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Construction of organic–inorganic hybrid composites derived from C₃N₅ incorporated with CeO₂ for enhanced photocatalytic hydrogen evolution†

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Energy scarcity and environmental issues can be effectively addressed via photocatalytic hydrogen production. The effective combination of semiconductor materials can prevent exciton recombination, making it a highly effective method for enhancing photocatalytic activity. This study details the synthesis of a conjugated polymer encapsulated with a metal oxide photocatalyst using a simple *ex situ* method. The encapsulation of the polymer with CeO₂ nanoparticles resulted in exceptional performance in H₂ production, exhibiting improved visible light absorption and a significant increase in charge transfer efficiency. This is attributed to the high charge transfer and reduced recombination in the composite. Moreover, photogenerated holes led to a substantial decline in the recombination rate of excitons and concomitant enhancement in the rate of photocatalytic H₂ production. Markedly, the observed hydrogen evolution for 10 wt% of CeO₂ doped C₃N₅ composites is 1256 μmol g⁻¹ h⁻¹, whereas for C₃N₅, it is 125 μmol g⁻¹ h⁻¹. Electrochemical analysis showed that the optimized composites exhibit a low electron–hole recombination rate, and UV-vis spectroscopic analysis showed improved visible light absorption resulting in excellent photocatalytic activity. Notably, the proposed system offers a novel strategy for hydrogen evolution via photocatalysis using CeO₂/C₃N₅ composites. Consequently, this research offers a new perspective on the design of organo–inorganic heterostructures and introduces a novel pathway to explore their catalytic capabilities.

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Introduction

Fossil fuels, including crude oil, natural gas, and coal, play an essential role in satisfying the world's energy needs.^{1,2} It is anticipated that global energy consumption will increase by nearly 55% and CO₂ emissions by 20% by 2024–2050.^{3,4} Global warming and fuel shortages that have resulted from the significant increase in the emission of greenhouse gases and the accelerated depletion of fossil fuels have compelled humanity to identify alternative renewable and pure energy sources.⁵ It is anticipated that the consumption of renewable energy will increase and reach approximately 247 exajoules by 2050. In contrast, the total consumption of renewable energy was 96 exajoules in 2024.⁶ Photocatalytic water splitting using semiconductors has been recognized as a sustainable and prospective green technology that has garnered significant attention owing to its ability to effectively convert sustainable solar energy into environmentally friendly hydrogen. A sustainable method for producing green hydrogen fuel is through photocatalytic water splitting, which utilizes pure solar energy.^{7–9}

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† Electronic supplementary information (ESI) available: Fig. S1. Rietveld analysis of the CeO₂ photocatalyst and composition of the CeO₂ photocatalyst. Fig. S2. FTIR spectra of C₃N₅, CeO₂ and C₃N₅/CeO₂ composites. Fig. S3. (a) SEM image (b) and (c) respective energy-dispersive spectroscopy ratio of the CeO₂/C₃N₅ composite (Ce, O, C, and N elements). Fig. S4. Survey XPS spectra of CeO₂, C₃N₅ and C₃N₅/CeO₂ composites. Fig. S5. High resolution XPS spectra of CeO₂ and C₃N₅ pristine materials: (a) C 1s, (b) N 1s, (c) Ce 3d, and (d) O 1s. Fig. S6. Photoluminescence spectra of the samples. Fig. S7. Mott–Schottky analysis of the samples: (a) CeO₂ and (b) C₃N₅. Fig. S8. XRD analysis of the C₃N₅/CeO₂ composite after and before the reaction study. Table S1. Physicochemical properties of the pristine catalyst used for photocatalytic application. Table S2. Elemental composition of C₃N₅, CeO₂ and C₃N₅/CeO₂ composites. Table S3. Activity comparison of some representative photocatalysts for photocatalytic hydrogen production. See DOI: <https://doi.org/10.1039/d4ya00476k>



