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Facile synthesis of an urchin-like Sb₂S₃ nanostructure with high photocatalytic activity[†]

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Herein, an urchin-like Sb₂S₃ nanostructure has been synthesized without a surfactant *via* a wet chemical method. The crystal structure, morphology, composition and optical properties were characterized using XRD, TEM, SEM, EDS, Raman spectroscopy, and diffuse reflectance absorption spectroscopy. The factors, including the reaction time, temperature, and ratio of the raw materials, influencing the evolution of the urchin-like morphology have been discussed, and a plausible formation mechanism for the urchin-like Sb₂S₃ has been proposed. The urchin-like Sb₂S₃ micro/nanostructure exhibits high catalytic performance towards the degradation of MB under visible light irradiation. The photodegradation ratio of MB is up to 99.32% under visible light irradiation of 130 min. Our synthesis method will be extended to prepare other photocatalysts.

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In recent years, nanomaterials with controllable size and novel morphologies have been extensively studied due to their unique physical and chemical properties.¹⁻⁵ For instance, AgBr nanoplates,⁶ AgCl octahedrons⁷ and NaTi₂(PO₄)₃ nanocubes⁸ show unique photocatalytic activities and electrochemical performances. Hierarchical flower-like SnO₂ nanospheres⁹ or TiO₂ nanorod arrays^{10,43} exhibit highly efficient gas sensing and photoelectrochemical properties.

As an important V-VI compound, antimony sulfide (Sb₂S₃) is considered as a promising material for energy conversion due to its suitable band gap (1.5-2.2 eV), which covers the range of the solar spectrum.11-13 To the best of our knowledge, Sb₂S₃ nanomaterials with different morphologies have been mainly synthesized using various surfactants.¹⁴ Surfactants can regulate the morphology and structure of the nanoparticles effectively by parceling on the surface of the particles through the coordination or charge effect.¹⁵ By adding surfactants such as cetyltrimethylammonium bromide (CTAB),16 polyethylene glycol (PEG),17 dodecyltrimethylammonium bromide (DTAB)18 and polyvinylpyrrolidone (PVP),19-21 some nanostructures including straw-tied-like, nanorod^{22,23}, nanosheet,²⁴ nanotube, dandelion-like, double cauliflower-like,25 bar,26 dumbbell-like, and nanowire bundle structures have been successfully prepared, which show potential applications in photocatalysis and energy storage areas. However, how to effectively synthesize

 Sb_2S_3 crystals and control their morphology *via* a simple method is still challenging for materials scientists. Herein, we report a simple wet chemical method to synthesize an urchinlike Sb_2S_3 nanostructure without any surfactant. Moreover, we have examined the photocatalytic activity of Sb_2S_3 by the degradation of MB under simulated visible light irradiation. Our study presents an Sb_2S_3 nanostructure with good application in the visible-light photocatalytic field.

The morphology and size of the products were revealed by the SEM and TEM images. Fig. 1a shows the SEM image of the urchins-like Sb₂S₃ prepared under refluxing conditions at 160 °C for 5 h. The as-prepared Sb₂S₃ products mainly consist of the uniform urchin-like structure. The magnified SEM image shows that the urchin-like structure consists of nanorods stretching radially from the same central point (Fig. 1b). The TEM image further confirms that the urchin-like Sb₂S₃ is made up of many individual nanorods with lengths about 10 μ m and diameters of 100 nm (Fig. 1b and c). The measured spacing of the crystallographic planes is 0.309 nm, which is consistent with the (320) plane lattice distance of orthorhombic Sb₂S₃ (Fig. 1d).

The XRD pattern of the obtained sample is shown in Fig. 1e. All the diffraction peaks in the XRD pattern can be readily indexed to orthorhombic Sb₂S₃ (JCPDS no. 42-1393) with the calculated lattice parameters of a = 1.120 nm, b = 1.128 nm and c = 0.383 nm. No characteristic peaks for impurities are observed. The fact that the reflection peaks of Sb₂S₃ are strong and sharp indicates that Sb₂S₃ is highly crystalline (Fig. 1e). The EDS spectrum reveals that the as-prepared products consist of S and Sb elements, and the observed C and Cu peaks are due to the carbon-coated Cu grid²⁷ (Fig. 1f). Furthermore, the elemental distribution ratio of S and Sb in the compound was

