMT have the zeta potentials of +32.5 and -32.5 mV, respectively, when the acidity of both of reactant suspensions are set to pH 3.2. Therefore, a maximum electrostatic interaction can be achieved at pH 3.2 between these two materials when they are mixed. As a result, PQAC-SiO₂ nanoparticles are rapidly attracted into the MMT interlayers to form the highly ordered layer-structured nanocomposites.

Fig. 6 shows the XRD patterns of layer-structured MMT/PQAC-SiO₂ nanocomposites at different weight ratios. All of the nanocomposites are observed to exhibit a diffraction peak at $2\theta = 7.3^\circ$ in their XRD patterns, which is assigned to the (001) reflection of MMT with a d -spacing of 1.2 nm determined by Bragg's law. It is well known that the pristine MMT layers can stack with one another by the attraction force between their cationic edges and anionic surfaces during the drying process due to their ionic nature, thus creating a d -spacing of 1.2–1.5 nm. This phenomenon could be typically observed from various MMT-based materials and is associated with the water content of MMT.⁴¹ Nevertheless, it is noteworthy in Fig. 6a that this diffraction peak exhibits a gradual decline in its intensity with increasing the weight fraction of PQAC-SiO₂ nanoparticles, suggesting the gradual elimination of ordered structure in the (001) plane of MMT. Meanwhile, a new diffraction peak appears at $2\theta = 0.84^\circ$ in their XRD patterns, which corresponds to a d -spacing of 10.6 nm in the MMT/PQAC-SiO₂ nanocomposites. Such a new diffraction peak is attributed to the formation of pillared structure supported by intercalating PQAC-SiO₂

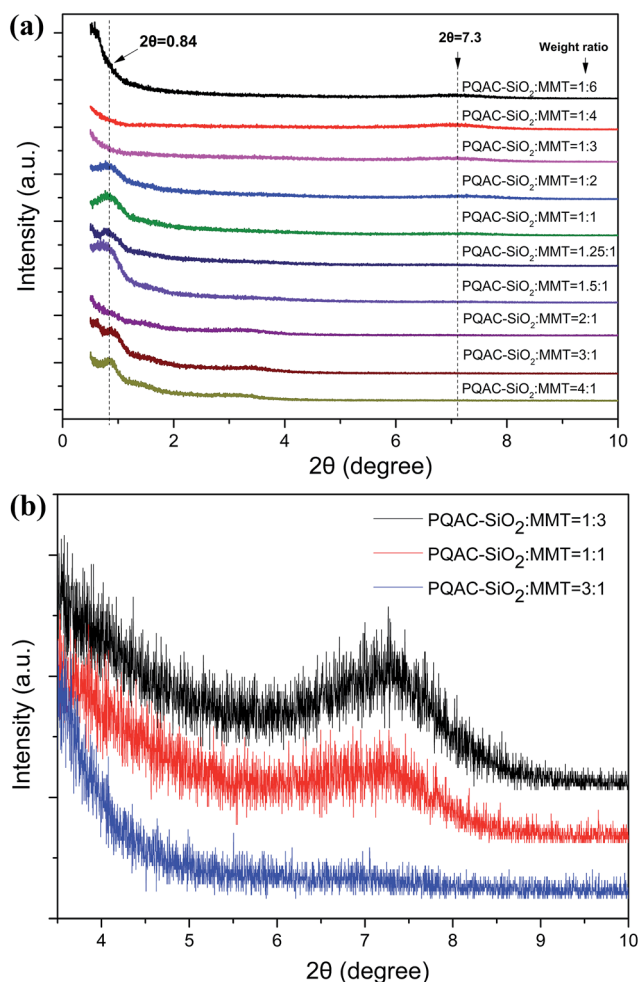


Fig. 6 X-ray diffraction patterns of MMT/PQAC-SiO₂ nanocomposites with different weight ratios of MMT/PQAC-SiO₂ nanocomposites.