Owing to its environmentally friendly and sustainable nature, photocatalysis has garnered significant interest among many other potential technologies. It is based on the photosynthesis process occurring in plants to transform low-density solar energy into high-density energy fuels.^{10–12} To achieve this, research communities are working extremely hard to investigate effective, reliable, affordable, and visible-light sensitive materials. Innovation based on photocatalysis is useful as a viable, sustainable, cost-effective, reliable and steady solution to the world's power issues.^{13–15} Solar energy can be converted into sustainable hydrogen energy *via* photocatalytic water splitting using semiconductors irradiated by visible light, which is a safe and environmentally friendly process. Goodeve and Kitchener (1938) discovered that TiO₂ could enhance dye removal efficiency under both vacuum and air conditions, and TiO₂ remained stable after the completion of the reaction; this marked the beginning of semiconductor-based photocatalysts.^{16,17} Fujishima and Honda (1972) noticed photoelectrical water splitting on TiO₂ films when solar light was being irradiated and the door to semiconductor-based photocatalyst innovation was opened as soon as the world became aware of this amazing accomplishment.¹⁸ However, the activity of the majority of photocatalysts is still constrained by the absence of active sites, severe recombination of photogenerated excitons, and diminished light-harvesting efficiency.^{19,20} Therefore, it is imperative to investigate novel photocatalysts with a broad spectral response that are extremely active to achieve sustainable hydrogen production. The modifiable band diagram, low toxicity, good thermal stability, and chemical stability of graphite carbon nitride (*g*-C₃N₅) have all been the subject of extensive research. In addition to that, the visible light absorption can be expanded and a rapid electron movement can be facilitated by the narrow band gap of *g*-C₃N₅.^{21–26} CeO₂ is a type of rare earth metal oxide that is non-toxic and low-cost having a cubic crystalline n-type material with a 2.5 eV band gap. In the CeO₂ unit cell, every Ce(IV) is linked in the octahedral interstitial with eight adjacent O²⁻, and each O²⁻ is linked with four adjacent Ce(IV), forming a tetrahedral structure; here, Ce³⁺ and Ce⁴⁺ are capable of coexisting indefinitely and converting to each other effortlessly.^{27–29} The efficacy of photocatalysts will be improved, resulting in the multivalence property of CeO₂, which enables the generation of robust interactions with other catalyst components for photocatalytic hydrogen production. Zou *et al.* have developed a CeO₂ hybridised *g*-C₃N₄ composite, which produced a hydrogen yield of 860 μmol g⁻¹ h⁻¹.³⁰ Sha *et al.* have discussed about the charge separation in CeO₂/MnO₂ nanoflakes with an efficiency of 540 μmol g⁻¹ h⁻¹ of hydrogen.³¹ Liu *et al.* have modified C₃N₅ with nickel oxide for improving the activity, which shows a yield of 357 μmol g⁻¹ h⁻¹ of hydrogen production.³² Still, there are no pertinent reports regarding the formation of a heterojunction between C₃N₅ and CeO₂ for photocatalytic hydrogen evolution. The heterostructures obtained by the chemical binding of C₃N₅ and CeO₂ composites effectively segregate and constrain the charge carrier recombination. As anticipated, the well-designed CeO₂/C₃N₅ heterojunction, substantially improves the photocatalytic hydrogen evolution, without the need of additional co-catalysts.

Materials

All chemicals and reagents 3-amino-1,2,4-triazole (C₂H₄N₄) (≥98%), cerium(III) nitrate hexahydrate (Ce(NO₃)₃·6H₂O) (≥98%), methanol (CH₃OH) (≥98%), ethanol (CH₃CH₂OH) (≥99.5%), triethanolamine (C₆H₁₅NO₃) (≥98%), lactic acid (C₃H₆O₃) (≥98%), ascorbic acid (C₆H₈O₆) (≥99%), sodium sulfide (Na₂S) (≥98%), and sodium sulfite (Na₂SO₃) (≥98%) were obtained from Sigma Aldrich, TCI, and SRL India.

Synthesis of C₃N₅

C₃N₅ was synthesized using a thermal polymerization method in a muffle furnace, as described in the literature by Wu *et al.* In a typical synthesis, 2 g of 3-amino-1,2,4-triazole was placed in a porcelain crucible and was gradually calcined at 550 °C with a heating rate of 2 °C min⁻¹ for a period of 3 h. Bulk C₃N₅ is a dark-yellowish product that was collected and crushed into a powder after naturally cooling to room temperature³³ (Scheme 1a).

