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Facile shape-controlled synthesis of lanthanum oxide with different hierarchical micro/nanostructures for antibacterial activity based on phosphate removal

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In this study, three La_2O_3 hierarchical micro/nanostructures, nanospindles, nanopolyhedra and nanospheres, were synthesized to remove phosphate from microbial growth media for bacterial inactivation as an antimicrobial strategy. The experiment results showed that the morphology of La_2O_3 hierarchical micro/nanostructures can be controlled by the concentration of the reactants, reaction temperature and cooling treatment. Meanwhile, the properties of the La_2O_3 were researched for phosphate removal and antibacterial activity. The results showed that the different morphologies of the La_2O_3 micro/nanostructures had different removal abilities for phosphate, and can differently inhibit growth of bacteria. Spherical La_2O_3 possesses the best removal ability and antibacterial activity, which indicates that the smaller the La_2O_3 hierarchical micro/nanoparticles are, the stronger the phosphate removal capacity. Compared with nanoparticles, the hierarchical micro/nanoparticles have a significant advantage: they not only have the properties of nanoparticles, but also are relatively stable, not easy to aggregate, and are easy to separate after the reaction. Therefore, the La_2O_3 hierarchical micro-nanomaterials may have very good application prospects for phosphate reduction in open water and to inhibit algae overgrowth.

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1 Introduction

Recently, lanthanum compounds have attracted considerable attention in hydrogen storage, electrode and sorbent materials, gate insulators and superconductors due to their versatility and multifunctionality.^{1–4} Meanwhile, it is known that lanthanum compounds, such as lanthanum carbonate (La_2CO_3), lanthanum hydroxide ($\text{La}(\text{OH})_3$), lanthanum oxide carbonate ($\text{La}_2\text{O}_2\text{CO}_3$),^{5–7} and mixtures of lanthanum compounds (La_2O_3 , $\text{La}_2\text{O}_2\text{CO}_3$ and $\text{La}_2(\text{OH})_3$),⁸ can bind to phosphate so strongly that they can form LaPO_4 and remove excess phosphate in bacterial or algal growth media,⁸ and lead to inhibit the growth of microorganisms and algae. As a typical representative, lanthanum carbonate has been developed as a medicine to reduce the excess phosphate in the human body.⁹ Based on the

existing reports on bacterial and algal limitation utilizing lanthanum species,^{10–12} we here synthesize three La_2O_3 hierarchical micro/nanostructures and investigate their phosphate removal ability and antibacterial activity.

It is widely accepted that phosphate is essential for growth of all organisms and serves as the main building block for nucleic acids, proteins, and energy carriers.¹³ Phosphate is the only form of phosphorus that can be directly assimilated by microorganisms. Firstly, the initial assimilation of inorganic phosphate for microorganisms proceeds *via* phosphorylation of ADP, which is affected by phosphate limitation in the environment.⁵ In addition, abundant phosphate is an essential condition to synthesize the cell wall of Gram-positive bacteria and the lipid composition of the membranes of Gram-negative bacteria. It could be affected strikingly within phosphate limitation environment.⁶ Consequently, the lack of phosphate often limits their growth, and efficient phosphate removal in water is expected to be a green strategy to prevent the growth of microorganisms and algae, which would avoid toxic substances released from antibacterial agents.^{14–16} On the other hand, reuse of secondary municipal effluent from wastewater treatment plants in water bodies could effectively alleviate freshwater resource shortage.⁹ During the wastewater treatment, excessive nutrients, *e.g.* phosphate, must be efficiently removed to prevent eutrophication.¹⁷

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Considering the strong combination of lanthanum to phosphate and the valid antimicrobial strategy resulted from nutrient starvation of phosphate removal, it is meaningful to explore a facile, high quality and low cost method to synthesize La_2O_3 with micro/nanostructure. In the current study, shape-controlled La_2O_3 hierarchical micro/nanostructures were prepared through the regulation of the concentration of the reactants and the reaction temperature. Three morphologies of La_2O_3 micro/nanostructures were obtained, and were utilized to compare antimicrobial activity. The results indicated that lanthanum oxide can provide a broadly applicable antimicrobial strategy.