nanoparticles into the interlayers of MMT, thus resulting in an enlarged interlayer space. Moreover, two diffraction peaks can be simultaneously observed from the MMT/PQAC-SiO₂ nanocomposite at the weight ratio of 1/2 as shown in Fig. 6, indicating the coexistence of the original layered structure of MMT and the intercalation one formed by PQAC-SiO₂ nanoparticles. However, with continually increasing the weight fraction of PQAC-SiO₂ nanoparticles, more and more PQAC-SiO₂ nanoparticles were intercalated into the interlayers of MMT until each of interlayers was full with the PQAC-SiO₂ nanoparticles. In this case, the diffraction peak at $2\theta = 7.3^\circ$ is found to disappear at a high weight fraction of PQAC-SiO₂ nanoparticles, whereas the peak at $2\theta = 0.84^\circ$ is noticeably enhanced as seen in Fig. 6b. The interlayer space of MMT/PQAC-SiO₂ nanocomposites could be determined as 9.4 nm by the new diffraction data according to Bragg's law, and meanwhile, the size of PQAC-SiO₂ nanoparticles could also be estimated as 9.4 nm.

The morphologies and microstructures of MMT/PQAC-SiO₂ nanocomposites were observed by SEM and then confirmed by TEM. Fig. 7 shows the SEM micrographs of layer-structured nanocomposites at different MMT/PQAC-SiO₂ weight ratios. It



is clearly observed in Fig. 7a–d that PQAC–SiO₂ nanoparticles are homogeneously dispersed on the surfaces of MMT lamellae due to the electrostatic self-assembly, and most of them are monodispersed. Furthermore, the distribution of nanoparticles seems to become denser with an increase of weight fraction of PQAC–SiO₂ nanoparticles. However, there are some agglomerated nanoparticles observed from Fig. 7a–d due to the overloading of PQAC–SiO₂ nanoparticles in MMT. On the other hand, the TEM micrographs in Fig. 8 confirm the uniform distribution of PQAC–SiO₂ nanoparticles within the interlayers of MMT.⁴² These nanoparticles are observed to exhibit a regular array between the MMT lamellae, and the boundary between

the nanoparticles is easily distinguished without overlapping because of their cationic nature. These results convincingly confirm that the formation of layer-structured nanocomposites based on multilayered MMT and PQAC–SiO₂ nanoparticles.

The layer-structural characteristics of MMT/PQAC–SiO₂ nanocomposites were further analyzed by N₂ adsorption-desorption experiments. Fig. 9 shows the representative N₂ adsorption-desorption isotherms of MMT/PQAC–SiO₂ nanocomposites as well as their specific surface areas calculated by the BJH method. As displayed in Fig. 9a, the PQAC–SiO₂ nanoparticles are found to present a typical Type IV isotherm with an H2 hysteresis loop according to the Brunauer–Deming–