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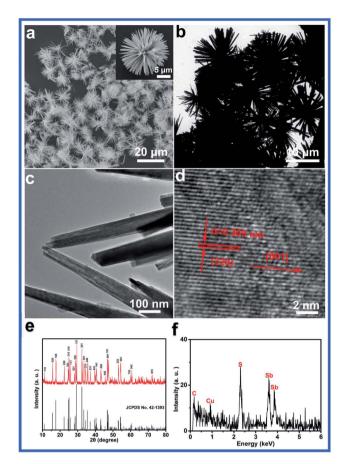


Fig. 1 (a) SEM, (b and c) TEM and (d) HRTEM images, (e) XRD pattern and (f) EDS spectrum of urchin-like ${\rm Sb}_2{\rm S}_3$.

found to be about 3 : 2 through the quantification calculation of the EDS peaks, which was consistent with the chemical formula of antimony sulfide.²⁸

The effects of the reaction time, reaction temperature and the ratio of the raw materials on the as-obtained products were carefully studied to investigate the formation of the urchin-like morphology. Fig. 2 reveals the SEM images of the as-obtained samples prepared at different reaction times. Initially, no product was formed. When the reaction time was prolonged to 2 h, amorphous antimony sulfide was formed (Fig. 2a). After 3 h, single units with slight splitting were observed (Fig. 2b). Upon increasing the reaction time, uniform urchin-like Sb₂S₃ was obtained. When the reaction time was further increased to 6 h, the urchin-like structure was transformed into a rod-flower structure in which the thorns of the urchin-like structure were thicker (Fig. 2c). As shown in Fig. 2d and e, after 9 h, the rodflower structure began to rupture and finally converted into irregular nanorods with lengths of about 15 µm, as shown in Fig. 2f.

The reaction temperature also has a significant effect on the morphologies of the final products. As shown in Fig. 3a, amorphous nanoparticles with an irregular shape were formed at 140 $^{\circ}$ C. When the temperature was increased to 150 $^{\circ}$ C, a simple splitting phenomenon appeared. Upon further

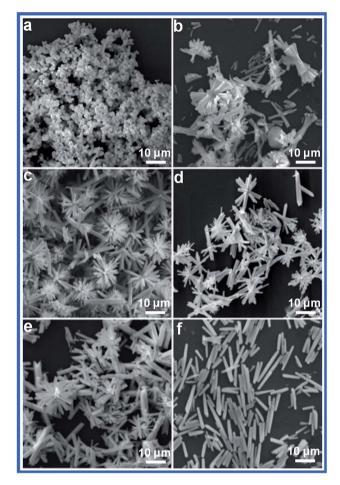


Fig. 2 SEM images of the Sb_2S_3 prepared at 160 °C for different times: (a) 2 h, (b) 3 h, (c) 6 h, (d) 9 h, (e) 12 h, and (f) 15 h.

increasing the temperature, the urchin-like micro/ nanostructure was formed. A further increase in the reaction temperature led to the formation of rod flowers. When the reaction temperature was 180 °C, some rod-like structures appeared.

We examined the effect of the ratio of the raw materials on the product. Only rod particles were observed at a ratio of S/Sb =1 : 1 (Fig. 4a). When the ratio was increased to 1 : 3, the urchin shapes were observed. A further increase in the concentration of the sulfur source led to the appearance of nest structures. Irregular spherical particles were finally generated when the ratio of the raw materials was 1 : 7.

Based on the experimental results, a plausible reaction for the synthesis of the Sb_2S_3 crystals can be interpreted as follows:

$$CH_3CSNH_2 + 2OH^- \rightarrow S^{2-} + CH_3COO^- + NH^{4+}$$
(1)

$$[Sb_2(C_4H_4O_6)_2]^{2-} \to 2Sb^{3+} + 2(C_4H_4O_6)^{4-}$$
(2)

$$2Sb^{3+} + 3S^{2-} \rightarrow Sb_2S_3\downarrow \tag{3}$$

According to the abovementioned experimental results, we speculated the formation mechanism of the urchin-like Sb_2S_3 structure. Sb_2S_3 is a highly anisotropic semiconducting material

