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## Anion vacancy correlated photocatalytic CO<sub>2</sub> to CO conversion over quantum-confined CdS nanorods under visible light†

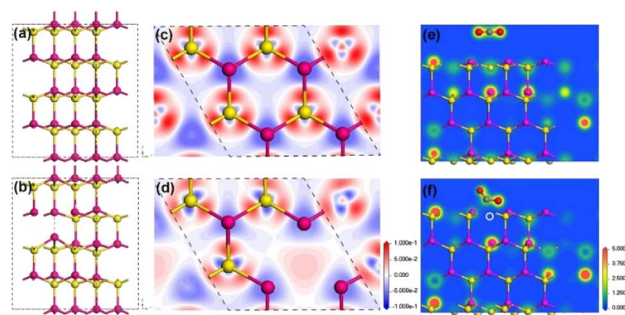
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Surface vacancies in II–VI semiconductors have been confirmed as active sites for photocatalytic CO<sub>2</sub> reduction. Here we take quantum-confined CdS nanorod as a model photocatalyst to correlate anion vacancies with photocatalytic CO<sub>2</sub> reduction performance. In terms of electronic structure change, CdS nanorods with more surface S vacancies give enhanced CO generation rates, which were confirmed by deliberately introducing S vacancies *via* a post-treatment.

Production of solar fuels or valuable chemicals from CO<sub>2</sub> reduction driven by solar energy remains one of the most promising pathways to relieve the energy crisis and environmental disruption.<sup>1–9</sup> Semiconductor nanocrystals containing II–VI elements have emerged as ideal candidates in the field of CO<sub>2</sub> photoreduction,<sup>10–14</sup> because of their large extinction coefficient, high probability of exciton generation and easy modification of surface properties.<sup>15,16</sup> Although most of the reported systems involve external active sites,<sup>17–19</sup> such as metal complexes<sup>20,21</sup> and surface ligands,<sup>22</sup> surface vacancies of II–VI nanocrystals were also found to work as active sites.<sup>23–25</sup> For example, zinc vacancies in ZnS QDs were effective binding sites for CO<sub>2</sub> photoreduction.<sup>26</sup> Anion vacancies on CdSe/CdS QDs could achieve highly efficient and selective CO evolution *via* photocatalytic CO<sub>2</sub> reduction.<sup>27</sup> Moreover, S vacancies formed *in situ* on CdS nanoparticles contributed to electrocatalytic CO<sub>2</sub> reduction.<sup>28</sup> These results prompted us to understand the inherent correlation of surface anion vacancies and catalytic

performance of CO<sub>2</sub> photoreduction by semiconductors nanocrystals.

In this contribution, we chose quantum-confined CdS nanorod as a model photocatalyst to explore the anion vacancy–activity relationship toward photocatalytic CO<sub>2</sub> reduction. First, a structural model of hexagonal CdS without or with surface S vacancies was built for density functional theory (DFT) calculations,<sup>27</sup> see details in the ESI.† Owing to the deficiency of S atoms, the four adjacent Cd atoms deviated from their original positions and demonstrated a small shift toward S vacancies with distinct lattice constriction as compared with pristine CdS (Fig. 1a and b). Electron density calculations (Fig. 1c and d) revealed that the electron redistribution around S vacancies occurred by removing a neutral S atom from the CdS supercell. The two unbound electrons became active and produced a localized electron trapping center because of positively charged S vacancies. As shown in Fig. 1e and f, a linear CO<sub>2</sub> molecule bent on a S vacancy site after geometric optimization



**Fig. 1** Optimized bulk lattice structure of 2 × 2 × 2 supercell hexagonal CdS without (a) and with S vacancies (b). Charge-density difference maps around the S site at the (001) cross section without (c) and with S vacancies (d). Electron density distribution around the adsorbed CO<sub>2</sub> site at the hexagonal CdS(001) facet (e) without and (f) with S vacancies. Cd, S, C and O atoms are represented by pink, yellow, gray and red balls, respectively. The S vacancy is indicated by a white dashed circle. The color level in (c) and (d) goes from a reduction (blue) to a gain in density (red) and the units are e<sup>−</sup> Å<sup>−2</sup>.