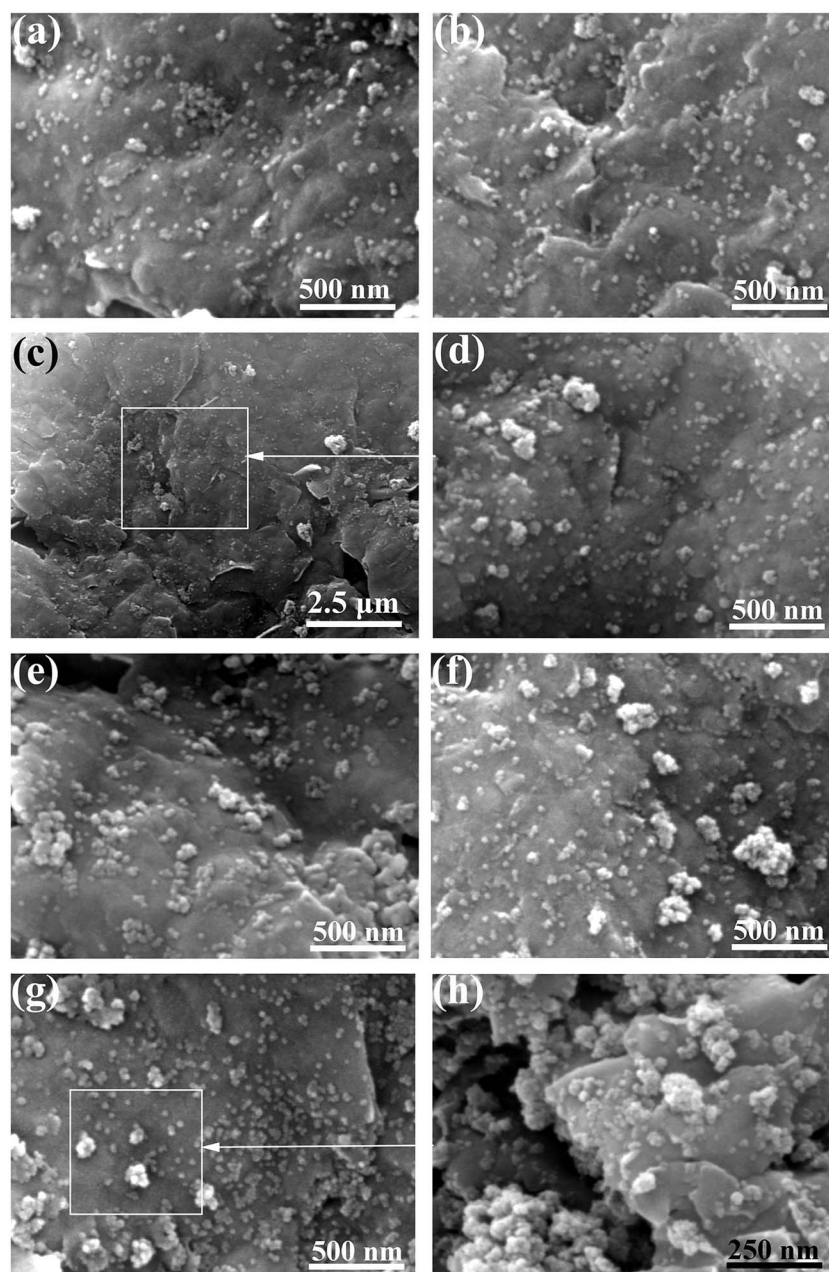


Fig. 7 SEM micrographs of MMT/PQAC–SiO₂ nanocomposites with MMT/PQAC–SiO₂ weight ratios of (a) 4/1, (b) 3/1, (c, d) 1/1, (e) 1/1.25 (f) 1/2 and (g, h) 1/3.

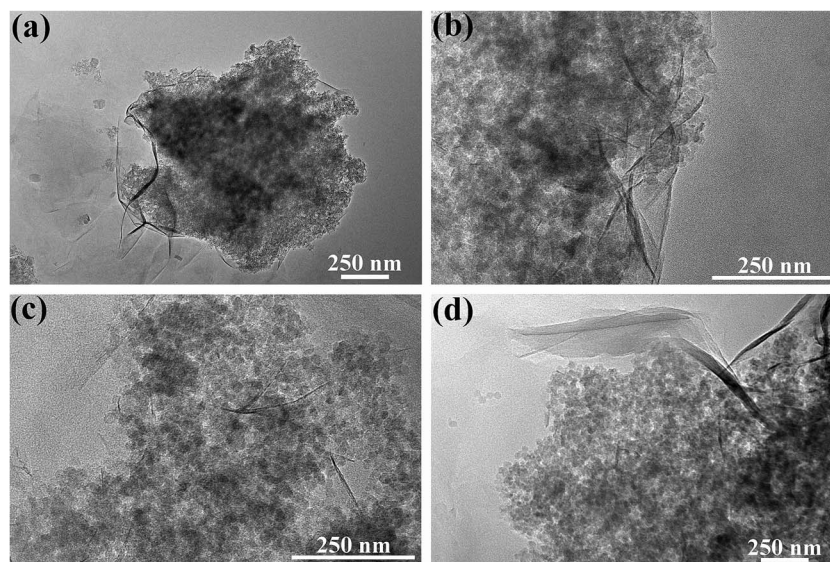


Fig. 8 TEM micrographs of MMT/PQAC-SiO₂ nanocomposites with MMT/PQAC-SiO₂ weight ratios of (a) 3/1, (b) 1/1, (c) 1/2, and (d) 1/3.