2 Experimental section

Material preparation

Series of morphologies micro-nanostructures of La_2O_3 were synthesized by a simple urea homogeneous precipitation process. All the chemicals, lanthanum nitrate hexahydrate $\text{La}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, and urea were of analytical pure grade. The morphologies of La_2O_3 were controlled by the adjustment of the concentration of the reactants and the reaction temperature. The synthesis process consists of two steps: synthesis of $\text{La}(\text{OH})\text{CO}_3$ precursor and La_2O_3 product. Firstly, spindle $\text{La}(\text{OH})\text{CO}_3$ was synthesized at 90 °C with the concentration ratio of 1 : 2 between $\text{La}(\text{NO}_3)_3$ and $\text{CO}(\text{NH}_2)_2$. While polyhedron $\text{La}(\text{OH})\text{CO}_3$ were prepared at 125 °C with the concentration ratio of 1 : 6 between $\text{La}(\text{NO}_3)_3$ and $\text{CO}(\text{NH}_2)_2$. In the same manner, when the concentration ratio between $\text{La}(\text{NO}_3)_3$ and $\text{CO}(\text{NH}_2)_2$ is 1 : 400, spherical $\text{La}(\text{OH})\text{CO}_3$ was successfully synthesized at 85 °C. Subsequently, the $\text{La}(\text{OH})\text{CO}_3$ precursor was calcined from room temperature to 800 °C with a heating rate of 1 °C min^{-1} and maintained at this temperature for 4 h. Then spherical micro/nanostructures La_2O_3 were obtained. Other La_2O_3 (spindles, polyhedrons) samples were prepared by a similar procedure with different $\text{La}(\text{OH})\text{CO}_3$ precursors.

Characterization

XRD measurements were determined using a D8 ADVANCE X-ray diffractometer at a scanning rate of 15° min^{-1} in the 2θ range from 10–80°, with graphite monochromatic Cu K α radiation ($\lambda = 0.15405$ nm). Perkin-Elmer 580B infrared spectrophotometer was used to measure Fourier transform infrared spectroscopy (FT-IR) spectra with the KBr pellet technique. Thermogravimetric and differential thermal analysis (TG-DTA) data were recorded with Thermal Analysis instrument (SDT 2960, TA Instruments, New Castle, DE) with the heating rate of 10 °C min^{-1} in an air flow of 100 mL min^{-1} . SEM micrographs were obtained using a field emission scanning electron microscope (FE-SEM, S-4800, Hitachi). Microstructures and morphologies were investigated using scanning electronic microscopy (SEM, JEOL JSM-6390LV), transmission electron microscope (TEM, JEM-2100) and field emission scanning electron microscopy (FESEM, SUPRA40). All the measurements were performed at room temperature (RT). Brunauer–Emmett–Teller (BET) (Tristar Micromeritics, Norcross, GA, USA) method

was used to determine the specific surface area at 77 K. The BET analysis was measured after out gassing for 90 minutes at 250 °C.

Phosphate absorption capacity of lanthanum oxide nanoparticles and antibacterial activity

To evaluate the binding affinity of the nanoparticles to phosphate, PO_4^{3-} was tested by colorimetry assay. Firstly, standard curve was drawn. Secondly, residual PO_4^{3-} was determined. Specifically, 10% ascorbic acid was freshly prepared as follow: 10 g ascorbic acid was dissolved in 100 mL ddH $_2\text{O}$. Then molybdate solution were prepared as follows: 13 g of $(\text{NH}_4)_6\text{Mo}_7\text{O}_{24} \cdot 4\text{H}_2\text{O}$ was dissolved in 100 mL of double distilled water (ddH $_2\text{O}$). Meanwhile, 0.35 g $\text{K}(\text{SbO})\text{C}_4\text{H}_4\text{O}_6 \cdot 1/2\text{H}_2\text{O}$ was dissolved in 100 mL of ddH $_2\text{O}$. The prepared $(\text{NH}_4)_6\text{Mo}_7\text{O}_{24}$ and $\text{K}(\text{SbO})\text{C}_4\text{H}_4\text{O}_6$ solutions were added to 300 mL of dilute sulfuric acid successively. Whereafter the mixed solution was stirred and stored in 4 °C. Then La_2O_3 micro/nanoparticles with different morphologies were added to 15, 25, and 50 $\mu\text{g mL}^{-1}$ phosphate solution, respectively. 1 mL of ascorbic acid (10%), 2 mL of molybdate solution and above La_2O_3 samples were mixed to determine residual PO_4^{3-} according to absorbance at 710 nm by 752 Ultraviolet Spectrophotometer.¹⁸

Escherichia coli (strain C43) and *Staphylococcus aureus* (strain TM300) were grown in Difco™ LB broth (Chemie Brunschwig) for 4 h at 37 °C and gently agitation to a concentration of about 10⁸ CFU mL^{-1} . This suspension was diluted to the required concentration with physiological saline (0.9 wt% NaCl in water) or where medium free (in particular phosphate free) was needed, repeatedly (5 times) centrifuged for 60 min at RCF = 1300 (Mistral 3000E, 2500 rpm). Subsequently, the supernatant was removed and the bacterial pellet was resuspended in minimal media. Minimal media was produced from 5 g glucose, 1 g NH_4Cl , 5.4 g NaCl, 0.1 g MgSO_4 and 0.02 g CaCl_2 hexahydrate filled up to 1 liter with ddH $_2\text{O}$ and sterilized by autoclaving at 121 °C for 15 min. To quantify the CFU load in a sample, a dilution row (10⁻¹–10⁻⁸) was plated in duplicate on dextrose agar plates (PDA, VWR BDH Prolabo). The plates were incubated at 37 °C for 24 h before readout.