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while remaining in the linear configuration on the perfect CdS(001) facet. The adsorption energy changed from  $-0.078$  eV on the perfect facet to  $-0.815$  eV on the S-vacancy facet. Meanwhile, the C=O bond was elongated from  $1.17$  Å to  $1.268$  Å and the O–C–O angle reduced from  $180^\circ$  to  $122.37^\circ$ , indicating that the CO<sub>2</sub> molecule was highly activated on S vacancies.<sup>6,29</sup> In contrast, the geometric structure of the CO<sub>2</sub> molecule almost remained unchanged on the perfect CdS surface. The S vacancies changed the local electronic structure and promoted CO<sub>2</sub> adsorption and activation.

Under the guidance of theoretical studies, we prepared quantum-confined CdS nanorods with different numbers of S-vacancies to work as photocatalysts towards CO<sub>2</sub> reduction under visible light. To our delight, with more surface S vacancies, the rate of CO generation was significantly enhanced, indicating the positive correlation between CO evolution and the amount of surface S vacancies. Furthermore, the deliberate introduction of S vacancies on CdS nanorods *via* a post-treatment procedure of exchanging Cd<sup>2+</sup> with Ag<sup>+</sup> ions could also improve the rate of CO evolution by about  $\sim 3$ -fold, further suggesting that the more surface S vacancies on CdS the higher the activity for CO<sub>2</sub> photoreduction.

Quantum-confined CdS nanorods with different amounts of S vacancies were synthesized *via* a modified hot injection method at high temperature,<sup>30,31</sup> see synthetic details in the ESI†. The obtained CdS nanorods with varied amounts of S vacancies were denoted as CS-1, CS-2, CS-3 and CS-4, respectively. Transmission electron microscopy (TEM) confirmed that the obtained CdS samples were in the shape of nanorods (Fig. 2a–d), in which the four CdS samples exhibited a similar size distribution with a length of  $\sim 20$  nm and a width of  $\sim 4$  nm. An enlarged aberration-corrected scanning transmission electron microscopy (STEM) image showed a lattice fringe of  $3.29$  Å, which was ascribed to the (002) facet of hexagonal CdS (Fig. S1, ESI†). The corresponding elemental mapping of a single CdS nanorod showed the coexistence of Cd and S elements (Fig. 2e), which was



Fig. 2 TEM images of (a) CS-1, (b) CS-2, (c) CS-3 and (d) CS-4 CdS samples. (e) STEM image of CdS nanorods and the corresponding elemental mapping, respectively.

further verified by full X-ray photoelectron spectroscopy (XPS) analysis (Fig. S2, ESI†). Moreover, the narrow and sharp peaks of the prepared samples in X-ray diffraction (XRD) patterns perfectly matched the standard pattern of hexagonal CdS (PDF No. 41-1049),<sup>31</sup> indicating the successful formation of CdS nanorods with identical shapes and crystalline structures (Fig. 3a).

The four CdS samples were then characterized in solution. As shown in Fig. 3b, the first absorption peak was centered at  $\sim 440$  nm in ultraviolet-visible (UV-vis) absorption spectra, because of the quantum confinement effects of CdS nanorods in the radial direction. A similar first absorption peak location suggested that these samples of the four CdS nanorods, either in width or length, were well maintained. Subsequently, electron paramagnetic resonance (EPR) was carried out. The observed signal centred at  $g = 2.0016$  (Fig. S3, ESI†) resulted from anion S vacancies, in line with the fact that anion vacancies in II–VI semiconductors gave rise to EPR signals near  $g = 2$ .<sup>32–35</sup> In addition, the line at  $g = 2.0016$  with slight asymmetry was attributed to S vacancies located at the surface of nanorods because surface defects with an anisotropic Zeeman interaction were actually anisotropic.<sup>35</sup> In CdS nanorods, the formation of S vacancies on the surface resulted in excess Cd sites associated with electron traps,<sup>36–38</sup> thus leading to the formation of positively charged trap states with paramagnetism.