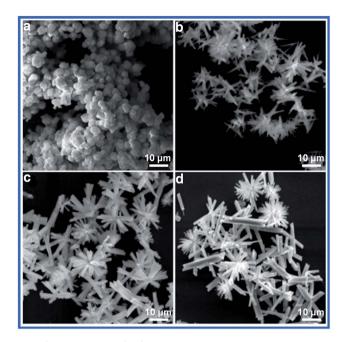
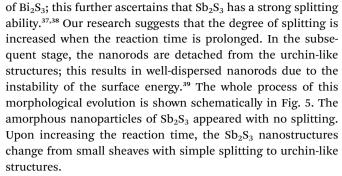


Fig. 3 SEM images of Sb_2S_3 prepared for 5 h at different reaction temperatures: (a) 140 °C, (b) 150 °C, (c) 170 °C and (d) 180 °C.

with infinite ribbon-like (Sb₄S₆) polymers and layers in its orthorhombic crystal structure.^{29,30} As already reported in the literature, Sb₂S₃ tends to easily form one-dimensional nanorods along the *c*-axis due to intermolecular attraction between the antimony and sulfur atoms.^{31–34} Because of the fast and anisotropic crystal growth, many newborn clusters agglomerate together to deposit on a particular particle face, causing crystal splitting when the solution is in the excess saturation state.^{35,36} Then, each of the nanorods starts to grow into urchin-like crystals. This is similar to the chain-like crystalline structure



The fact that the optical properties of the as-prepared Sb₂S₃ are determined using the absorption spectrum (Fig. 6a) provides a simple and effective method to estimate the band gap of the urchin-like Sb₂S₃.^{40,41} The band gap value of 1.64 eV determined by other researchers is quite comparable to the values reported in the bibliographical information.42 Furthermore, it can be seen that optical absorption occurs in nearly all the visible light region; this suggests that the as-synthesized Sb₂S₃ can be stimulated by visible light. Raman spectrum of the urchin-like Sb₂S₃ is shown in Fig. 6b. The appearance of sharp peaks at 147, 198, 257, 305, and 452 cm^{-1} suggests the formation of a highly crystalline product, which is consistent with the XRD results.43 Due to suitable band gap of urchin-like Sb₂S₃, the photocatalytic activity of urchin-like Sb₂S₃ was assessed by depositing an organic dye (MB) in an aqueous solution under visible light irradiation. The concentration of MB dye was monitored at 665 nm using an UV/vis spectrophotometer. From the blank experiment (Fig. S2[†]), we can see a slight drop in the curve due to self-degradation of MB dye under visible light irradiation. Therefore, we can conclude that the dye is almost photo-stable under these conditions. It can be clearly observed from Fig. 6c that the organic dye is slightly degraded by Sb₂S₃ in the dark. According to Fig. 6d, the organic dye could be degraded by only 23.1% using H₂O₂, whereas the degradation efficiency for MB dye was 99.34% in the presence of Sb₂S₃ and H₂O₂ after 130 min; this indicated that the degradation of the organic dye was mainly attributed to the excellent

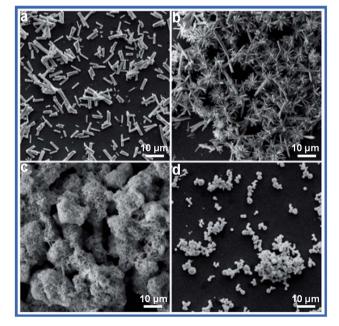


Fig. 4 SEM images of Sb_2S_3 prepared with different ratios of S and Sb: (a) S/Sb = 1 : 1, (b) S/Sb = 1 : 3, (c) S/Sb = 1 : 5, and (d) S/Sb = 1 : 7.

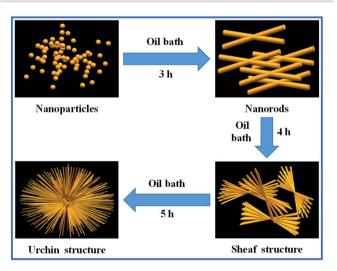


Fig. 5 A schematic of the formation of the urchin-like Sb₂S₃ structure.

