


 Cite this: *RSC Adv.*, 2017, 7, 23535

Received 15th March 2017

Accepted 14th April 2017

DOI: 10.1039/c7ra03063k

rsc.li/rsc-advances

Polydopamine nanotubes-templated synthesis of TiO₂ and its photocatalytic performance under visible light†

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Herein, we demonstrated that an efficient TiO₂ visible-light photocatalyst can be fabricated by templating mesoporous polydopamine (PDA) nanotubes. Because PDA is a nitrogen-doped carbon source, PDA nanotubes as templates elicited distinct advantages for obtained mesoporous TiO₂. First, they introduced a codopant of carbon and nitrogen into anatase TiO₂ that narrowed its band gap. Second, they could be graphitized in an optimized calcination process to form graphene-like carbon layers on TiO₂ surfaces with high conductivity, which assisted the separation of photogenerated electron–holes of TiO₂. These features endow TiO₂@C-PDA (400-2) excellent photocatalytic performance for the degradation of methyl orange under visible-light irradiation (>420 nm).

1. Introduction

Porous scaffolds can provide catalysts with high accessibility and high diffusion rates of reactant and product. Also, catalysts can be stabilized through encapsulation in hollow nanostructures. Therefore, porous nanostructures have important applications as scaffolds for bio-, electro- and photocatalysts.^{1–5} Porous carbon nanostructures as catalyst carriers have attracted considerable attention because of their good thermal and chemical stability, low cost, large surface area, and versatile fabrication methods. In particular, carbon scaffolds can improve catalyst properties significantly. For example, hybrid TiO₂@carbon has demonstrated promising photocatalytic ability under visible-light irradiation due to a narrowed band gap and accelerated charge transfer in hybrids.^{6–13} The band gap of TiO₂ can be narrowed further by introducing a codopant for the TiO₂ lattice (*e.g.* carbon, nitrogen-codoped TiO₂). Thus, carbon nanomaterials doped with heteroatoms show superior features as scaffolds.^{14–16} Besides the doping effects mentioned above, a carbon scaffold can also modulate the structure and morphology of TiO₂ and further influence its photocatalytic efficiency. Recent studies have shown that a highly conductive carbon layer on TiO₂ is beneficial for fast electron transfer in a photocatalytic process of TiO₂@carbon.^{15,17}

Polydopamine (PDA) is a promising precursor for nitrogen-doped carbon nanomaterials.^{18–20} The reactivity of functional groups on PDA gives it high absorbency of metal ions, which can be transformed further to useful metal oxides or metal

nanoparticles depending on the conditions of carbonization or calcination. Previously, we reported successful fabrication of tubular PDA nanostructures,²¹ which could be templates for the production of various catalysts. Because tubular TiO₂ has demonstrated very high efficiency as flow-through photocatalytic membranes,²² and has shown efficient solar water splitting after doping with carbon,⁷ we investigated the fabrication and properties of tubular TiO₂ *via* templating PDA nanotubes (TiO₂@C-PDA). Using it as a proof of concept, we hoped to reveal what effects PDA templates imparted for TiO₂ as a photocatalyst. We observed that the photocatalytic ability of TiO₂ could be improved significantly *via* scaffolding of PDA nanotubes. The latter, as TiO₂-scaffolds, demonstrated versatile advantages through integration of morphological and structural control, multidoping and easily adjustable graphite content in the carbon layer.

2. Experimental sections

2.1 Materials

Dopamine hydrochloride (purity 98%) was obtained from Sigma-Aldrich. Analytical curcumin, ethanol, hydrochloric acid (12 M), terephthalic acid and titanium(IV) isopropoxide (TTIP) were purchased from Sinopharm Chemical Reagent Beijing Co. Ltd. Tris(hydroxymethyl) aminomethane (Tris)–HCl buffer (1.5 M, pH 8.8) was obtained from Shanghai Double-Helix Biotech. Co. Ltd. All chemicals were used without further purification. Deionized water (Millipore Q >18 MΩ cm at 25 °C) was used to prepare aqueous solutions or as non-solvent.

2.2 Preparation of PDA nanotubes

According to our previous report,²¹ 100 mg curcumin and 500 mg dopamine hydrochloride were dispersed in 100 mL

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra03063k



ethanol under stirring, and then 400 mL deionized water was added. When precipitation was complete, 3 mL (Tris)-HCl buffer solution (1.5 M, pH 8.8) was added in this suspension. After gentle stirring for 24 h, the sediment was separated and cleaned by fresh water thrice, and collected by sedimentation or centrifugation. Then, the sediment was soaked in ethanol to dissolve curcumin crystals. When curcumin crystals were removed completely, the sediment was washed by deionized water and freeze-dried to obtain PDA nanotubes.