Given the distinct electron binding energies of inner and surface atoms, XPS analysis, a powerful technique to provide the composition information of nanomaterials, was utilized to reveal the surface stoichiometry of CdS nanorods. Taking the ratio of their respective integrated XPS signals,<sup>39,40</sup> we calculated the percentages of surface Cd and S atoms on CdS nanorods, see details in the ESI (Fig. S4 and Table S1, ESI†). The ratios of surface atoms to whole atoms for Cd and S in CdS nanorods were determined from integrated XPS data. As summarized in



Fig. 3 (a) XRD patterns of CS-1, CS-2, CS-3 and CS-4 CdS nanorods. (b) UV-vis absorption spectra and (c) steady-state PL spectra (excitation: 400 nm) of CdS nanorods for CS-1, CS-2, CS-3 and CS-4 in *n*-hexane at room temperature.

Table 1, the ratios of surface Cd to S atoms for various CdS samples were determined to be 3.24, 3.40, 3.70 and 3.88 from CS-1 to CS-4. Obviously, the surface Cd/S ratios of all samples were greater than 1, indicating surface Cd-rich properties.<sup>27</sup> Thus, the existing defects in CdS were electron traps-anion S vacancies,<sup>41</sup> and the increased Cd/S ratio from CS-1 to CS-4 suggested the enhanced amounts of S vacancies.

The steady-state photoluminescence (PL) spectroscopic analysis (Fig. 3c) displayed two emissions including a narrow band at  $\sim 457$  nm and a broad band at  $\sim 630$  nm, which were radiative recombinations of photogenerated electrons and holes at the band edge and the trap sites, respectively.<sup>36,42</sup> To scrutinize the broad emission, the intensity of the narrow bands in the spectra was normalized. The significant enhancement of the trap-state emission at  $\sim 630$  nm from CS-1 to CS-4 indicated the increased amounts of surface trap states in these samples.<sup>43</sup> Apparently, this trend was well consistent with the surface Cd/S ratio variation from CS-1 to CS-4, implying that the emission peak at  $\sim 630$  nm was radiative recombination of electron-hole at S vacancies. In addition, the corresponding PL lifetimes of the four samples decreased from 4.9 ns to 2.6 ns (Fig. S5 and Table S2, ESI†), suggesting the promoted electron-hole recombination at surface S vacancies.

Then, photocatalytic CO<sub>2</sub> reduction experiments were carried out under ambient conditions, see details in the ESI.† The initial photocatalytic experiments were performed with triethylamine (TEA) as a sacrificial reagent under visible light ( $\lambda = 450$  nm). Excitingly, CdS nanorods could produce CO gas as the sole carbon-containing product under visible light as no other reduction products were observed. Moreover, the CO generation rate was gradually improved from  $\sim 295$  mmol g<sup>-1</sup> to  $\sim 513$  mmol g<sup>-1</sup> with increased S vacancies from CS-1 to CS-4 under long-time irradiation (Fig. 4a). Meanwhile, negligible change was observed in the selectivity of CO, calculated according to the equation  $\eta = [n_{\text{CO}}/(n_{\text{CO}} + n_{\text{H}_2})] \times 100\%$ , with variation in S vacancy amounts (Fig. 4b). Obviously, rich surface S vacancies showed vital influence on CO<sub>2</sub> reduction, which coincided with our DFT prediction. In order to further validate the origin of generated CO, an isotopic experiment using <sup>13</sup>CO<sub>2</sub> as the reactant was performed, and the obtained gaseous products were analyzed by gas chromatography-mass spectrometry (GC-MS). A signal at  $m/z = 29$  was observed (Fig. S6, ESI†), which could be assigned to the photogenerated <sup>13</sup>CO. This result directly confirmed the photoreduction of CO<sub>2</sub> to CO.