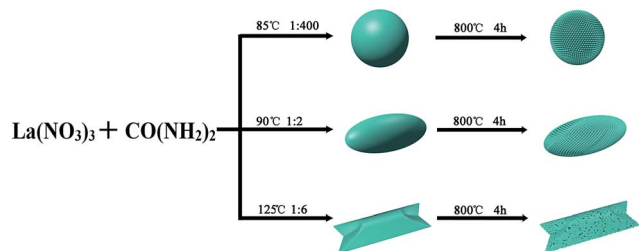
3 Results and discussion

Phase structure, morphology and formation mechanism of La_2O_3

Polyhedral, spherical and spindles-like La_2O_3 micro/nanomaterials with hierarchical micro/nanostructures were synthesized by a simple precipitation method with urea and lanthanum nitrate. The synthesis process consists of two steps: synthesis of $\text{La}(\text{OH})\text{CO}_3$ and La_2O_3 (Scheme 1).

The morphology and microstructure of the products were characterized by SEM and TEM (shown in Fig. 1). The SEM images reveal that three different morphologies of La_2O_3 samples are successfully obtained, including polyhedral, spherical and spindles-like micro-nanostructures. The sizes of the samples are about 1 μm , 10 μm and 12 μm for spherical





Scheme 1 Schematic illustration for preparation of micro/nano-structures La_2O_3 with a simple precipitation method using urea and lanthanum nitrate.

La_2O_3 , polyhedral La_2O_3 and spindles La_2O_3 respectively. From Fig. 1 A(b), B(e) and C(h), the HRTEM images reveal that the surfaces of samples are very rough. It can be further observed that the samples consist of many even smaller nanoparticles with the size of 8–15 nm (inset of Fig. 1A(b), 2B(e) and 2C(h)),

showing that the three morphologies of La_2O_3 possess hierarchical micro-nanostructures. The average diameters of the smaller nanoparticles are 8.58 ± 0.08 nm for spherical La_2O_3 (Fig. 1 A(c)), 15.58 ± 0.08 nm for polyhedral La_2O_3 (Fig. 1B(f)) and 13.45 ± 0.08 nm for spindles-like La_2O_3 (Fig. 1C(i)), respectively. In addition, the HRTEM shows that the values of inter planar lattice spacing are 0.29 nm, 0.31 nm and 0.34 nm, respectively, corresponding to the (101), (100) and (110) plane of La_2O_3 . The results further show that the prepared La_2O_3 nanocrystals belong to cubic crystalline phase, which are in agreement with the XRD results (inset of Fig. 1A(b), 2B(e) and 2C(h)). In addition, the existence of detectable diffraction rings in the selected-area electron diffraction (SAED) pattern of three hierarchical nano- La_2O_3 (inset of Fig. 1A(b), 2B(e) and 2C(h)) further reveal the formation of polycrystalline products. These results further confirm the formation of the hierarchical nano- La_2O_3 .

The growth mechanism of the nanostructures is investigated by FESEM and TEM in time-dependent experiments. The results