Synthesis of CeO₂

CeO₂ was synthesized by a thermal polymerization method in a muffle furnace, as described in the literature by Song *et al.* Briefly, 2.0 g cerium(III) nitrate hexahydrate (Ce(NO₃)₃·6H₂O) was calcined at 500 °C for 2 h in the muffle furnace with a heating rate of 2 °C min⁻¹. CeO₂ is a light-yellowish product that was collected and crushed into a powder after naturally cooling to room temperature³⁴ (Scheme 1b).

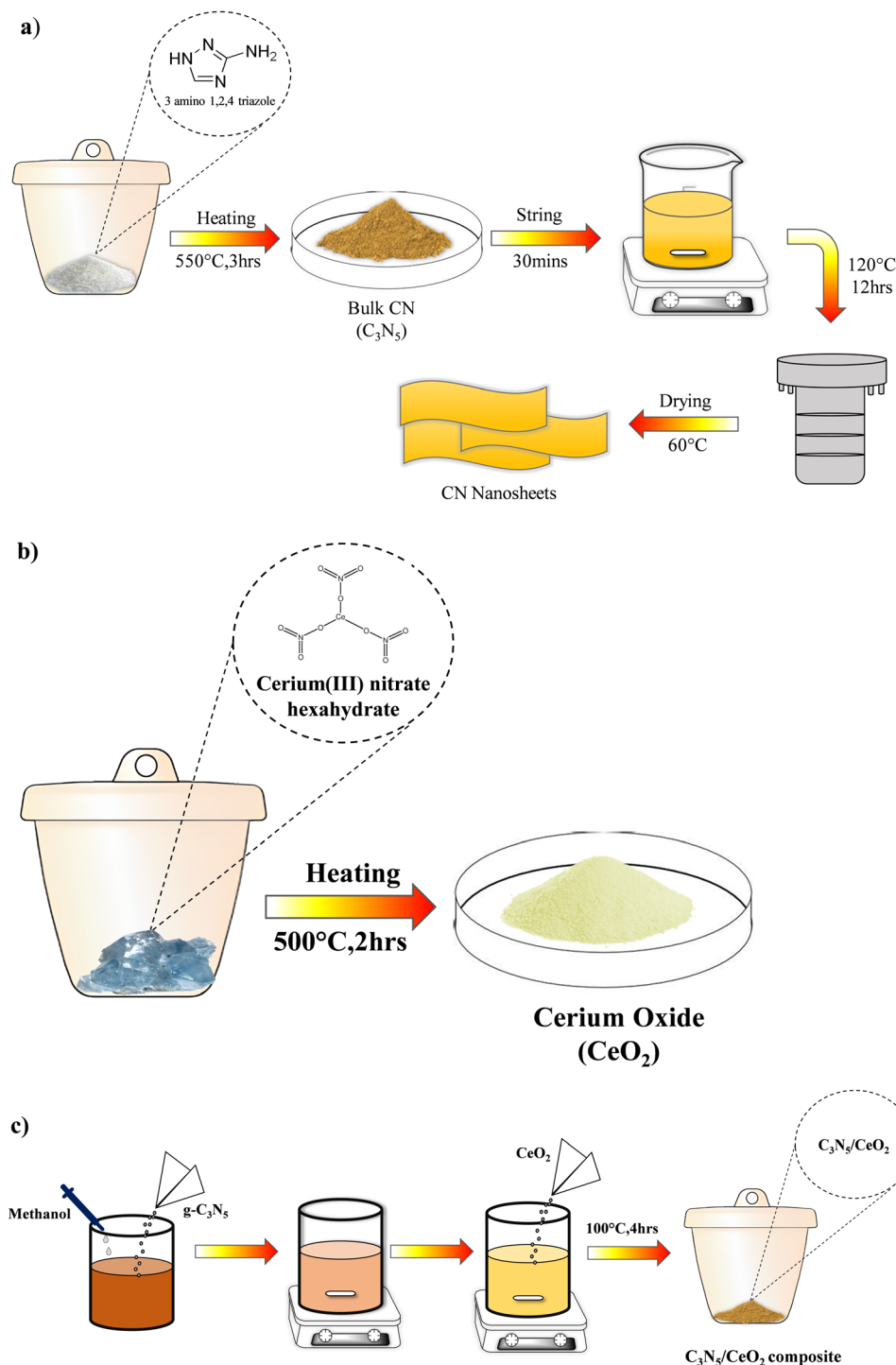
Synthesis of C₃N₅/CeO₂ composite

The C₃N₅/CeO₂ heterostructures were synthesized as follows: 100 mg of C₃N₅ was mixed in 25 mL of methanol under an ultrasonic setting for 30 min, and then a sufficient amount of CeO₂ was immersed into the suspensions. The mixture was vigorously stirred at room temperature, followed by half an hour of ultrasonic treatment to remove the organic solvent. The resultant dark yellowish powder was then calcined for 4 h at 100 °C with a 2 °C min⁻¹ heating rate. The composite catalysts were designated as 5 wt% C₃N₅/CeO₂, 10 wt% C₃N₅/CeO₂, 15 wt% C₃N₅/CeO₂, and 20 wt% C₃N₅/CeO₂ by various weight percentages of the polymer content (5, 10, 15, 20 wt%) in the composite (Scheme 1c). The term “C₃N₅/CeO₂” refers to 10 wt% C₃N₅/CeO₂ in the subsequent text.

Characterisation

The phase purity and crystallinity of the prepared photocatalysts were assessed through X-ray diffraction utilizing a PANalytical X'Pert particle diffractometer with Cu K α radiation ($\lambda = 1.5406$ Å). Shimadzu IRTracer-100 was used to carry out the Fourier-transform infrared spectroscopy (FT-IR). The morphology of the nanocomposites was examined utilizing high-resolution transmission electron microscopy (HRTEM, JEOL Japan, JEM-2100 Plus) and field emission scanning electron microscopy (FE-SEM) (FEI Quanta FEG 200). UV-vis diffuse reflectance (UV-DRS) data were acquired through the utilization of a Shimadzu UV-3600i Plus UV-vis spectrophotometer. The photoluminescence (PL) spectra were analysed using a fluorescence spectrophotometer (Horiba, Fluorolog-QM).