The valence-band (VB) XPS analysis (Fig. S7, ESI†) was utilized to determine the VB potentials of CdS, which were



Fig. 4 (a) Profile of photocatalytic CO<sub>2</sub> reduction activity and (b) the corresponding CO selectivities for CS-1, CS-2, CS-3 and CS-4. (c) The rate of photocatalytic CO<sub>2</sub> reduction, and (d) the corresponding selectivity of CO using original CdS (denoted as pristine), a physical mixture of CdS and AgNO<sub>3</sub> (denoted as pristine + AgNO<sub>3</sub>), and CdS after Ag substitution (denoted as Cd<sub>0.95</sub>Ag<sub>0.1</sub>S) as the photocatalysts. Error bars represent mean  $\pm$  s.d. of at least three independent experiments.

+1.49, +1.53, +1.61 and +1.61 V *versus* the normal hydrogen electrode (NHE) for CS-1, CS-2, CS-3 and CS-4 (Table 1), respectively. The band gaps of the four samples were calculated in the range of  $\sim 2.70$  eV to  $\sim 2.74$  eV by using the Tauc plot (Fig. S8, ESI†). Thus, the conduction band (CB) positions were calculated to be  $-1.21$  V,  $-1.21$  V,  $-1.13$  V and  $-1.13$  V *vs.* NHE for CS-1, CS-2, CS-3 and CS-4, respectively. Accordingly, the driving force for CO<sub>2</sub> reduction was almost identical with the varied amounts of S vacancies, thus excluding the influence of band-position change on CO<sub>2</sub> reduction.

In order to further confirm the above results, anion S vacancies were deliberately introduced on CdS nanorods *via* a post-treatment method.<sup>35,44,45</sup> AgNO<sub>3</sub> was here chosen as a Ag<sup>+</sup> precursor to realize the cation-exchange reaction under ambient conditions,<sup>39</sup> see details in the ESI.† TEM images indicated the unchanged morphology of CdS nanorods after exchanging 0.05% Cd<sup>2+</sup> with Ag<sup>+</sup> ions, denoted as Cd<sub>0.95</sub>Ag<sub>0.1</sub>S (Fig. S9, ESI†). The first absorption peak indicated similar band gaps (Fig. S10, ESI†). However, the emission intensity at  $\sim 630$  nm dramatically enhanced (Fig. S11, ESI†), suggesting the increase of trap states in the form of anion S vacancies. These results matched well with those of a previous report,<sup>45</sup> in which chemically induced Ag<sup>+</sup> ions could achieve electronic tuning of CdS nanorods. Subsequently, photocatalytic CO<sub>2</sub> reduction experiments were carried out with pristine CdS and Cd<sub>0.95</sub>Ag<sub>0.1</sub>S nanorods under identical conditions. As shown in Fig. 4c, the CO generation rate was improved from  $\sim 7.9$  mmol g<sup>-1</sup> h<sup>-1</sup> to  $\sim 22.5$  mmol g<sup>-1</sup> h<sup>-1</sup> after Ag<sup>+</sup> ion exchange, almost a 3-fold enhancement. Meanwhile, the selectivity of CO increased from  $\sim 75\%$  to  $\sim 84\%$  (Fig. 4d), indicating an effective pathway to

Table 1 Element ratios of Cd and S and band-energy levels in CdS samples

Samples	Surface Cd (%)	Surface S (%)	Surface Cd/S ratio	VB <i>vs.</i> NHE (V)	CB <i>vs.</i> NHE (V)
CS-1	41.24	37.94	3.24	1.49	-1.21
CS-2	47.59	26.84	3.40	1.53	-1.21
CS-3	47.36	39.07	3.70	1.61	-1.13
CS-4	43.14	34.61	3.88	1.61	-1.13



control CO<sub>2</sub> reduction activity and selectivity. DFT simulations also showed that CO<sub>2</sub> adsorption occurred at the trap states but not on Ag or Cd sites on the surface, indicating that CO<sub>2</sub> activation and reduction still proceeded at the anion vacancies in Cd<sub>0.95</sub>Ag<sub>0.1</sub>S nanorods (Table S3 and Fig. S12†). In contrast, a physical mixture of CdS nanorods and AgNO<sub>3</sub> (denoted as pristine + AgNO<sub>3</sub>) exhibited a similar CO generation rate to pristine CdS, further confirming that the improved CO generation was a result of increased anion traps, but not Ag species.