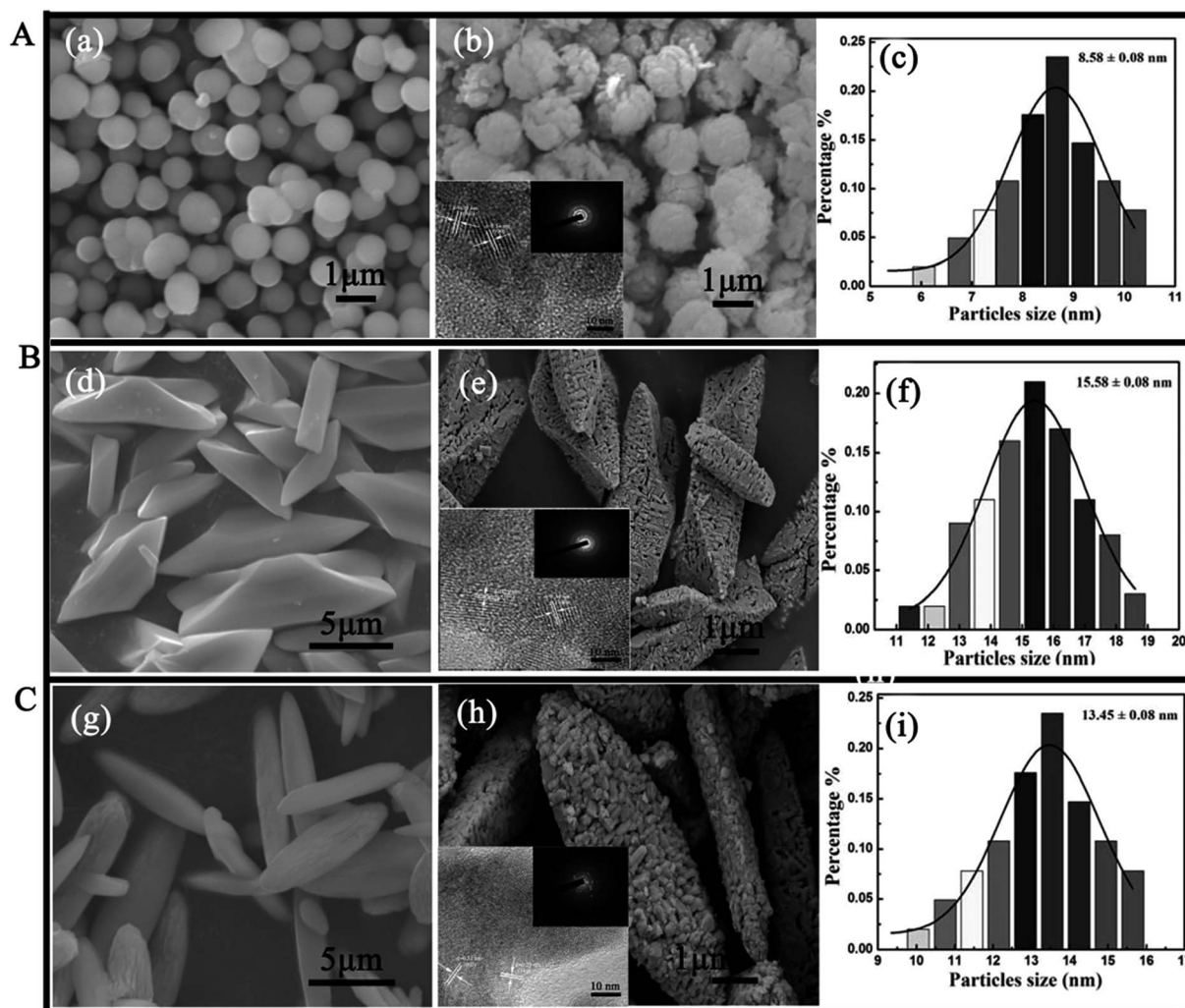


Fig. 1 SEM and particle size distribution of La_2O_3 and LaOHCO_3 . A(a) Spherical LaOHCO_3 ; A(b) spherical La_2O_3 (inset: lattice fringes and SAED patterns); A(c) size distribution analysis of hierarchically spherical nano- La_2O_3 ; B(d) polyhedral LaOHCO_3 ; B(e) polyhedral La_2O_3 (Inset: lattice fringes and SAED patterns); B(f) Size distribution analysis of hierarchically polyhedral nano- La_2O_3 ; C(g) spindles-like LaOHCO_3 ; C(h) spindles-like La_2O_3 (inset: lattice fringes and SAED patterns); C(i) Size distribution analysis of hierarchically spindles-like nano- La_2O_3 .



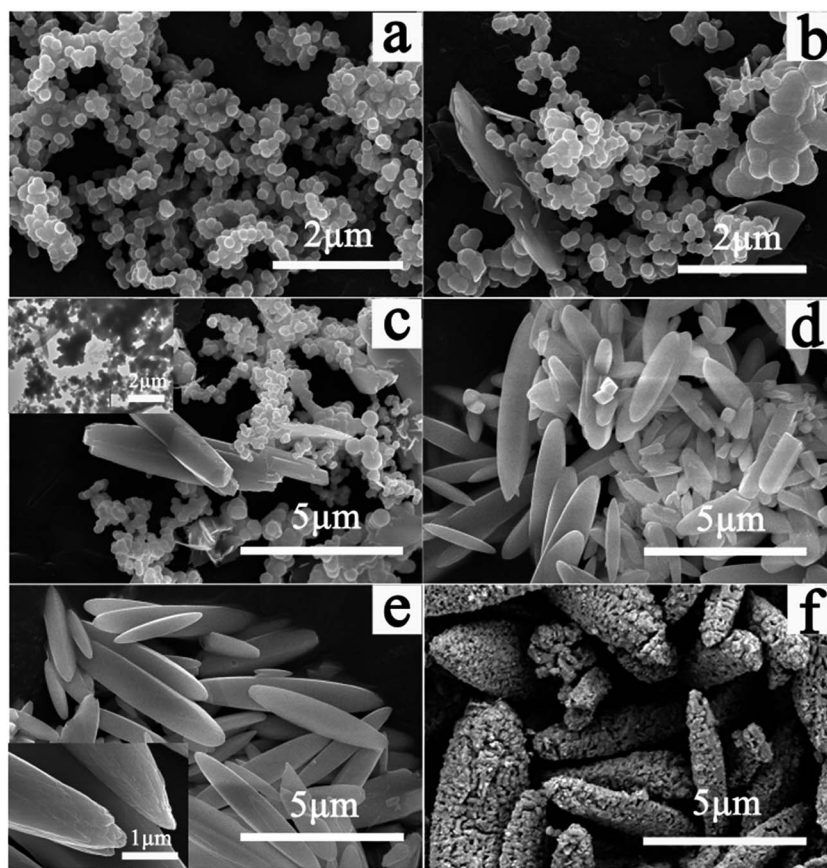
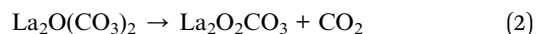
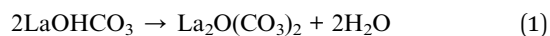