Scheme 1 (a). Schematic representation of the synthesis of C₃N₅. (b) Schematic representation of the synthesis of CeO₂. (c) Schematic representation of the synthesis of the C₃N₅/CeO₂ composite.

Photoelectrochemical measurements

The CHI 760E electrochemical workstation was employed to acquire the Mott–Schottky, impedance and transient photocurrent responses. Ag/AgCl and a platinum wire serve as the reference and counter electrode, respectively. As an electrolyte, a 0.5 M Na₂SO₄ aqueous solution is utilized. The slurry was prepared through the deposition of 2.5 g of the catalyst

dispersed in a mixture of deionized water and anhydrous ethanol (1:1) and coated on the working electrode (glassy carbon electrode) for electrochemical studies.

Photocatalytic studies

The photocatalytic hydrogen generation reaction was performed in direct sunlight. The reaction was carried out in a



Kjeldhal flask containing 5 mg of the catalyst dispersed in 50 ml of 5% sacrificial agent. The dissolved gases were eliminated from the reaction solution by purging with nitrogen gas for 15 minutes. The reaction setup was kept under sunlight, and an offline gas chromatograph (Shimadzu GC-2014 equipped with a Molecular Sieve [5 Å column]) analysed hydrogen generation with a TCD detector after the reaction setup was kept in sunlight for a period of 3 hours. The mean solar light intensity was ascertained to be 80 000 lux using a lux meter. While the pH of the reaction mixture was adjusted using solutions of NaOH and H₂SO₄, various sacrificial agents were employed when necessary.