2.3 Synthesis of TiO₂@C-PDA samples

Titanium(IV) isopropoxide (0.1 g) was mixed with HCl (0.3 g, 12 M) under vigorous stirring for 5 min. Then, PDA nanotubes (0.1 g) were immersed in this mixture under stirring. Intensive stirring for 10 min enabled full absorption of titanium sol by PDA nanotubes. Then, the “mud-like” mixture was aged in a closed container at 70 °C for 24 h to further hydrolyze TTIP. The aged product was Ti(IV)-PDA gel. To carbonize PDA and crystallize TiO₂, Ti(IV)-PDA gel was heated further in a tube furnace from room temperature to 500 °C at a heating rate of 5 °C min⁻¹ under a continuous flow of argon (flow rate: 50 mL min⁻¹), maintained at 500 °C for 3 h, and cooled to room temperature. The obtained product was named TiO₂@PDA. Then, in air, the TiO₂@PDA powder was heated in a muffle furnace from room temperature to 400 °C under a heating rate of 5 °C min⁻¹ and maintained at 400 °C for 1, 2 and 3 h, respectively. Calcinated products were ground before further characterizations and experiments. Obtained samples were denoted as TiO₂@C-PDA (400-1), TiO₂@C-PDA (400-2) and TiO₂@C-PDA (400-3) based on the temperature and duration of calcination. Besides, pristine TiO₂ was prepared by directly heating the HCl-hydrolyzed TTIP gel at 500 °C for 3 h in a muffle furnace in air.

2.4 Characterization and measurements

Morphological characterizations were conducted through scanning electron microscopy (SEM; SU8010, Hitachi) at 15 kV equipped with an energy-dispersive spectroscopy (EDS) detector. Transmission electron microscopy (TEM) was operated on an H-7650B device (Hitachi) at an accelerating voltage of 80 kV. High-resolution transmission electron microscopy (HRTEM) images were obtained on a FEI G2 F20 microscope (Tecnai) with an accelerating voltage of 200 kV.

The chemical states and species of elements were verified using an X-ray photoelectron spectrometer (Axis Ultra; Kratos Analytical) with monochromatic Al K α ($h\nu = 1486.7$ eV) radiation as the excitation and at an X-ray power of 150 W. All spectra were calibrated using the C 1s peak (284.7 eV). Fourier transform infrared (FTIR) spectroscopy was done using a Tensor 27 FTIR spectrophotometer (Bruker). Crystallization characterizations of calcinated products were performed with a XRD-7000 diffractometer (Shimadzu) in reflection mode using a Cu target ($\lambda = 0.15418$ nm) as the incident X-ray. Accelerating voltage of 40 kV, current of 40 mA, and 2θ -step of 0.013° were selected. The scan was collected between 20° and 80°. Raman spectra were acquired by a Raman spectrometer (Xplore PLUS;

Horiba Scientific). The spot size of a 532 nm incident laser for signal collection was about 25 × 25 μ m. Spectra were calibrated using the 520 cm⁻¹ line of a silicon wafer, and the laser beam was focused by an optical microscope with a 10× objective lens, 1200 grooves per mm grating, and 100 μ m slit. UV-visible absorption spectra of TiO₂ and TiO₂@C-PDA samples were collected using a UV-Vis spectrophotometer (U-3900; Hitachi) equipped with a diffuse reflectance accessory, and BaSO₄ powder was used as the reflectance standard. Zeta potential values of TiO₂@C-PDA in aqueous solution at different pH values were measured using a Zetasizer Nano ZS90 (Malvern) at 25 °C. Measurements were performed five times for each sample. Thermogravimetric analysis (TGA) was carried out on a TGA Q50 (TA Instruments) system from room temperature to 800 °C under air at a heating rate of 10 °C min⁻¹. Brunauer-Emmett-Teller (BET) surface area was determined from N₂ adsorption isotherm data collected at 77 K using a Belsorp-Mini II (MicrotracBEL) system. Before testing, ≈ 100 mg samples were dried under a N₂ atmosphere at 300 °C for ≥ 12 h. The specific surface area was calculated using the BET method, and the pore volume and pore-size distribution were obtained using the Barrett-Joyner-Halenda (BJH) model.

2.5 Photocatalytic activity tests

The photocatalytic activity of various samples on degradation of methyl orange (MO) aqueous solution was investigated. A 300 W Xe arc lamp (CHF XM150, Beijing Trust Tech) equipped with a wavelength cutoff filter for $\lambda > 420$ nm and positioned ≈ 8 cm above the aqueous suspension. Briefly, 100 mg of sample was dispersed in 100 mL MO aqueous solution at 10 mg L⁻¹. The solution was stirred for 1 h to ensure establishment of an adsorption-desorption equilibrium between TiO₂ and MO in suspension before irradiation. At a given interval of 30 min, 3 mL of the suspension was sampled, and then the catalyst and MO solution were separated by centrifugation. The concentration of MO in the sample was determined by monitoring absorbance at 461 nm in a UV absorption spectrum with a UV-3600 spectrophotometer (Shimadzu).

Photoluminescence (PL) spectra of TiO₂@C-PDA aqueous suspensions were measured at room temperature using a fluorescence spectrophotometer (F-4600; Hitachi). The excitation wavelength was 420 nm.

To test the amount of hydroxyl radical (\cdot OH) generated over TiO₂@C-PDA photocatalysts under visible-light irradiation, 40 mg of tested samples was suspended in 80 mL of aqueous solution containing NaOH (0.01 M) and terephthalic acid (3 mM).²³ Before exposure to visible-light irradiation, the solution was stirred for 1 h in the dark to ensure establishment of an adsorption-desorption equilibrium. Then, the mixture was illuminated with a 300 W Xe arc lamp (CHF XM150, Beijing Trust Tech) equipped with a wavelength cut-off filter of $\lambda > 420$ nm. During illumination, at 30 min intervals, 3 mL of the mixture was removed and centrifuged. The fluorescence spectra of obtained supernatants were recorded at room temperature using a fluorescence spectrophotometer (F-4600; Hitachi) with a Xe lamp as the excitation source. The intensity of the emission



peak at 426 nm of 2-hydroxyterephthalic acid (excitation, 315 nm) was used to evaluate the ability of photocatalysts to produce $\cdot\text{OH}$ radicals under light irradiation.