Fig. 2 The growth mechanisms of spindles-like La_2O_3 within different points-in-time: (a) 0.5 h; (b) 1 h; (c) 2 h (inset: TEM); (d) 3 h; (e) 4 h; (f) precursors calcined from room temperature to 800 °C for 4 h.

reveal that three different morphologies of La_2O_3 samples, including polyhedral, spherical and spindles-like, are initially derived from spherical morphology, which show similar growth mechanisms to each other. We take spindles-like La_2O_3 for example to explain the growth mechanism in details (shown in Fig. 2). Initially, the $\text{La}(\text{OH})\text{CO}_3$ precursors of spherical morphology are formed within 0.5 hour reaction time (Fig. 2a), similarly to the formations of polyhedral and spherical La_2O_3 samples. Then, more $\text{La}(\text{OH})\text{CO}_3$ precursor products of spherical morphology assemble to form spindles-like morphology *via* oriented attachment within 1–3 hours (Fig. 2b–e). Meanwhile, non hollow structures are observed by characterization of TEM (inset of Fig. 2c). Then, we find spindles-like $\text{La}(\text{OH})\text{CO}_3$ precursors consist of small particles (inset of Fig. 2e). Finally, the $\text{La}(\text{OH})\text{CO}_3$ precursors are calcined from room temperature to 800 °C and maintained at this temperature for 4 h to form La_2O_3 samples with micro/nanostructures (Fig. 2f). The growth mechanisms of polyhedral and spherical products are also researched in time-dependent experiments. The similar growth mechanisms are observed.

XRD patterns of three samples are very similar, one of which is depicted in Fig. 3A as a representation. From Fig. 3A, a strong intensity peak is detected at a diffraction angle of 30°, which is assigned to (101) plane of La_2O_3 . The other five peaks are assigned to (100), (002), (102), (110) and (103) lattice planes

belonging to cubic crystalline phase of La_2O_3 (JCPDS no. 26-0815). The results indicate that the pure La_2O_3 crystals are successfully prepared by the simple method. Representative FTIR spectra of spherical La_2O_3 and $\text{La}(\text{OH})\text{CO}_3$ precursor are shown in Fig. 3B. In the spectrum of $\text{La}(\text{OH})\text{CO}_3$, the bands at 1486 and 1412 cm^{-1} are attributed to the stretching vibration mode of the C–O bond, and the flexural vibration of CO_3^{2-} appear at 859 cm^{-1} , 720 cm^{-1} , 695 cm^{-1} . In the spectrum of La_2O_3 , the La–O stretching vibration is at about 653 cm^{-1} , showing La_2O_3 is prepared successfully. The thermogravimetric analysis (TGA) and differential temperature analysis (DTA) traces disclosed the formation process of La_2O_3 from pyrolysis of the $\text{La}(\text{OH})\text{CO}_3$ precursor. The TGA curves of the as-prepared $\text{La}(\text{OH})\text{CO}_3$ are shown in Fig. 3C. It can be noticed that there are three weight loss steps around 250 °C to 300 °C, 450 °C to 500 °C and 700 to 790 °C. For $\text{La}(\text{OH})\text{CO}_3$, the weight loss between 250 °C to 300 °C could be attributed to the removal of water, accompanying the formation of $\text{La}_2\text{O}(\text{CO}_3)_2$ (see reaction (1)). Then $\text{La}_2\text{O}(\text{CO}_3)_2$ decomposes to form La_2O_3 and release CO_2 , approximately to lose weight 28% (see reaction (2) and (3))