Because of the sluggish thermodynamic and kinetic nature of the linear CO<sub>2</sub> molecule, its activation is considered as a key step for further conversion. The anion S vacancies play two crucial roles in this process: one is as active sites for CO<sub>2</sub> adsorption and activation (*vide ante* DFT calculation results). The other is as effective electron trap states, which give rise to energy states energetically localized within the nanorods' energy gap and capture charge carriers from delocalized electronic states. The trap states reduce the mobility and the diffusion length of charge carriers, as illustrated in Fig. 5. In addition, the presence of trap states decreases interfacial charge transfer resistance, which is demonstrated by the smaller arc radius of the electrochemical impedance spectroscopy (EIS) Nyquist plot of vacancy-rich CdS nanorods (Fig. S13, ESI†).<sup>38</sup> With visible light irradiation, photogenerated electrons at the CB of CdS are captured by trap states, and reduce adsorbed CO<sub>2</sub> molecules at S vacancies on the CdS surface to generate COOH\* intermediates through a proton-coupled electron transfer process. Subsequently, the COOH\* intermediate is reduced by photoelectrons to generate OH\* and CO\* intermediates. Finally, OH\* combines with a proton to produce H<sub>2</sub>O and CO\* detaches from the surface to evolve the product of CO.<sup>25,46</sup>

In summary, we have demonstrated that rich anion S vacancies on quantum-confined CdS nanorods can effectively enhance photocatalytic CO<sub>2</sub> reduction activity. CdS nanorods with more S vacancies show enhanced CO<sub>2</sub> photoreduction activity. Combined with electronic tuning *via* the cation-exchange reaction of Ag<sup>+</sup> substituting Cd<sup>2+</sup> ions, the improved activity is a result of the promoted CO<sub>2</sub> adsorption and activation, the facilitated interfacial charge transfer, and reduced mobility and diffusion length of charge carriers in the presence of high density of surface S vacancies. Understanding the influence of surface S vacancies on CdS photocatalytic CO<sub>2</sub>