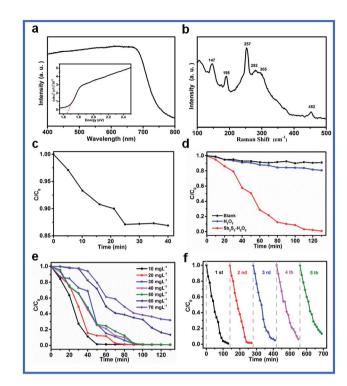


Fig. 6 (a) The diffuse reflectance absorption spectrum of the asprepared Sb₂S₃ sample (the inset presents the corresponding plots of $(\alpha hv)^2$ versus photon energy). (b) The room-temperature Raman spectrum. (c) The degradation curve for MB in the dark. (d) The degradation curves for MB using the different catalyst systems (e) the degradation curves at different MB concentrations. (f) The cycling runs for the degradation of MB.

photocatalytic activity of Sb₂S₃. The photoluminescence spectra of the as-prepared urchin-like Sb₂S₃ shows a low signal, suggesting the good charge separation properties of our samples (Fig. S1a^{\dagger}). Fig. S1b^{\dagger} shows the *I*-*T* curves of the FTO/Sb₂S₃ photoelectrode, of which the photocurrent intensity is 36.057 μ A cm⁻² at a potential of 0 V versus Ag/AgCl, and this fast photocurrent response is consistent with the PL results. Fig. 6e shows the photocatalytic degradation of an aqueous solution of MB at different concentrations by urchin-like Sb₂S₃. However, with an increase in the concentration of MB, the degradation efficiency is reduced. Furthermore, the cycling performance of urchin-like Sb₂S₃ was measured using the same photodegradation process. The photocatalyst was obtained by centrifugation and washed with deionized water after each cycle. The degradation ratio decreased slightly after reuse for 5 times; the slight decay in the photodegradation activity was partially ascribed to the inevitable loss of the photocatalyst during the washing and centrifugation process; this suggested that Sb₂S₃ exhibited prominent photocatalytic stability (Fig. 6f).

The promising photocatalytic activity of Sb_2S_3 can be mainly attributed to its relatively narrow band gap (1.64 eV), making it a perfect photocatalyst to efficiently absorb and utilize visiblelight energy. As is known, the urchin-like nanostructure has been extensively investigated due to its potential applications in many areas such as in varistors, electronic devices, UVabsorbers, and catalysis.^{47,48} This kind of nanostructure has

a high specific surface area, which endows it with a large contact area with the dye and superior light harvesting efficiency in the field of photocatalysis.49 Moreover, the single crystalline nanorod-assembled urchin-like nanostructure is favorable for fast electron transfer and thus facilitates electron and hole separation.^{46,50,51} In particular, when urchin-like Sb₂S₃ is irradiated by visible light of energy greater than its band gap, electron-hole pairs will be generated and partially separated. The electrons and holes react with the adsorbed surface substances, such as O₂ and OH⁻, to form the reactive species $\cdot O^{2-}$, $\cdot OH$, which are the major oxidative species in the decomposition of organic pollutants. Then, the oxidative species degrade the organic pollutant into small molecules such as CO2 and H2O.44,45 p-Benzoquinone (BQ) and isopropyl alcohol (IPA) were used as additive agents to trap the active species $\cdot O_2^{-1}$ and ·OH, respectively. Fig. S2[†] shows the degradation curves for MB after the addition of the different capture agents. It can be seen that IPA has a significant inhibitory effect on the degradation of MB, whereas the effect of BQ is less; this indicates that \cdot OH plays a major role in the photocatalytic process.

The degradation of the organic dye was accomplished *via* a series of parallel and consecutive redox reactions as show in eqn (2.1)-(2.5).

$$Sb_2S_3 + h\nu \to Sb_2S_3 (h_{VB}^+ + e_{CB}^-)$$
 (2.1)

$$O_2 + e_{CB}^- \rightarrow \cdot O_2^- \tag{2.2}$$

$$H_2O \rightarrow H^+ + OH^- \tag{2.3}$$

$$OH^- + h_{VB}^+ \rightarrow OH$$
 (2.4)