Results and discussion

Fig. 1 shows the crystal structures of CeO₂, C₃N₅, and the 10 wt% CeO₂/C₃N₅ nanocomposite revealed by the XRD patterns (Fig. 1a). The CeO₂ peaks in the XRD pattern can be found at 28.6°, 33.1°, 47.5°, 56.4°, and 59.1°, 69.2°, 76.6° and 79.2°, which correspond to the (111), (200), (220), (311), (222), (400), (331) and (420) planes, respectively, of the fluorite crystal

structure (JCPDS 34-0394) (Fig. 1b). The large peak (002) in the C₃N₅ XRD pattern, which is centered at 27.5 degrees, reveals the material's amorphous nature (JCPDS 87-1526). The XRD pattern of the CeO₂/C₃N₅ nanocomposites reveals peaks that correlate to the synthesis of both CeO₂ and C₃N₅. Rietveld analysis of CeO₂ is obtained to understand the atomic percentage of the elements (Fig. S1, ESI†) composition. In the nanocomposite, the peak intensities of CeO₂ increased and the C₃N₅ peak intensity decreased, indicating some degree of crystallographic disorder brought on by the addition of C₃N₅. The stretching vibration of the surface free amino group and the hydroxyl group of water is represented by the wide peak of pure C₃N₅ at 3000–3600 cm⁻¹ in (Fig. S2, ESI†). The typical stretching vibration of C–N and C–N heterocycles is responsible for the peaks at 1235 cm⁻¹, 1316 cm⁻¹, 1411 cm⁻¹, 1571 cm⁻¹, and 1637 cm⁻¹. The characteristic stretching vibration of the s-triazazine ring unit is represented by the peak at 805 cm⁻¹. The characteristic peaks of C₃N₅ and CeO₂ remained unaltered in the composite, as evidenced by the presence of all characteristic vibration peaks in the corresponding CeO₂/C₃N₅ composites.

The UV-Vis diffuse reflectance spectra of the modified samples are illustrated in Fig. 1(c). The CeO₂ shows an absorption