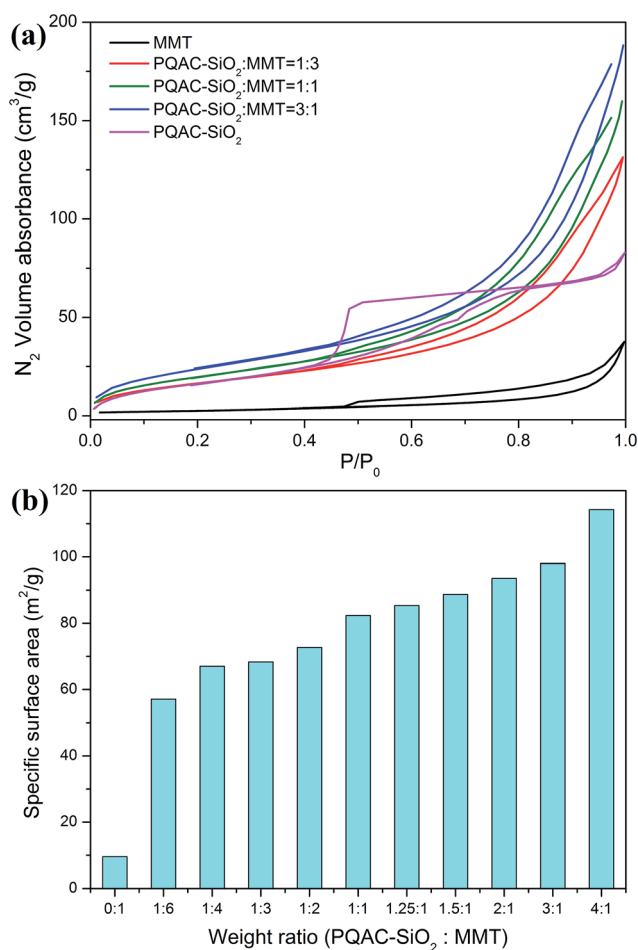


Fig. 9 (a) The N₂ adsorption-desorption isotherms and (b) specific surface areas of pure MMT, PQAC-SiO₂ nanoparticles and MMT/PQAC-SiO₂ nanocomposites.

Deming-Teller (BDDT) classification. Such a characteristic isotherm is ascribed to the existence of microporous structure caused by the aggregation of PQAC-SiO₂ nanoparticles.⁴³ On the other hand, both pristine MMT and MMT/PQAC-SiO₂ nanocomposites exhibit a gradual N₂-adsorption increase with pressure in the relative pressure (P/P_0) range of 0–0.48 and then a desorption delay in the P/P_0 range of 0.48–0.9 in their N₂ adsorption-desorption isotherms, which can be assigned to a Type IV isotherms with a typical H4 hysteresis loop.⁴ The MMT/PQAC-SiO₂ nanocomposites are found to present a monolayer adsorption on the surfaces of MMT in the initial stage of isotherms at low P/P_0 values and then a multilayer adsorption in the P/P_0 range of 0.05–0.48. When the pressure increases to high values ($P/P_0 = 0.48$ –0.9), a steep increment appears due to capillary condensation within the interlayers of MMT/PQAC-SiO₂ nanocomposites followed by a saturation of N₂ gas as the interlayers were filled with liquid nitrogen. These results confirm the existence of a layered structure within the MMT/PQAC-SiO₂ nanoparticles. Furthermore, it is noticeable in Fig. 9b that the specific surface areas of MMT/PQAC-SiO₂ nanocomposites exhibit a significant increase compared to that of pristine MMT, and the higher the weigh fraction of PQAC-SiO₂ nanoparticles, the larger the specific surface areas. This phenomenon is attributed to the introduction of PQAC-SiO₂ nanoparticles with a larger surface area as well as the enlargement of interlayer space of MMT.