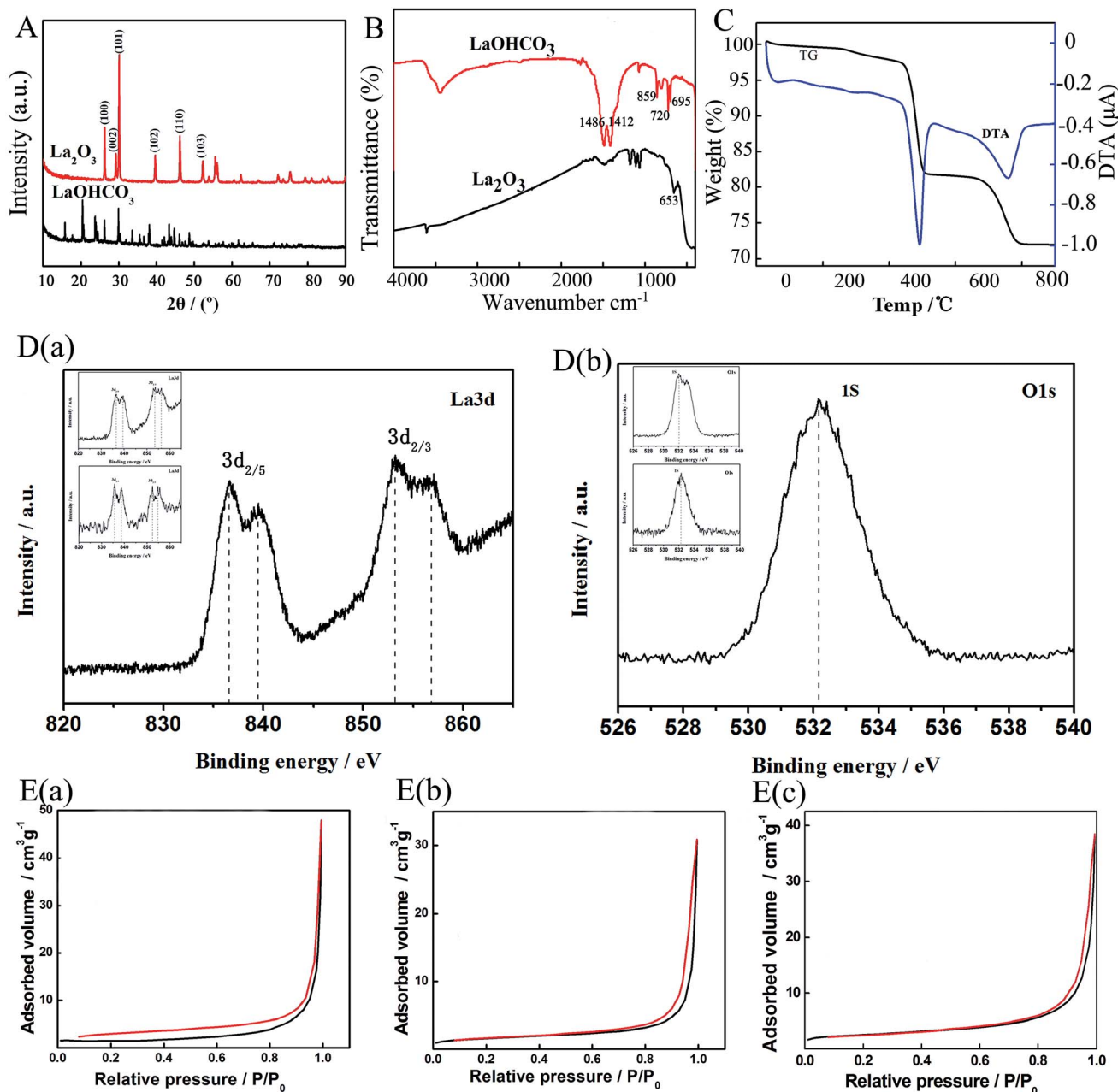


Fig. 3 (A) XRD patterns of polyhedral La_2O_3 and LaOHCO_3 . The top red line depicts the diffraction pattern of La_2O_3 nanocrystals, whereas the lower black line corresponds to LaOHCO_3 nanocrystals; (B) FTIR of the spindles La_2O_3 and LaOHCO_3 ; (C) the thermogravimetric (TG) and DTA of LaOHCO_3 ; (D) XPS spectra of (a) La 3d, (b) O 1s; (E) N_2 -sorption isotherms and pore size distribution curves (inset) for different morphology micro-nano La_2O_3 . (a) Spherical La_2O_3 , (b) polyhedral La_2O_3 , (c) spindle-like La_2O_3 .