reduction activity would provide new insights for tuning CO<sub>2</sub> photoreduction performance under mild conditions.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 X. Li, J. Yu, M. Jaroniec and X. Chen, *Chem. Rev.*, 2019, **119**, 3962–4179.
- 2 D. Devasia, A. J. Wilson, J. Heo, V. Mohan and P. K. Jain, *Nat. Commun.*, 2021, **12**, 2612.
- 3 Y. Wang, X. Shang, J. Shen, Z. Zhang, D. Wang, J. Lin, J. C. S. Wu, X. Fu, X. Wang and C. Li, *Nat. Commun.*, 2020, **11**, 3043.
- 4 Y. A. Wu, I. McNulty, C. Liu, K. C. Lau, Q. Liu, A. P. Paulikas, C.-J. Sun, Z. Cai, J. R. Guest, Y. Ren, V. Stamenkovic, L. A. Curtiss, Y. Liu and T. Rajh, *Nat. Energy*, 2019, **4**, 957–968.
- 5 Y.-J. Xu, *Angew. Chem., Int. Ed.*, 2021, **60**, 2–25.
- 6 H. Li, J. Zhao, L. Luo, J. Du and J. Zeng, *Acc. Chem. Res.*, 2021, **54**, 1454–1464.
- 7 G. Wang, C. T. He, R. Huang, J. Mao, D. Wang and Y. Li, *J. Am. Chem. Soc.*, 2020, **142**, 19339–19345.
- 8 H. L. Wu, X. B. Li, C. H. Tung and L. Z. Wu, *Adv. Mater.*, 2019, e1900709.
- 9 J. Albero, Y. Peng and H. García, *ACS Catal.*, 2020, **10**, 5734–5749.
- 10 P. Zhang, S. Wang, B. Guan and X. W. D. Lou, *Energy Environ. Sci.*, 2019, **12**, 164–168.
- 11 K. K. Sakimoto, A. B. Wong and P. Yang, *Science*, 2016, **351**, 74–77.
- 12 X.-B. Li, Z.-K. Xin, S.-G. Xia, X.-Y. Gao, C.-H. Tung and L.-Z. Wu, *Chem. Soc. Rev.*, 2020, **49**, 9028–9056.
- 13 B. Su, L. Huang, Z. Xiong, Y. Yang, Y. Hou, Z. Ding and S. Wang, *J. Mater. Chem. A*, 2019, **7**, 26877–26883.
- 14 G. Chen, F. Wei, Z. Zhou, B. Su, C. Yang, X. F. Lu, S. Wang and X. Wang, *Sustainable Energy Fuels*, 2023, **7**, 381–388.
- 15 X.-B. Li, C.-H. Tung and L.-Z. Wu, *Angew. Chem., Int. Ed.*, 2019, **58**, 10804–10811.
- 16 J. Wang, T. Xia, L. Wang, X. Zheng, Z. Qi, C. Gao, J. Zhu, Z. Li, H. Xu and Y. Xiong, *Angew. Chem., Int. Ed.*, 2018, **57**, 16447–16451.
- 17 F. Xu, K. Meng, B. Zhu, H. Liu, J. Xu and J. Yu, *Adv. Funct. Mater.*, 2019, **29**, 1904256.



Fig. 5 Schematic illustration of CdS nanorods with S vacancies for photocatalytic CO<sub>2</sub> to CO conversion.