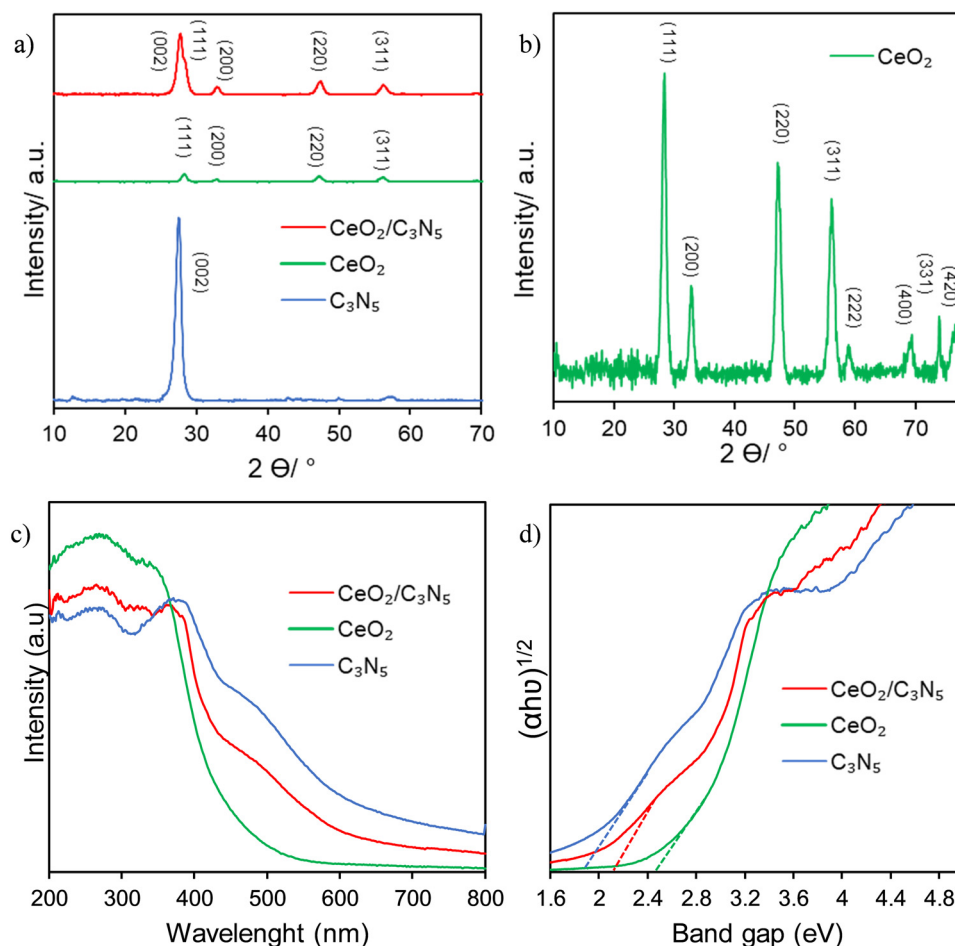


Fig. 1 (a) XRD patterns of C₃N₅, CeO₂ and CeO₂/C₃N₅ composites. (b) XRD patterns of CeO₂. (c) UV-vis absorbance spectra and (d) optical band gap using relative Tauc plots.



below 500 nm. Nevertheless, the photo response of exposed CeO_2 is expanded to the visible region when it is combined with C_3N_5 . The band gap positions of CeO_2 and $g\text{-C}_3\text{N}_5$ are calculated as 2.5 eV and 1.9 eV, respectively, using the Tauc plot, as illustrated in Fig. 1(d). This may be attributed to the fact that C_3N_5 effectively functions in absorbing significant quantities of visible light particles. This confirms the efficacy of C_3N_5 for visible light harvesting.

The morphologies of CeO_2 , C_3N_5 and $\text{CeO}_2/\text{C}_3\text{N}_5$ have been closely observed using SEM analysis. The SEM images show the presence of both CeO_2 incorporated into C_3N_5 . The impact of CeO_2 on the morphological structure of C_3N_5 is clearly visible in the composite (Fig. 2(a–c)). The TEM images reveal the composite formation between CeO_2 and C_3N_5 in Fig. 2(e–g). As demonstrated, the micromorphology of $g\text{-C}_3\text{N}_5$ sheets is clearly observed, and the presence of CeO_2 particles is confirmed. The influence of CeO_2 decoration on the structure of C_3N_5 is clearly visible in the TEM images. Fig. 2(e) shows clear (111) and (220) crystal planes with lattice fringes with a spacing of 0.21 nm and 0.27 nm, respectively. The crystallite size of CeO_2 and C_3N_5 is calculated as 7.13 nm and 8.55 nm (Table S2, ESI[†]). The acquired images confirmed the existence of crystallized CeO_2 nanoparticles encircled by an exceedingly thin layer of C_3N_5 , which reflects the composite's extremely low concentration. (Fig. S3, ESI[†]). The images show that the CeO_2 nanoparticles were evenly distributed on the host photocatalyst, which shows that the CeO_2 nanoparticles were evenly distributed on the host photocatalyst, which is a sign of a good deposition process.

By employing X-ray photoelectron spectroscopy (XPS), the electronic chemical states and surface elemental composition of CeO_2 , C_3N_5 and $\text{CeO}_2/\text{C}_3\text{N}_5$ were ascertained (Fig. 3). The

CeO_2 , C_3N_5 and $\text{CeO}_2/\text{C}_3\text{N}_5$ composite, as indicated by the XPS survey spectra presented in Fig. S4 (ESI[†]), comprise the elements Ce, O, C and N. The C 1s spectra of C_3N_5 depict the binding energies at 284.9 eV, 287.6 eV and 288.3 eV, corresponding to C–C/C=C, N=C and C–NH bonds, respectively. In the high resolution of N 1s spectra, two main peaks at 402.3 eV and 404.1 eV are attributed to CN=C and C–N=N–C groups, respectively. In addition, the peak at 406.9 eV is generated by N–H bonds. In the O 1s spectra, the peak labelled at 528.8 eV is attributed to the lattice oxygen in CeO_2 . Apart from this, a broad peak is deconvoluted at 532.3 eV and 538 eV, which is attributed to oxide defects and adsorbed oxygen, respectively. The high-resolution Ce 3d spectrum of CeO_2 is deconvoluted. Ce^{4+} is the dominant state, and its characteristic spectrum is represented by the peak with high intensity at 914.2 eV. Furthermore, the Ce 3d_{5/2} spin-orbit is represented by the peaks at 879.8 eV, 884.7 eV and 887.1 eV. The Ce 3d_{3/2} spin-orbit is shown by the peaks at 903.7 eV, 900.4 eV, 899.3 eV, and 895.4 eV (Fig. S5, ESI[†]). The sample surface is not completely oxidized, as evidenced by the presence of $\text{Ce}^{3+}/\text{Ce}^{4+}$ oxidation states on the surface of $\text{CeO}_2/\text{C}_3\text{N}_5$ composites (Table S2, ESI[†]).