3. Results and discussion

3.1 Morphology and phase structure

Fig. 1 demonstrates images of original PDA nanotubes (Fig. 1a and d) as well as TiO_2 @C-PDA nanotubes before (Fig. 1b and e) and after calcination (Fig. 1c and f). The tubular morphology of original PDA nanotubes was maintained in TiO_2 @C-PDA even after calcination in air for several hours (Fig. 1c and f), whereas the carbon skeleton was almost “burned out”, leaving TiO_2 as the main component in nanotubes. The average length of original PDA nanotubes was more than several tens of micrometers, but it decreased to several micrometers in hybrids after calcination. The average diameters of original PDA nanotubes, TiO_2 @PDA nanotubes and TiO_2 @C-PDA nanotubes were 465 ± 40 , 366 ± 57 , and 210 ± 30 nm, respectively. The thicknesses of tube walls for these PDA, TiO_2 @PDA and TiO_2 @C-PDA nanotubes were 107 ± 19 , 99 ± 17 , and 56 ± 12 nm, respectively, suggesting considerable shrinkage of TiO_2 @PDA after calcination in air due to TiO_2 crystallization and burning out of carbon scaffolds. Element mapping of TiO_2 @C-PDA showed the smooth distribution of carbon, nitrogen on a single TiO_2 nanotube (Fig. 1g–j).

To remove extra amorphous carbon in TiO_2 @PDA, calcination in air was conducted at 400°C for 1, 2 and 3 h, respectively. Fig. 2 shows the TEM and HRTEM images of TiO_2 @C-PDA (400-2). Fig. 2a and b show the typical morphologies of a calcinated TiO_2 @C-PDA (400-2) nanostructure. The tube wall was mesoporous and composed of dark domains (Fig. 2b). All dark domain nanoparticles were well-crystallized (Fig. 2c), and the size distribution of domains was narrowed. The layer distance

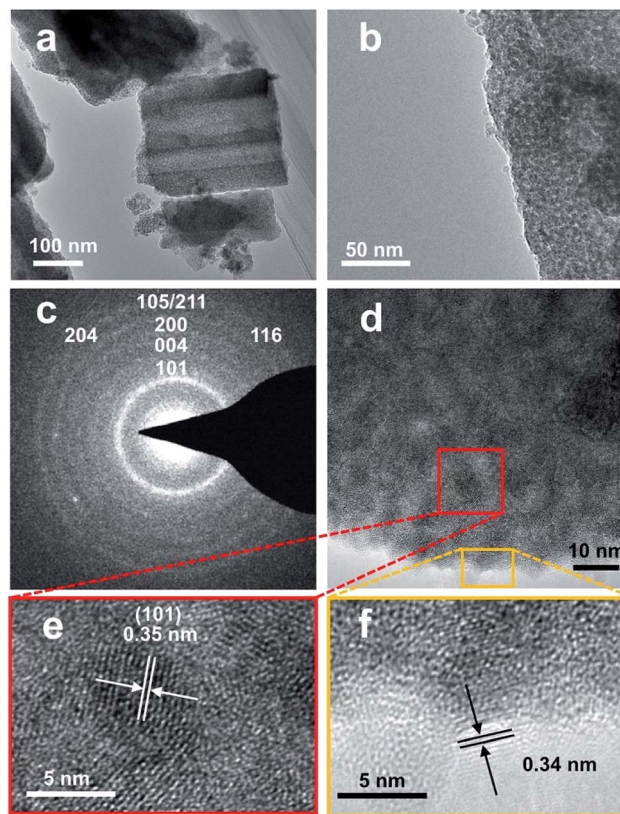


Fig. 2 (a and b) TEM images of TiO_2 @C-PDA (400-2), (c) SAED pattern of a TiO_2 crystal, (d–f) HRTEM images of TiO_2 @C-PDA (400-2). Lattice fringes of anatase (e) and graphene-like carbon coverings (f).

of an identical crystal was 0.35 nm, which is a typical feature of an anatase phase (101) lattice (Fig. 2e). Dark domains were TiO_2 nanocrystals (Fig. 2c), whereas the light part around the TiO_2 domain was a carbon layer (Fig. 2d). The carbon layer covering TiO_2 nanocrystals was thick and amorphous in TiO_2 @C-PDA (400-1) (Fig. S1†) but was graphene-like after longer burning. Also, 0.34 nm was the distance between graphite (002) layers (Fig. 2f). The graphitization effect in calcination for carbonized PDA could be clearly observed even without TiO_2 . Fig. 3 shows HRTEM images of the residue of carbonized PDA nanotubes after calcination in air at 400°C for 2 h. Graphite-like ordered carbon layers were concentrated in this residue. The measured lattice spacing of 0.18 nm corresponded to the (102) facet of graphitic carbon,²⁴ and that of 0.21 nm belonged to the (1100)