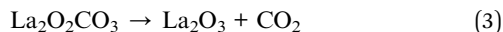


Fig. 3D presents the X-ray photoelectron spectroscopies of La_2O_3 . These spectra exhibit characteristic La 3d and O 1s peaks. As indicated in Fig. 3D(a), the La 3d spectrum shows two peaks at the binding energy 839.48 and 856.77 eV, corresponding to La 3d_{5/2} and 3d_{3/2}, respectively. Meanwhile, the peaks located at a binding energy of around 528.9 and 531.65 eV are both corresponding to the La_2O_3 and H_2O spectrum (Fig. 3D(b)). These results are in accordance with the previous reports on La_2O_3 .¹⁹

Brunauer–Emmett–Teller (BET) gas adsorption measurements were used to characterize pore volume, pore diameter and specific surface area (see Fig. 3E). The N_2 adsorption/desorption isotherms of the synthesized La_2O_3 exhibit typical type-IV hysteresis, indicating the presence of pores (Fig. 3E). The samples are mainly mesoporous (inset of Fig. 3E). The BET surface area is measured to be $18.76 \text{ m}^2 \text{ g}^{-1}$ for spherical La_2O_3 , $12.43 \text{ m}^2 \text{ g}^{-1}$ for polyhedral La_2O_3 and $8.14 \text{ m}^2 \text{ g}^{-1}$ for polyhedral La_2O_3 , respectively.

To determine absorption capacity of La_2O_3 micro-nanostructures for phosphate, ascorbic acid and molybdate



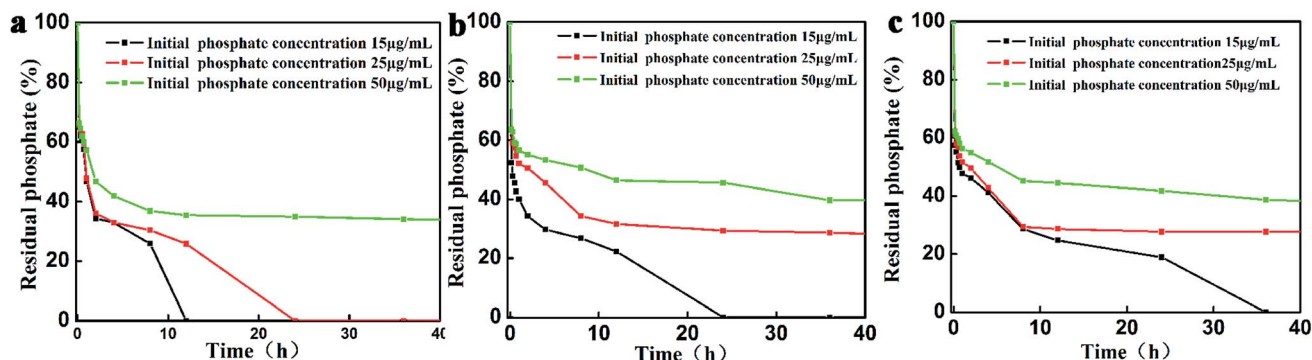


Fig. 4 Phosphate absorption capacity of different morphology micro-nano La_2O_3 . (a) Spherical La_2O_3 , (b) spindle-like La_2O_3 , (c) polyhedral La_2O_3 . Black wire, red wire and green wire: initial phosphate concentration is 15 $\mu\text{g mL}^{-1}$, 25 $\mu\text{g mL}^{-1}$ and 50 $\mu\text{g mL}^{-1}$ respectively.

solution were employed to measure phosphate group. The initial concentrations of phosphate were 15 $\mu\text{g mL}^{-1}$, 25 $\mu\text{g mL}^{-1}$ and 50 $\mu\text{g mL}^{-1}$, respectively. As shown in Fig. 4, the residual phosphate reduce when La_2O_3 was added. It is worth highlighting that phosphate is absolutely removed by spherical La_2O_3 for only 12 h (Fig. 4a black wire) when initial phosphate concentration is 15 $\mu\text{g mL}^{-1}$, while 24 h for spindles-like micro-nano La_2O_3 (Fig. 4b black wire), 35 h for polyhedral La_2O_3 (Fig. 4c black wire). Moreover, when initial phosphate concentration is 25 $\mu\text{g mL}^{-1}$ (red wire), phosphate removal have been accomplished completely by spherical La_2O_3 after 24 h, but accomplished 68% by spindle-like La_2O_3 and 78% by polyhedral La_2O_3 in the meantime. Likewise, superior removal capacity of

spherical La_2O_3 is observed, when initial phosphate concentration is 50 $\mu\text{g mL}^{-1}$. The results show that spherical La_2O_3 has best phosphate removal capacity from water, which attributes to that the spherical La_2O_3 has the smallest particle sizes and the largest BET surface area.