- 18 J. Di, C. Chen, C. Zhu, P. Song, M. Duan, J. Xiong, R. Long, M. Xu, L. Kang, S. Guo, S. Chen, H. Chen, Z. Chi, Y.-X. Weng, H. Li, L. Song, M. Wu, Q. Yan, S. Li and Z. Liu, *Nano Energy*, 2021, **79**, 105429.
- 19 Y. L. Men, Y. You, Y. X. Pan, H. Gao, Y. Xia, D. G. Cheng, J. Song, D. X. Cui, N. Wu, Y. Li and S. Xin, *J. Am. Chem. Soc.*, 2018, **140**, 13071–13077.
- 20 M. F. Kuehnell, C. D. Sahm, G. Neri, J. R. Lee, K. L. Orchard, A. J. Cowan and E. Reisner, *Chem. Sci.*, 2018, **9**, 2501–2509.
- 21 M. F. Kuehnell, K. L. Orchard, K. E. Dalle and E. Reisner, *J. Am. Chem. Soc.*, 2017, **139**, 7217–7223.
- 22 Y.-X. Feng, H.-J. Wang, J.-W. Wang, W. Zhang, M. Zhang and T.-B. Lu, *ACS Appl. Mater. Interfaces*, 2021, **13**, 26573–26580.
- 23 H. Cao, J. Xue, Z. Wang, J. Dong, W. Li, R. Wang, S. Sun, C. Gao, Y. Tan, X. Zhu and J. Bao, *J. Mater. Chem. A*, 2021, **9**, 16339–16344.
- 24 Z.-W. Wei, H.-J. Wang, C. Zhang, K. Xu, X.-L. Lu and T.-B. Lu, *Angew. Chem., Int. Ed.*, 2021, **60**, 16622–16627.
- 25 P.-P. Luo, X.-K. Zhou, Y. Li and T.-B. Lu, *ACS Appl. Mater. Interfaces*, 2022, **14**, 21069–21078.
- 26 H. Pang, X. Meng, P. Li, K. Chang, W. Zhou, X. Wang, X. Zhang, W. Jevasuwan, N. Fukata, D. Wang and J. Ye, *ACS Energy Lett.*, 2019, **4**, 1387–1393.
- 27 Q. Guo, F. Liang, X.-B. Li, Y.-J. Gao, M.-Y. Huang, Y. Wang, S.-G. Xia, X.-Y. Gao, Q.-C. Gan, Z.-S. Lin, C.-H. Tung and L.-Z. Wu, *Chem*, 2019, **5**, 2605–2616.
- 28 B. Qin, Y. Li, H. Wang, G. Yang, Y. Cao, H. Yu, Q. Zhang, H. Liang and F. Peng, *Nano Energy*, 2019, **60**, 43–51.
- 29 A. Wagner, C. D. Sahm and E. Reisner, *Nat. Catal.*, 2020, **3**, 775–786.
- 30 L. Carbone, C. Nobile, D. M. Giorgi, D. F. Sala, G. Morello, P. Pompa, M. Hytch, E. Snoeck, A. Fiore, R. I. Franchini, M. Nadasan, F. A. Silvestre, L. Chiodo, S. Kudera, R. Cingolani, R. Krahne and L. Manna, *Nano Lett.*, 2007, **7**, 2942–2950.
- 31 J. Yang, J.-H. Zeng, S.-H. Yu, L. Yang, G.-e. Zhou and Y.-t. Qian, *Chem. Mater.*, 2000, **12**, 3259–3263.
- 32 J. Schneider and A. Räuber, *Solid State Commun.*, 1967, **5**, 779–781.
- 33 B. K. Meyer, P. Omling, E. Weigel and G. Müller-Vogt, *Phys. Rev. B*, 1992, **46**, 15135–15138.
- 34 S. D. Setzler, M. Moldovan, Z. Yu, T. H. Myers, N. C. Giles and L. E. Halliburton, *Appl. Phys. Lett.*, 1997, **70**, 2274–2276.
- 35 A. J. Almeida, A. Sahu, A. Riedinger, D. J. Norris, M. S. Brandt, M. Stutzmann and R. N. Pereira, *J. Phys. Chem. C*, 2016, **120**, 13763–13770.
- 36 H. Fujiwara, H. Hosokawa, K. Murakoshi, Y. Wada and S. Yanagida, *J. Phys. Chem. B*, 1997, **101**, 8270–8278.
- 37 W. N. Nan, Y. A. Niu, H. Y. Qin, F. Cui, Y. Yang, R. C. Lai, W. Z. Lin and X. G. Peng, *J. Am. Chem. Soc.*, 2012, **134**, 19685–19693.
- 38 C. Pu and X. Peng, *J. Am. Chem. Soc.*, 2016, **138**, 8134–8142.
- 39 Z.-J. Li, J.-J. Wang, X.-B. Li, X.-B. Fan, Q.-Y. Meng, K. Feng, B. Chen, C.-H. Tung and L.-Z. Wu, *Adv. Mater.*, 2013, **25**, 6613–6618.
- 40 H. H. Wei, C. M. Evans, B. D. Swartz, A. J. Neukirch, J. Young, O. V. Prezhdo and T. D. Krauss, *Nano Lett.*, 2012, **12**, 4465–4471.
- 41 Y. Gao and X. G. Peng, *J. Am. Chem. Soc.*, 2015, **137**, 4230–4235.
- 42 X. Zhang, Z. Zhao, W. Zhang, G. Zhang, D. Qu, X. Miao, S. Sun and Z. Sun, *Small*, 2016, **12**, 793–801.
- 43 C. Pu and X. Peng, *J. Am. Chem. Soc.*, 2016, **138**, 8134–8142.
- 44 D. H. Son, S. M. Hughes, Y. Yin and A. Paul Alivisatos, *Science*, 2004, **306**, 1009–1012.
- 45 P. K. Jain, B. J. Beberwyck, L.-K. Fong, M. J. Polking and A. P. Alivisatos, *Angew. Chem., Int. Ed.*, 2012, **51**, 2387–2390.
- 46 D. D. Zhu, J. L. Liu and S. Z. Qiao, *Adv. Mater.*, 2016, **28**, 3423–3452.