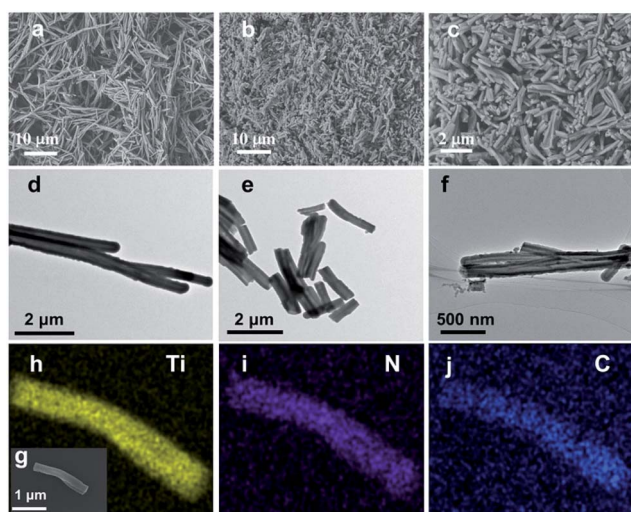


Fig. 1 (a and d) SEM and TEM images of long PDA nanotubes. (b and e) SEM and TEM images of TiO_2 @PDA nanotubes. (c and f) SEM and TEM images of TiO_2 @C-PDA (400-2). (g) SEM image of a TiO_2 @C-PDA nanotube. Also shown are images of titanium (h), nitrogen (i) and carbon (j) mapping.

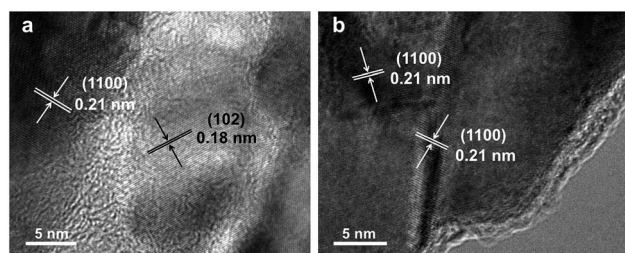


Fig. 3 HRTEM images (a and b) of the residue of carbonized-PDA after calcination in air for 2 h at 400°C .



in-plane lattice fringes of graphene.^{25,26} Carbon with different degrees of graphitization shows different reactivity in burning; the one with a higher degree of graphitization survives longer during burning than its amorphous counterpart. Therefore, calcination in our experiments was not only a simple step to remove extra carbon, but also an optimal process to select a conductive graphite layer remaining on TiO₂ nanoparticles. The presence of a graphite layer on TiO₂@C-PDA (400-2) may be favorable for the efficient separation of photogenerated electron-hole pairs on TiO₂.¹⁷

Interfacial interactions through chelated- and bidentate-binding modes²⁷ and electron transfer in catechol/TiO₂-anatase nanostructures have been confirmed.^{28,29} The advantage of PDA nanotubes as scaffolds for TiO₂ is their inherent ability to chelate with Ti(IV) and TiO₂.^{30–32} The strong interaction between Ti(IV) and the catechol groups of PDA promotes complete and tight contact between a carbon layer and TiO₂ (Scheme 1). Further structural characterizations of TiO₂@C-PDA were conducted. Fig. 4 presents X-ray diffraction (XRD) patterns of these hybrids. All samples showed pure anatase features that were in agreement with HRTEM observations. The (101) diffraction peaks in TiO₂@C-PDA (400-1) and TiO₂@C-PDA (400-2) showed clear shifts to smaller degrees as compared with that of pure TiO₂. This result indicates expansion of the TiO₂ lattice by introduction of foreign atoms after calcination. No obvious lattice expansion was observed in TiO₂@C-PDA (400-3), suggesting that a prolonged calcination time decreases the content of foreign species in the host TiO₂ lattice. The corresponding crystal sizes in different samples were calculated through application of the Scherrer equation to the peak at 25.3° (Table 1). The crystallite size decreased from 18.3 nm in pure TiO₂ to 7.7 nm in TiO₂@C-PDA (400-3). The precursor of Ti(IV) was absorbed and locked in these micropores on PDA tube walls. This spatial confinement was maintained even in its transformation to TiO₂ crystals, which is similar to when TiO₂ is generated in micelles.³³ The bidentate bonding of

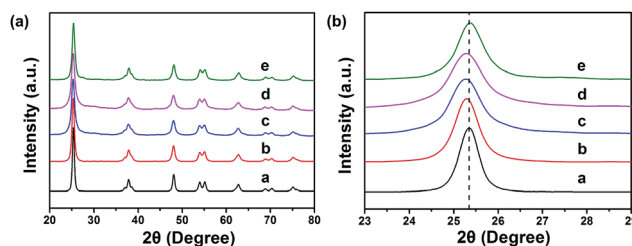


Fig. 4 (a) XRD patterns of TiO₂ and TiO₂@C-PDA. (a) TiO₂, (b) TiO₂@PDA, (c) TiO₂@C-PDA (400-1), (d) TiO₂@C-PDA (400-2), (e) TiO₂@C-PDA (400-3). (b) Partly magnified XRD patterns of (a). XRD patterns exhibit the diffraction peaks of anatase TiO₂.