The $\text{CeO}_2/\text{C}_3\text{N}_5$ composite is more mobile and has a greater resistance to charge recombination, as indicated by the reduced arc radius of the electrochemical impedance spectroscopy (EIS) Nyquist plot in comparison to that of CeO_2 and C_3N_5 (Fig. 4a). Mott–Schottky analyses were performed on CeO_2 and $g\text{-C}_3\text{N}_5$ to clarify the flat band potential of the pristine materials. Possibly, the positive slope represented an n-type semiconductor (Fig. 4c and d). In contrast to the redox potential of H^+/H_2 , the flat band positions of C_3N_5 and CeO_2 concerning the Ag/AgCl electrode were -1.42 eV and -1.08 eV, respectively,

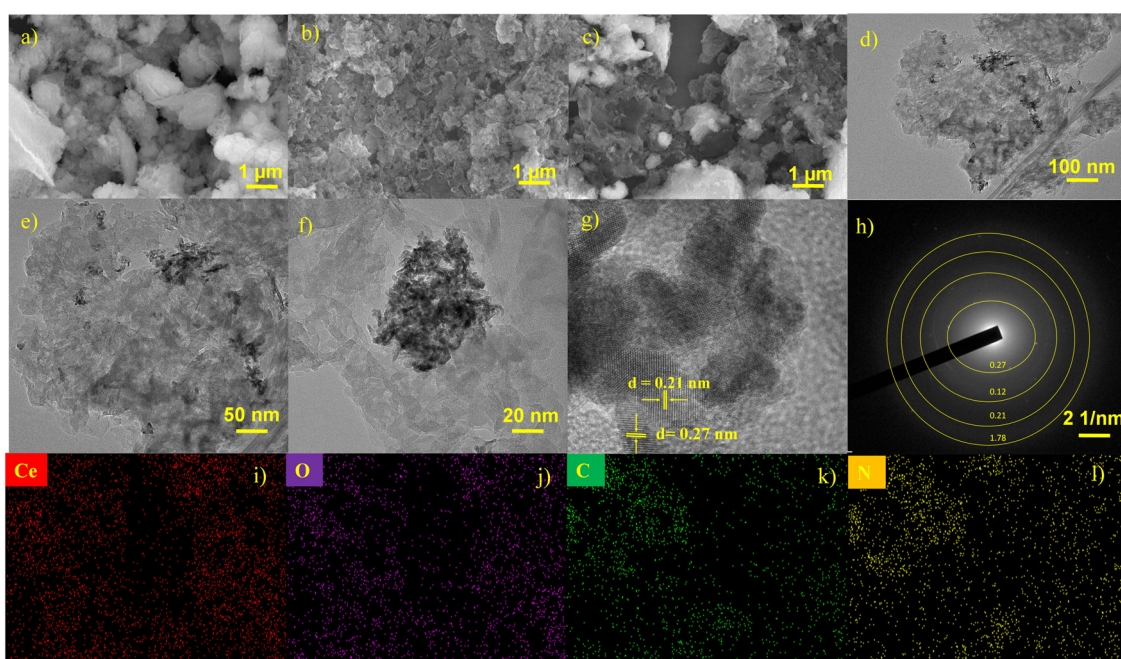


Fig. 2 SEM images of (a) CeO_2 , (b) C_3N_5 , and (c) $\text{CeO}_2/\text{C}_3\text{N}_5$ composites; (d)–(g) high-resolution TEM (HRTEM) images of $\text{CeO}_2/\text{C}_3\text{N}_5$ composites; (h) clear fringes with lattice spacings. (i)–(l) EDS mapping images of Ce, O, C, and N elements.