3.3. Thermal stability

The thermal stabilities of MMT/PQAC-SiO₂ nanocomposites were investigated by TGA, and the resulting TGA and derivate TG (DTG) thermograms are depicted in Fig. 10. It is observed that pristine MMT exhibits two distinct weight losses at 25–100 °C and 600–650 °C due to the desorption of surface free water and interlayer water, respectively.⁴⁴ On the other hand, MMT/PQAC-SiO₂ nanocomposites are found to exhibit a two-



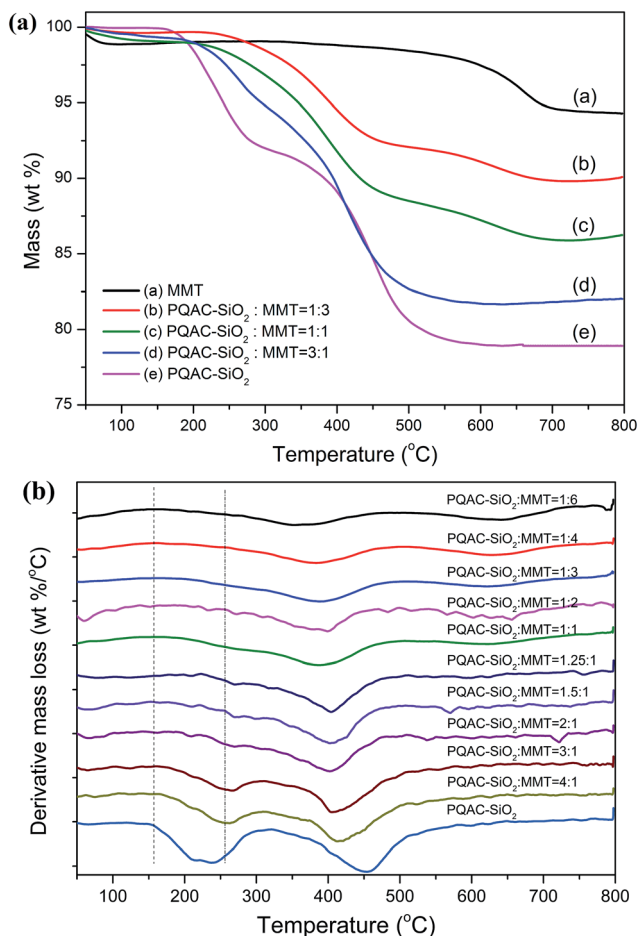


Fig. 10 (a) TGA and (b) DTG thermograms of MMT, PQAC-SiO₂ nanoparticles and MMT/PQAC-SiO₂ nanocomposites.

step decomposition behavior in the presence of PQAC-SiO₂ nanoparticles. The first degradation is attributed to the decomposition of quaternary ammonium ions on the PQAC-SiO₂ nanoparticles and the second one is ascribed to the pyrolysis of alkane chains of PQAC as mentioned previously. However, the initial weight loss at 25–100 °C is not observed from the MMT/PQAC-SiO₂ nanocomposites, indicating that these nanocomposites have been dried completely. It is noteworthy in Fig. 10a that the initial decomposition temperature of MMT/PQAC-SiO₂ nanocomposites is improved significantly to approximately 200 °C compared to that of pristine PQAC-SiO₂ nanoparticles. Such an improvement in initial decomposition temperature is ascribed to the fact that the PQAC-SiO₂ nanoparticles are well protected through an interlayer effect when intercalated into the interlayers of MMT.^{37,41} Moreover, as depicted in Fig. 10b, both the initial decomposition temperature and the maximum rapid degradation temperature tend to increase with increasing the weight fraction of PQAC-SiO₂ nanoparticles. Meanwhile, it is interesting to note that the weight loss of interlayer water gradually disappears with increasing the weight fraction of PQAC-SiO₂ nanoparticles. This may be due to the fact that the formation of pillaring structure

by PQAC-SiO₂ nanoparticles enlarges the interlayer spacing and thus makes the interlayer water easily evaporate during the drying process. These results suggest that the intercalation of PQAC-SiO₂ nanocomposites into MMT can effectively improve the thermal stability of MMT/PQAC-SiO₂ nanocomposites through the interlayer effect and therefore enhances the durability and reliability of PQAC-SiO₂ nanoparticles when used as an antibacterial agent.