Table 1 Various structural parameters of TiO₂@C-PDA samples

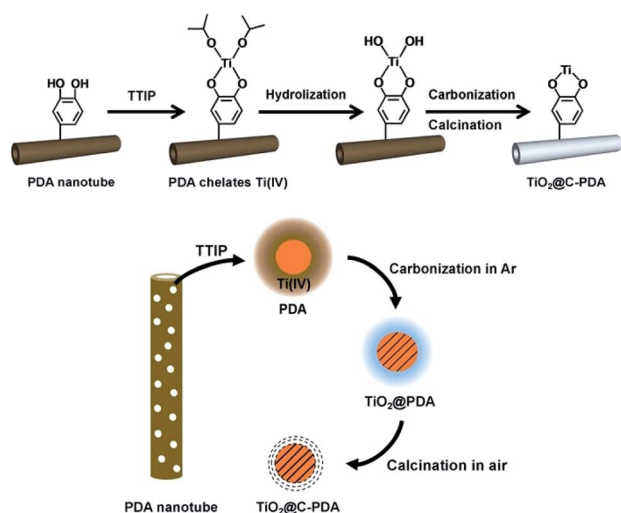
Samples name	Crystal size (nm)	BET surface area (m ² g ⁻¹)	Pore size (nm)	Band gap (eV)
Pristine TiO ₂	18.3	74.9	21.5	3.17
TiO ₂ @C-PDA (400-1)	10.7	81.8	5.63	2.49
TiO ₂ @C-PDA (400-2)	8.3	70.7	5.95	2.57
TiO ₂ @C-PDA (400-3)	7.7	68.0	7.02	2.85

catechol groups in PDA to TiO₆²⁻ has been hypothesized to also contribute to the size and crystal control, which is similar in the case of SO₄²⁻.³⁴ Because of this feature as well as inherent functional groups and microporous nanostructures, PDA nanotubes show better size-control compared with that of TiO₂@C core-shell composites.³⁵ No phase transformation from anatase to rutile occurred until the calcination temperature was increased to 800 °C, suggesting an extra-stabilized TiO₂ crystal (Fig. S2†).

Raman spectroscopy can provide invaluable contributions towards understanding the material structure and its influence on the functional properties. In anatase TiO₂ (Fig. 5), Raman peaks corresponding to the E_g mode (143 cm⁻¹) became broader and significantly blue-shifted to 152 cm⁻¹ for TiO₂@C-PDA (400-1), and 147 cm⁻¹ for TiO₂@C-PDA (400-2). This effect is contributed mainly by a change in the crystal size of poorly crystalline materials^{36,37} and change in carrier density.^{17,38} Raman spectroscopy further confirmed the strong surface interaction between the carbon layer and TiO₂ in TiO₂@C-PDA.

3.2 Elemental analyses and optical properties

To investigate the composition and chemical state of the elements in TiO₂@C-PDA after calcination, X-ray photoelectron spectroscopy (XPS) characterizations were conducted for all samples (Fig. 6, S3 and Table S1†). Calcination effectively removed large amounts of carbon in TiO₂@PDA. Meanwhile, the amount of nitrogen was also sacrificed in calcination, from 8.93% in TiO₂@PDA down to 1.54% in TiO₂@C-PDA (400-1) (Table S1†). In the high-resolution XPS spectra of the N 1s core-level for three samples (Fig. 6a, c and e), two peaks, one located at 398.6 eV, and the other at 400.0 eV, were identified. The weak peak located at 398.6 eV was attributed to the N species in the



Scheme 1 Fabrication of TiO₂@C-PDA nanotubes based on PDA nanotubes (schematic).



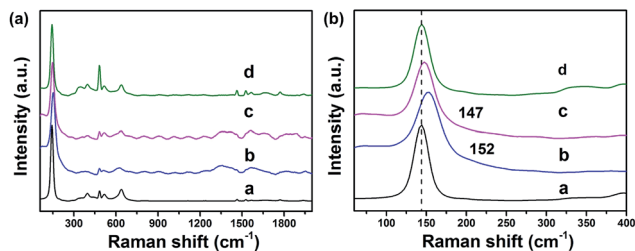


Fig. 5 (a) Raman spectra of TiO_2 and $\text{TiO}_2\text{@C-PDA}$. (a) TiO_2 , (b) $\text{TiO}_2\text{@C-PDA}$ (400-1), (c) $\text{TiO}_2\text{@C-PDA}$ (400-2), (d) $\text{TiO}_2\text{@C-PDA}$ (400-3). (b) Partly magnified Raman spectra of (a). The original peak at 143 cm^{-1} in TiO_2 was shifted to 152 cm^{-1} and 147 cm^{-1} in $\text{TiO}_2\text{@C-PDA}$ (400-1) and $\text{TiO}_2\text{@C-PDA}$ (400-2), respectively.