To investigate the effect of phosphate addition and phosphate starvation on microorganism growth, the growth curve of *Escherichia coli* (a Gram-negative bacteria) and *Staphylococcus aureus* (a Gram-positive bacteria) were analyzed. Meanwhile, residual phosphate was calculated after microorganism grows for 8 h. As shown in Fig. 5a, the phosphate addition is beneficial to microorganisms growth (both Gram-positive and Gram-negative bacteria) (Fig. 5a and b). It is about 32 times and 26

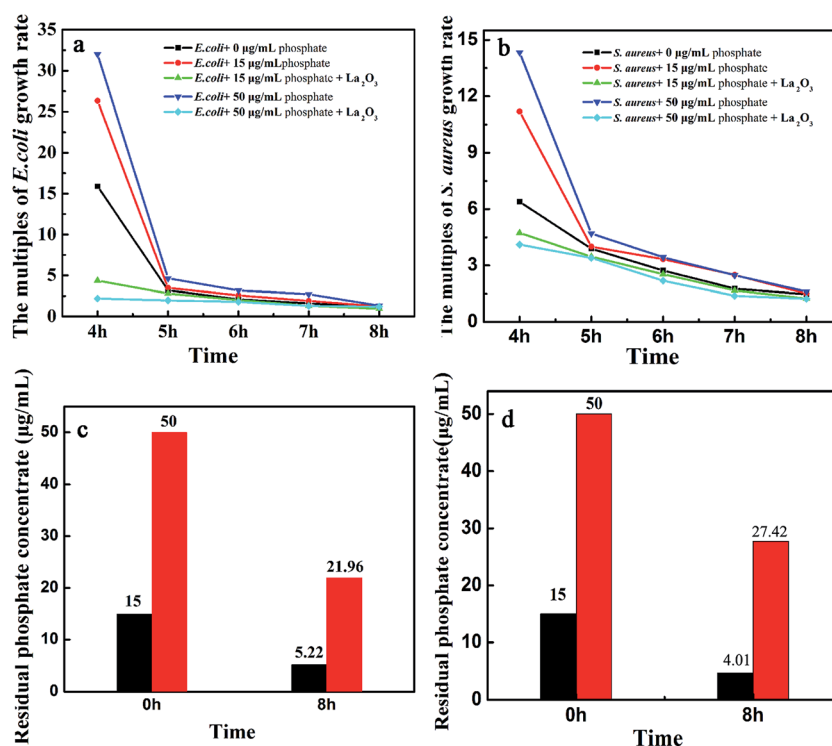


Fig. 5 Antibacterial activity (*Escherichia coli* and *Staphylococcus aureus*) and phosphate adsorption of spherical La_2O_3 (a, c) *Escherichia coli*, (b, d) *Staphylococcus aureus*.



times larger than the growth rate of *E. coli* in 2 h when adding phosphate ($15 \mu\text{g mL}^{-1}$, red line and $50 \mu\text{g mL}^{-1}$, blue line), showing that phosphate addition to minimal medium could be beneficial to growth of *Escherichia coli* (Fig. 5a) and *Staphylococcus aureus* (Fig. 5b) when growth less than 5 h. However, the benefit of phosphate is suppressed following the addition of La_2O_3 . The growth rates of *Escherichia coli* (Gram-negative bacteria) and *Staphylococcus aureus* (Gram-positive bacteria) were drawn in Fig. 5a–d. From Fig. 5a, it is only 5 and 2.5 times larger than the growth rate of *E. coli* in 2 h when adding La_2O_3 to medium in 4 h although the existence of phosphate ($15 \mu\text{g mL}^{-1}$ and $50 \mu\text{g mL}^{-1}$). The results show that the growth rates of *E. coli* and *S. aureus* are obviously limited within 5 h when La_2O_3 is added, attributed to effective binding of La_2O_3 to phosphate, namely that phosphate starvation results in limited microbial growth. Given this, phosphate removal ability capacity of three morphology micro-nano La_2O_3 could be a key point to compare antimicrobial properties of La_2O_3 . Due to superior removal capacity of spherical La_2O_3 , spherical La_2O_3 exhibited better antibacterial property than other two morphologies La_2O_3 . It means that phosphate absorption capacity of La_2O_3 is consistent with the antibacterial property of La_2O_3 .

4 Conclusion

We successfully obtained shape-controlled (spindle, polyhedron, and sphere) micrometer-scaled La_2O_3 , hierarchically assembled by nanoparticles, through a facile, high quality and low prices method. Meanwhile, they have strong removal ability for phosphate and excellent antibacterial property, especially for sphere micro-nano structure La_2O_3 . These hierarchical La_2O_3 micro-nanomaterials may have very good application prospects in water pollution of eutrophication, or reduce excess phosphate in human body in biomedical areas.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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