3.4. Antibacterial performance

A time-dependent antibacterial study was conducted to evaluate the antibacterial activities of MMT/PQAC-SiO₂ nanocomposites against *E. coli* and *S. aureus* as the model microorganisms of Gram-negative and Gram-positive bacteria, respectively. A colony counting method was adopted to determine the inhibition of bacteria growth at different incubation time under the antibacterial action of MMT/PQAC-SiO₂ nanocomposites. Fig. 11 shows the survival rates of two bacteria as a function of contact time upon exposure to MMT/PQAC-SiO₂ nanocomposites. It seems that the MMT/PQAC-SiO₂ nanocomposites exhibit the almost same antibacterial effect on *E. coli* and *S. aureus*. As shown in Fig. 11, *E. coli* and *S. aureus* show a drastic decrease in survival rate during the initial four hours when contacting the nanocomposites, and then, their decrements tend to decline. After contacting with the nanocomposites for 7 h, both *E. coli* and *S. aureus* were exterminated completely. This sterilization process can also be observed visibly from the digital photographs of the colonies of bacteria incubated on agar plates in Fig. 12. These photographs clearly demonstrate a significant reduction in the number of bacterial colonies with an increase of contact time. This result confirms an excellent antibacterial effectiveness of MMT/PQAC-SiO₂ nanocomposites against Gram-positive and Gram-negative bacteria.

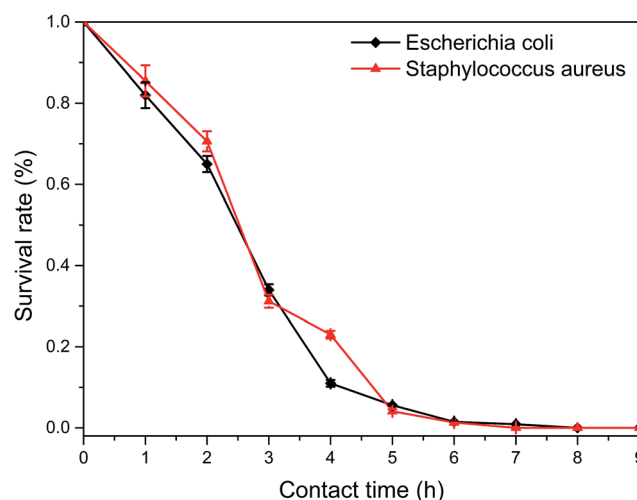


Fig. 11 Plots of the survival rates of model microorganisms as a function of contact time with the MMT/PQAC-SiO₂ nanocomposites as an antibacterial agent.



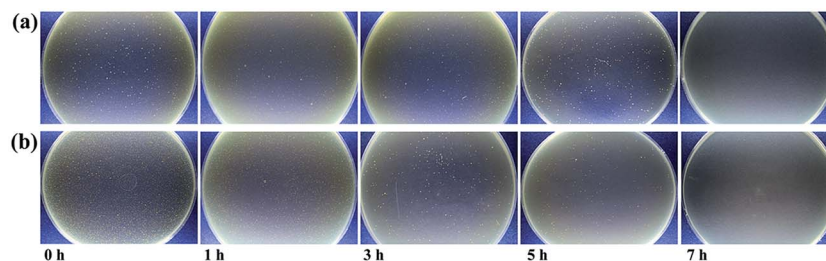


Fig. 12 Optical photographs of the Petri dishes loading (a) *E. coli* and (b) *S. aureus* at different contact time.