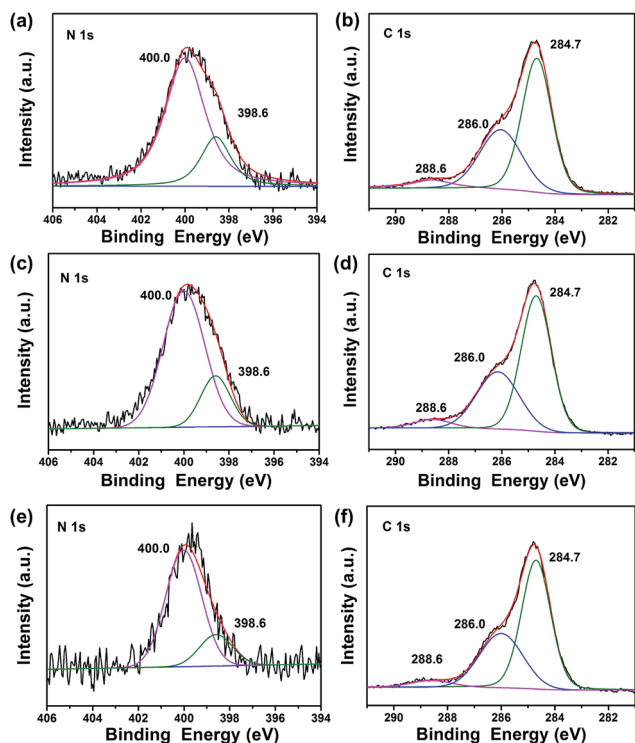


Fig. 6 (a) N 1s XPS spectra of $\text{TiO}_2\text{@C-PDA}$ (400-1), (b) C 1s XPS spectra of $\text{TiO}_2\text{@C-PDA}$ (400-1). (c) N 1s XPS spectra of $\text{TiO}_2\text{@C-PDA}$ (400-2), (d) C 1s XPS spectra of $\text{TiO}_2\text{@C-PDA}$ (400-2). (e) N 1s XPS spectra of $\text{TiO}_2\text{@C-PDA}$ (400-3), (f) C 1s XPS spectra of $\text{TiO}_2\text{@C-PDA}$ (400-3).

$\text{C}=\text{N}$ form from carbonized PDA,³⁹ which is present on the surface of TiO_2 . The strong N 1s peak around 400.0 eV , widely observed in samples prepared *via* wet processes,^{40–43} could be assigned to interstitial N species bonding with the lattice O in the host TiO_2 . On the basis of the relative XPS area, the contents of the interstitial N species in $\text{TiO}_2\text{@C-PDA}$ (400-1), (400-2) and (400-3) were estimated to be 1.2, 0.98, and 0.49%, respectively. Given the relatively small space of the interstitial site in the host TiO_2 , occupation of the N species at the interstitial site would induce an expansion of the host TiO_2 lattice, a notion that was well supported by XRD measurement given above.

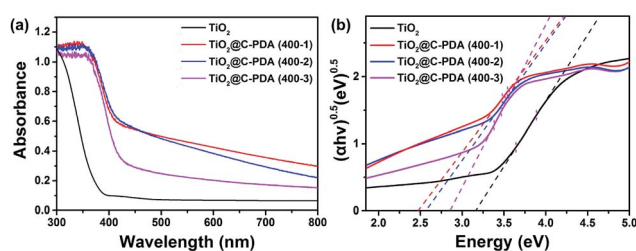


Fig. 7 (a) UV-vis diffuse reflectance spectra of TiO_2 and $\text{TiO}_2\text{@C-PDA}$ (400-1, 2, 3). (b) Band-gap evaluation from Tauc plots of TiO_2 and $\text{TiO}_2\text{@C-PDA}$ (400-1, 2, 3).

As shown in Fig. 6b, d and f, the high-resolution XPS spectra of C 1s for all samples consisted of three peaks located at 284.7 , 286.0 , and 288.6 eV , respectively. In combination with the HRTEM measurements given above, we believe that the main peak at 284.7 eV originated from both the surface graphene layer and adventitious carbon in XPS. The amount of surface carbon was calculated further based on the weight loss of samples in air (Fig. S4†). The carbon covered on TiO_2 was 3.8, 1.6 and 1.1 wt% for $\text{TiO}_2\text{@C-PDA}$ (400-1), $\text{TiO}_2\text{@C-PDA}$ (400-2) and $\text{TiO}_2\text{@C-PDA}$ (400-3), respectively. The contribution at 286.0 eV was due to the C–N of carbonized PDA.³⁹ The peak at 288.6 eV indicated formation of the Ti–O–C bond⁴⁴ most likely by substitution of the titanium atom instead of the oxygen atom in the host TiO_2 lattice. On the basis of the relative XPS area, the doping amounts of carbon in $\text{TiO}_2\text{@C-PDA}$ (400-1), $\text{TiO}_2\text{@C-PDA}$ (400-2) and $\text{TiO}_2\text{@C-PDA}$ (400-3) were 1.45, 0.84 and 0.82%, respectively. The content of the doped C species at the substitutional site and the N species at the interstitial site in TiO_2 lattice decreased with increasing calcination time, whereas the amount of change of carbon cover was in accordance with them.

Fig. 7a presents the UV-vis diffuse reflectance spectra of various $\text{TiO}_2\text{@C-PDA}$ samples compared with pristine TiO_2 without using PDA nanotubes as templates. From the Tauc plot (Fig. 7b), the corresponding band gap values of $\text{TiO}_2\text{@C-PDA}$ (400-1), (400-2) and (400-3) were estimated to be 2.49, 2.57, and 2.85 eV, respectively, as compared with the band gap (3.17 eV) of pure TiO_2 . Earlier investigations indicated that doping N at an interstitial site or carbon at a titanium site in a TiO_2 lattice induces a localized level above the valence band (VB) or below the conduction band (CB) in the forbidden band, respectively, leading to extension of the host TiO_2 's absorption to the visible-light region.^{45–47} Thus, both the interstitial N and substitutional C dopants in the host TiO_2 , as evidenced by the XPS measurements given above, were responsible for the observed reduction in the band gap.

In addition, the surface area, pore size and surface charge of $\text{TiO}_2\text{@C-PDA}$ samples were measured (Fig. S5 and S6†). The surface charge of these samples showed no clear difference from that of pure TiO_2 (Fig. S6†). The crystal sizes, BET surface area, pore size and band gaps of all samples are summarized in Table 1.