$$MB + \cdot OH + \cdot O^{2-} \rightarrow CO_2 + H_2O + \dots \qquad (2.5)$$

In summary, an urchin-like Sb₂S₃ nanostructure was successfully prepared without surfactants using a wet chemical method. The corresponding crystal splitting formation mechanism of urchin-like Sb₂S₃ was tentatively suggested. The narrow band gap of the urchin-like Sb₂S₃ was evaluated to be 1.64 eV, which was close to the best photoelectric conversion value reported to date. In addition, urchin-like Sb₂S₃ exhibited excellent photocatalytic performance; the degradation efficiency for the organic dye was 99.34% after exposure to visible-light irradiation for 130 min. The high photocatalytic activity of Sb₂S₃ was mostly due to its narrow band gap and wide absorption range of visible light. Our investigation demonstrates that Sb₂S₃ with urchin-like nanostructures has great potential to be applied in the degradation of organic contaminants.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 N. E. Kelly, S. O. Lee and K. D. Harris, *J. Am. Chem. Soc.*, 2001, **123**, 12682.
- 2 A. M. Morales and C. M. Lieber, Science, 1998, 279, 208.
- 3 X. Peng, L. Manna and W. Yang, Nature, 2000, 404, 59.
- 4 A. P. Alivisatos, Science, 1996, 271, 933.
- 5 H. Dai, E. W. Wong and Y. Z. Lu, Nature, 1995, 375, 769.
- 6 H. Wang, J. Gao, T. Q. Guo, R. M. Wang and L. Guo, *Chem. Commun.*, 2012, **48**, 275.
- 7 H. Wang, X. F. Lang, R. Hao, L. Guo, J. H. Li, L. H. Wang and X. D. Han, *Nano Energy*, 2016, **19**, 8.
- 8 J. Yang, H. Wang, P. F. Hu, J. J. Qi, L. Guo and L. H. Wang, *Small*, 2015, **31**, 3744.
- 9 H. Wang, Q. Q. Liang, W. J. Wang, Y. R. An, J. H. Li and L. Guo, *Cryst. Growth Des.*, 2011, **11**, 2942.
- 10 X. T. Wang, C. H. Liow, D. P. Qi, B. W. Zhu, W. R. Leow, H. Wang, X. D. Chen and S. Z. Li, *Adv. Mater.*, 2014, **21**, 3506.
- 11 L. Amy, G. Q. Lu, T. John and J. Yates, *Chem. Rev.*, 1995, **95**, 735.
- 12 B. Thompson and J. Fréchet, *Angew. Chem., Int. Ed.*, 2008, 47, 58.
- 13 W. U. Huynh, J. J. Dittmer and A. P. Alivisatos, *Science*, 2002, 295, 2425.
- 14 Z. Qiao, Z. L. Li and J. Y. Li, J. Cryst. Growth, 2009, 311, 3651.
- 15 Q. Zhang, Q. A. Zhu and X. F. Sun, *Chin. J. Inorg. Chem.*, 2008, 24, 547.
- 16 Y. Y. Zhu, P. Nie, L. F. Shen, S. Y. Dong, Q. Sheng, H. S. Li, H. F. Luo and X. G. Zhang, *Nanoscale*, 2015, 7, 3309.
- 17 H. S. Hou, M. J. Jing, Z. D. Huang, Y. C. Yang, Y. Zhang, J. Chen and X. B. Ji, ACS Appl. Mater. Interfaces, 2015, 7, 19362.
- 18 Y. B. Zhao and A. Manthiram, Chem. Mater., 2015, 27, 6139.
- 19 B. Li, H. Wang, B. W. Zhang, P. F. Hu, C. Chen and L. Guo, ACS Appl. Mater. Interfaces, 2013, 5, 12283.
- 20 H. J. Zhang, M. Ge, L. T. Yang, Z. Zhou and W. Chen, *J. Phys. Chem. C*, 2013, **117**, 10285.
- 21 C. S. Yan, G. Chen, D. H. Chen and J. Pei, *CrystEngComm*, 2014, **16**, 3965.
- 22 Y. Li, X. Wei, B. Zhu, H. Wang, Y. Tang, T. C. Sum and X. Chen, *Nanoscale*, 2016, **8**, 11284.
- 23 Y. Li, J. Feng, H. Li, X. Wei, R. Wang and A. Zhou, *Int. J. Hydrogen Energy*, 2016, **41**, 4096.
- 24 Y. Li, X. Wei, X. Yan, J. Cai, A. Zhou, M. Yang and K. Liu, *Phys. Chem. Chem. Phys.*, 2016, **18**, 10255.
- 25 N. Maiti, S. H. Im, Y. H. Lee and S. I. Seok, *ACS Appl. Mater. Interfaces*, 2012, **4**, 4787.
- 26 S. S. Yao, J. Cui, Z. H. Lu, Z. L. Xu, L. Qin, J. Q. Huang, Z. Sadighi, F. Ciucci and J. K. Kim, *Adv. Energy Mater.*, 2017, 7, 241.