4. Conclusions

The positively charged PQAC-SiO₂ nanoparticles were synthesized by grafting PQAC onto the surface of silica nanoparticles and their chemical composition and structure were confirmed by FTIR spectroscopy, XPS and TGA. The layer-structured MMT/PQAC-SiO₂ nanocomposites were successfully prepared by intercalating PQAC-SiO₂ nanoparticles into the interlayers of MMT through an electrostatic self-assembly method. The layered structure of MMT/PQAC-SiO₂ nanocomposites was confirmed by XRD patterns, N₂ adsorption-desorption analysis, SEM and TEM, and the optimum weight ratio of 2/1 between PQAC-SiO₂ nanoparticles and MMT was obtained on the basis of these characterization results. Further studies by N₂ adsorption-desorption experiments of nanocomposites confirmed the existence of a layered structure within the MMT/PQAC-SiO₂ nanocomposites. The intercalation of PQAC-SiO₂ nanoparticles into MMT also improved their thermal stability, resulting in a remarkable increment in onset thermal decomposition from 160 °C to 200 °C in comparison with PQAC-SiO₂ nanoparticles. The MMT/PQAC-SiO₂ nanocomposites also exhibited an excellent antibacterial capability against *E. coli* and *S. aureus* as the model microorganisms of Gram-negative and Gram-positive, respectively.

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References

- 1 H. H. Murray, *Appl. Clay Sci.*, 2000, **17**, 207–221.
- 2 M. Kotal and A. K. Bhowmick, *Prog. Polym. Sci.*, 2015, **51**, 127–187.
- 3 A. C. Lopes, P. Martins and S. Lanceros-Mendez, *Prog. Surf. Sci.*, 2014, **89**, 239–277.
- 4 B. Li, H. Mao, X. Li, W. Ma and Z. Liu, *J. Colloid Interface Sci.*, 2009, **336**, 244–249.
- 5 M. Polverejan, Y. Liu and T. J. Pinnavaia, *Chem. Mater.*, 2002, **14**, 2283–2288.
- 6 J. Ahenach, P. Cool and E. F. Vansant, *Phys. Chem. Chem. Phys.*, 2000, **2**, 5750–5755.
- 7 A. De Stefanis and A. A. G. Tomlinson, *Catal. Today*, 2006, **114**, 126–141.
- 8 N. Bitinis, M. Hernandez, R. Verdejo, J. M. Kenny and M. A. Lopez-Manchado, *Adv. Mater.*, 2011, **23**, 5229–5236.
- 9 Y. S. Han and S. Yamanaka, *J. Solid State Chem.*, 2006, **179**, 1146–1153.
- 10 Y. Kameshima, Y. Tamura, A. Nakajima and K. Okada, *Appl. Clay Sci.*, 2009, **45**, 20–23.
- 11 N. Khalfoui-Boutoumi, H. Boutoumi, H. Khalaf and B. David, *Appl. Clay Sci.*, 2013, **80**, 56–62.
- 12 H. Mao, B. Li, L. Yue, L. Wang, J. Yang and X. Gao, *Appl. Clay Sci.*, 2011, **53**, 676–683.
- 13 M. C. Margarita Darder and E. Ruiz-Hitzky, *Chem. Mater.*, 2003, **2003**, 3774–3780.
- 14 J. J. Lin, W. C. Lin, S. D. Li, C. Y. Lin and S. H. Hsu, *ACS Appl. Mater. Interfaces*, 2013, **5**, 433–443.
- 15 M. Lavorgna, I. Attianese, G. G. Buonocore, A. Conte, M. A. Del Nobile, F. Tescione and E. Amendola, *Carbohydr. Polym.*, 2014, **102**, 385–392.
- 16 G. A. Sotiriou, A. Meyer, J. T. Knijnenburg, S. Panke and S. E. Pratsinis, *Langmuir*, 2012, **28**, 15929–15936.
- 17 L. Zhou, H. Chen, X. Jiang, F. Lu, Y. Zhou, W. Yin and X. Ji, *J. Colloid Interface Sci.*, 2009, **332**, 16–21.
- 18 R. K. Khalil, *World J. Microbiol. Biotechnol.*, 2013, **29**, 1839–1850.
- 19 E. Obłak, A. Piecuch, A. Krasowska and J. Łuczyński, *Microbiol. Res.*, 2013, **168**, 630–638.
- 20 M. Tischer, G. Pradel, K. Ohlsen and U. Holzgrabe, *ChemMedChem*, 2012, **7**, 22–31.
- 21 K. Hegstad, S. Langsrud, B. T. Lunestad, A. A. Scheie, M. Sunde and S. P. Yazdankhah, *Microb. Drug Resist.*, 2010, **16**, 91–104.
- 22 B. Dizman, M. O. Elasri and L. J. Mathias, *J. Polym. Sci., Part A: Polym. Chem.*, 2006, **44**, 5965–5973.
- 23 A. W. Smith, *Adv. Drug Delivery Rev.*, 2005, **57**, 1539–1550.
- 24 X. Ye, J. Feng, J. Zhang, X. Yang, X. Liao, Q. Shi and S. Tan, *Colloids Surf., B*, 2017, **149**, 322–329.
- 25 A. Dong, Q. Zhang, T. Wang, W. Wang, F. Liu and G. Gao, *J. Phys. Chem. C*, 2010, **114**, 17298–17303.
- 26 B. Song, C. Wu and J. Chang, *Acta Biomater.*, 2012, **8**, 1901–1907.
- 27 G. S. Alvarez, C. Helary, A. M. Mebert, X. Wang, T. Coradin and M. F. Desimone, *J. Mater. Chem. B*, 2014, **2**, 4660–4670.
- 28 N. Y. Hebakkar, S. Acharya and T. N. Rao, *J. Colloid Interface Sci.*, 2011, **364**, 24–30.
- 29 N. Hao, X. Chen, K. W. Jayawardana, B. Wu, M. Sundhoro and M. Yan, *Biomater. Sci.*, 2016, **4**, 87–91.