3.3 Photocatalytic performance of $\text{TiO}_2\text{@C-PDA}$ nanotubes

The photodegradation of MO was used to evaluate the photocatalytic activity of $\text{TiO}_2\text{@C-PDA}$ samples under visible light



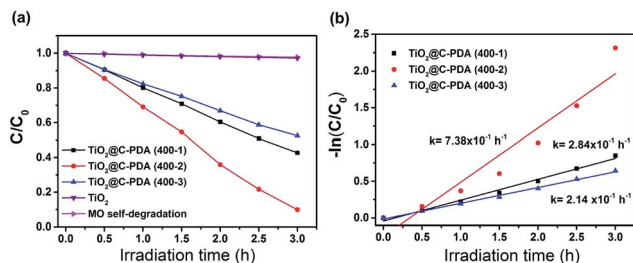


Fig. 8 (a) Photocatalytic degradation of MO in the presence of TiO₂ and TiO₂@C-PDA (400-1, 2, 3) under visible-light irradiation ($\lambda > 420$ nm). (b) Linear extrapolations of TiO₂@C-PDA (400-1, 2, 3) in photocatalytic degradation.

(>420 nm) (Fig. 8 and S7†). TiO₂@C-PDA (400-2) showed remarkable photocatalytic activity, only 10% MO remained after 3 h. TiO₂@C-PDA (400-1) and TiO₂@C-PDA (400-3) had similar photocatalytic activities, leaving $\approx 43\%$ and $\approx 53\%$ MO in reaction systems after 3 h of irradiation under visible light (>420 nm). The photocatalytic oxidation of MO was a pseudo-zero-order reaction. Its apparent reaction rate constant was derived from the linear slope of the relationship between $\ln(c/c_0)$ and kt , where c_0 and c are the concentrations of initial solution and after t (h) of irradiation, respectively. TiO₂@C-PDA (400-2) had the fastest reaction rate constant (0.738 h^{-1}), whereas the constants of TiO₂@C-PDA (400-1) and TiO₂@C-PDA (400-3) were 0.284 h^{-1} and 0.214 h^{-1} , respectively. The photocatalytic activity of TiO₂@C-PDA (400-2) did not decrease after four cycles of photodegradation, demonstrating the good stability of this catalyst (Fig. S8†). A comparison of XRD patterns of this photocatalyst before and after photodegradation experiments also supported this finding (Fig. S9†).

The photocatalytic performance is strongly dependent on the extent of separation of electron-hole pairs over the excited semiconductor photocatalyst. Investigations have shown that a graphene-like carbon layer can promote remarkable separation of the electron-hole pairs over the excited TiO₂ through an electron transfer from the CB to the graphene-like carbon layer. According to the HRTEM and XPS results shown above,

graphene-like carbon layers were obviously present in TiO₂@C-PDA (400-2) rather than in other samples. The effect of the graphene-like carbon layer in TiO₂@C-PDA (400-2) on separation of the photoexcited electron-hole pairs was further evidenced by PL spectra (Fig. 9). The intensity of PL emission of TiO₂@C-PDA (400-2) was decreased significantly compared with those of TiO₂@C-PDA (400-1) and (400-3), suggesting efficient separation of the electron-hole pairs in the presence of the graphene-like carbon layer. Unsurprisingly, TiO₂@C-PDA (400-2) exhibited the highest photocatalytic activity for MO degradation among these samples.

Recently, an isotopic labeling study showed that the hole-driven oxidation of H₂O in aqueous TiO₂ suspensions is the main source of the free $\cdot\text{OH}$ radical,⁴⁸ which plays an important part during the degradation of water pollutants in illuminated TiO₂ photocatalysts,⁴⁹ such as in case of MO degradation.⁵⁰ Therefore, the rate of formation of $\cdot\text{OH}$ radicals is an important parameter to understand the reaction mechanism of illuminated photocatalysts. Quantitative detection of $\cdot\text{OH}$ radicals of TiO₂@C-PDA under visible-light irradiation was conducted to investigate its photocatalytic mechanism. Terephthalic acid is an identical probe molecule for measuring $\cdot\text{OH}$ radicals.²³ The change in the fluorescence intensity of supernatant solutions of various illuminated TiO₂ photocatalysts is presented in Fig. 10. The amount of $\cdot\text{OH}$ radicals from TiO₂@C-PDA (400-2) was highest and paralleled their best photocatalytic performance. The higher amount of $\cdot\text{OH}$ radicals from TiO₂@C-PDA (400-2) under visible-light illumination revealed their increased ability to generate holes because of fast separation and transfer of electrons.

The advantage of TiO₂@C-PDA hybrids could be due to: (i) formation of Ti–O–C and interstitial N to vary the lattice structure of TiO₂;⁷ (ii) surface contact of TiO₂ and a carbon layer with a high degree of graphitization.¹⁷ Ti–O–C, similar to Ti–C, can extend light absorption to longer wavelengths.^{11,45} Conversely, electron transfer from a CB of TiO₂ to a carbon layer (as a sensitizer) benefits charge separation.^{51,52} Optimization of charge separation requires a carbon layer on TiO₂ of adequate thickness, better conductivity and close contact. Single-walled

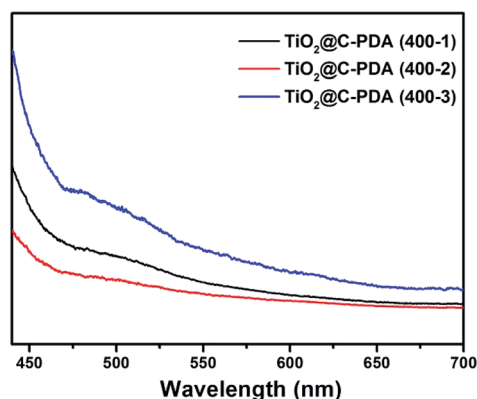


Fig. 9 PL spectra of TiO₂@C-PDA (400-1, 2, 3) using an excitation wavelength of 420 nm.