- 27 G. Y. Chen, B. Deng and G. B. Cai, *J. Phys. Chem. C*, 2007, **112**, 672.
- 28 X. H. Xiong, G. H. Wang, Y. W. Lin, Y. Wang, X. Ou, F. H. Zheng, C. H. Yang, J. H. Wang and M. L. Liu, ACS Nano, 2016, 10, 10953.
- 29 I. Moraseró and J. Bisquert, *J. Phys. Chem. Lett.*, 2010, **116**, 1579.
- 30 G. Y. Chen, B. Deng and G. B. Cai, *J. Phys. Chem. C*, 2008, **112**, 672.
- 31 S. S. Yao, J. Cui, Z. H. Lu, Z. L. Xu, L. Qin, J. Q. Huang, Z. Y. Sa, F. Ciucci and J. K. Kim, *Adv. Energy Mater.*, 2017, 7, 1602149.
- 32 J. Yang, J. H. Zeng, S. H. Yu, L. Yang, Y. H. Zhang and Y. T. Qian, *Chem. Mater.*, 2000, **12**, 2924.
- 33 X. F. Duan, Y. Huang, Y. Cui, J. F. Wang and C. M. Liber, *Nature*, 2001, **409**, 66.
- 34 H. M. Huang, S. Mao, H. Feick, H. Yan, Y. Wu, H. Kind, E. Weber, R. Russo and P. D. Yang, *Science*, 2001, 292, 1897.
- 35 C. An, K. Tang and Q. Yang, Inorg. Chem., 2003, 42, 8081.
- 36 B. Gates, B. Mayers, Y. Wu, Y. Sun and B. Cattle, *Adv. Funct. Mater.*, 2010, **10**, 679.
- 37 J. Tang and A. P. Alivisatos, Nano Lett., 2006, 6, 2701.
- 38 L. Tian, H. Y. Tan and J. J. Vittal, *Cryst. Growth Des.*, 2008, 8, 734.
- 39 J. Ota, P. Roy, S. K. Srivastava, B. B. Nayak and A. K. Saxena, *Cryst. Growth Des.*, 2008, **8**, 6.
- 40 X. W. Zheng, Y. Xie, L. Y. Zhu, X. C. Jiang, Y. B. Jia, W. H. Song and Y. P. Sun, *Inorg. Chem.*, 2002, 41, 455.
- 41 P. H. Zhang, C. X. Liu, X. L. Qi and X. Dai, *Nat. Phys.*, 2009, 5, 438.
- 42 H. F. Dong, J. P. Lei, H. X. Ju, F. Zhi, H. Wang, W. Guo, Z. Zhu and F. Yan, *Angew. Chem., Int. Ed.*, 2012, **124**, 4685.
- 43 C. An, K. Tang and Q. Yang, Inorg. Chem., 2003, 42, 8081.
- 44 J. Kimling, M. Maier, B. Okenve, V. Kotaidis and H. Ballot, *Environ. Sci. Technol.*, 2006, **32**, 15700.
- 45 M. Zhou, L. Lei and Q. Dai, Chem. Commun., 2007, 2645.
- 46 X. Tang, J. Qian, Z. Wang, H. Wang, Q. Feng and G. Liu, J. Colloid Interface Sci., 2008, 330, 386.
- 47 X. Yu, Z. Zhao, J. Zhang, W. Guo, L. Li, H. Liu and Z. L. Wang, *CrystEngComm*, 2017, **19**, 129.
- 48 H. Wang, Y. Bai, Q. Wu, W. Zhou, H. Zhang, J. Li and L. Guo, *Phys. Chem. Chem. Phys.*, 2011, **13**, 7008.
- 49 J. H. Pan, X. Z. Wang, Q. Huang, C. Shen, Z. Y. Koh, Q. Wang, A. Engel and D. W. Bahnemann, *Adv. Funct. Mater.*, 2014, 24, 95.
- 50 J. Liu, Z. Guo, W. Wang, Q. Huang, K. Zhu and X. Chen, *Nanoscale*, 2011, 3, 1470.
- 51 A. Hui, J. Ma, J. Liu, Y. Bao and J. Zhang, *J. Alloys Compd.*, 2017, **696**, 639.