- 30 Q. Qiu, T. Liu, Z. Li and X. Ding, *J. Mater. Chem. B*, 2015, **3**, 7203–7212.
- 31 X. S. Zhou, Y. X. Yin, L. J. Wan and Y. G. Guo, *Adv. Energy Mater.*, 2012, **2**, 1086–1090.
- 32 F. Z. Tu, X. Xu, P. Z. Wang, L. Si, X. S. Zhou and J. C. Bao, *J. Phys. Chem. C*, 2017, **121**, 3261–3269.
- 33 A. B. Bourlinos, R. Herrera, N. Chalkias, D. D. Jiang, Q. Zhang, L. A. Archer and E. P. Giannelis, *Adv. Mater.*, 2005, **17**, 234–237.
- 34 X. Zhang, X. Wang and D. Wu, *Energy*, 2016, **111**, 498–512.
- 35 D. Q. Wu, H. C. Cui, J. Zhu, X. H. Qin and T. Xie, *J. Mater. Chem. B*, 2016, **4**, 2606–2613.
- 36 F. Li, X. Wang and D. Wu, *Energy Convers. Manage.*, 2015, **106**, 873–885.
- 37 T. F. Tseng and J. Y. Wu, *J. Mater. Chem. C*, 2009, **113**, 13036–13044.
- 38 N. Andreu, D. Flahaut, R. Dedryvere, M. Minvielle, H. Martinez and D. Gonbeau, *ACS Appl. Mater. Interfaces*, 2015, **7**, 6629–6636.
- 39 K. Fujii, S. Hayashi and H. Kodama, *Chem. Mater.*, 2003, **15**, 1189–1197.
- 40 Y. Zheng and A. Zaoui, *Solid State Ionics*, 2011, **203**, 80–85.
- 41 A. Hilonga, J. K. Kim, P. B. Sarawade, D. V. Quang, G. N. Shao, G. Elineema and H. T. Kim, *Mater. Lett.*, 2012, **80**, 168–170.
- 42 X. Liu, Y. C. Du, L. Y. Hu, X. S. Zhou, Y. F. Li, Z. H. Dai and J. C. Bao, *J. Phys. Chem. C*, 2015, **119**, 5848–5854.
- 43 N. Ahmad, S. T. Hussain, B. Muhammad, N. Ali, S. M. Abbas and Z. Ali, *Prog. Nat. Sci.: Mater. Int.*, 2013, **23**, 374–381.
- 44 B. Wu, S. Qi and X. Wang, *Polym. Test.*, 2010, **29**, 717–722.