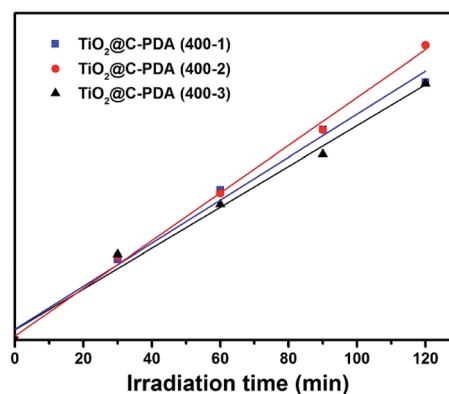


Fig. 10 Time dependence of the fluorescence intensity at 426 nm generated from 2-hydroxyterephthalic acid in the presence of TiO₂@C-PDA (400-1, 2, 3) under visible-light irradiation ($\lambda > 420$ nm).



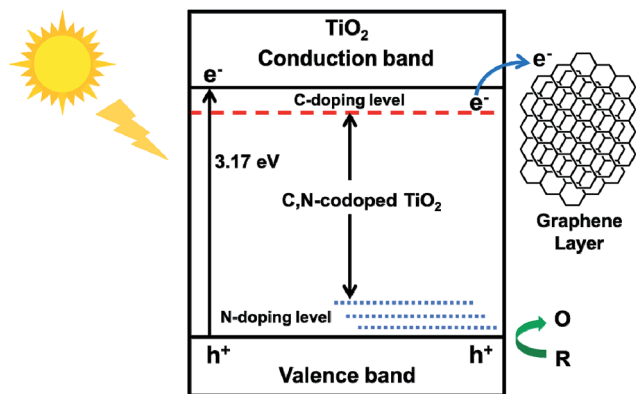


Fig. 11 Electron transition and visible-light photocatalytic process of $\text{TiO}_2\text{@C-PDA}$ (schematic).

carbon nanotubes (SWCNTs) and graphene are highly conductive, so the photoexcitation charges from TiO_2 can be rapidly separated and stabilized in $\text{TiO}_2\text{@SWCNT}^{53}$ and $\text{TiO}_2\text{@graphene}^{10}$. However, the contact area between TiO_2 and graphene or carbon nanotubes is limited. Zhang *et al.* reported an *in situ* polymer encapsulation-graphitization method to improve the contact between TiO_2 and carbon coverings, and further programmed the thickness of a carbon shell on a TiO_2 belt.¹⁷ However, the obtained $\text{TiO}_2\text{@C}$ showed an effect only under UV-light irradiation. Mesoporous PDA as a scaffold for TiO_2 enlarged the contact area between a graphene-like carbon layer and TiO_2 , and provided a codopant for TiO_2 . Fig. 11 illustrates the electron transition and visible-light photocatalytic mechanism of $\text{TiO}_2\text{@C-PDA}$. Moreover, theoretical calculations⁵⁴ and experimental studies⁵⁵ have indicated that codoping of non-metals and metals show synergistic effects on narrowing the band gap of TiO_2 . The catechol and amine groups on PDA nanotubes have strong interactions with various metal ions that can be transferred to multiple metal and non-metal dopants, thus showing the great versatility of PDA as a unique nitrogen-doped carbon scaffold for various catalysts. Visible light-induced photodecomposition is a green and renewable process for removing organic contaminants in water. Simple fabrication of $\text{TiO}_2\text{@C-PDA}$ photocatalysts endows it with great potential for industrial applications. It may have important and beneficial consequences on the environment and economy.

4. Conclusions

Herein, tubular and mesoporous TiO_2 were fabricated *via* sol-gel and calcination processes in the presence of PDA nanotubes. The obtained $\text{TiO}_2\text{@C-PDA}$ nanotubes were co-doped by carbon and nitrogen, which effectively narrowed the band gap of $\text{TiO}_2\text{@C-PDA}$. The crystal size of TiO_2 was confined through tightly covered PDA scaffolds, which efficiently suppressed crystal growth in calcination. Through optimized calcination, carbon cover on TiO_2 was thinned and graphitized. Such thinning is convenient for taking advantage of a conductive carbon layer while avoiding the blocking effect from a thick carbon layer and, in turn, results in efficient separation of the

photogenerated electron–holes of TiO_2 . The produced $\text{TiO}_2\text{@C-PDA}$ (400-2) with optimal structural and compositional advantages showed excellent photodegradation efficiency for MO under visible-light irradiation (>420 nm). PDA scaffolds can bring versatile benefits for TiO_2 photocatalysts.

Acknowledgements

The authors gratefully acknowledge the National Natural Science Foundation of China (Grant Numbers 21374132 and 21273281) for financial support.

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