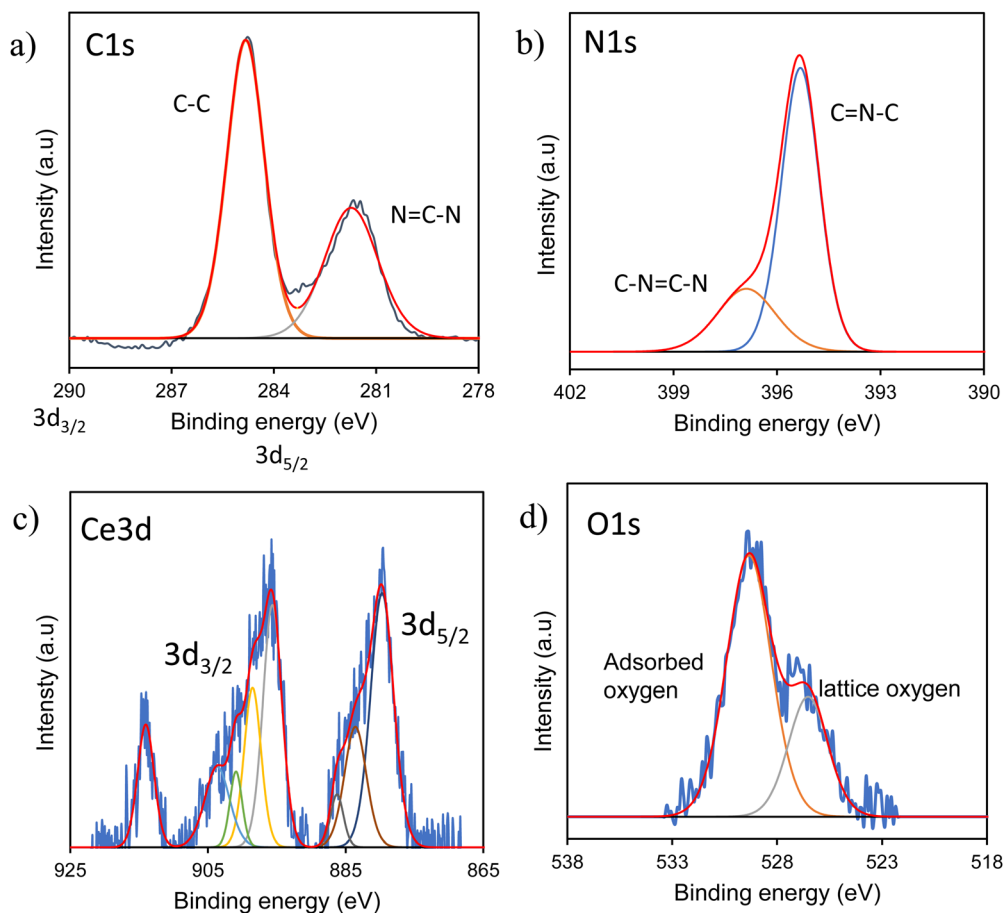


Fig. 3 High-resolution XPS spectra of $\text{CeO}_2/\text{C}_3\text{N}_5$ composites: (a) C 1s, (b) N 1s, (c) Ce 3d, and (d) O 1s.

both of which were more negative (eqn (S1), ESI[†]). The conduction band potential (CB) and flat band in n-type semiconductors are well established. Additionally, as demonstrated by the transient photocurrents in Fig. 4(b), $\text{CeO}_2/\text{C}_3\text{N}_5$ demonstrates a significantly higher photocurrent density than the pristine materials when subjected to intermittent light irradiation. The improved charge transfer and separation efficacy of the $\text{CeO}_2/\text{C}_3\text{N}_5$ nanocomposite is illustrated by its increased photocurrent response, which is accompanied by a decreased rate of photogenerated charge carrier recombination. The PL intensity of the composite is lower than that of individual counterparts, which implies that it has a higher charge carrier separation efficiency (Fig. S6, ESI[†]). In summary, the photoelectrochemical data indicated that the photoelectron transport efficacy of the catalyst can be enhanced through the formation of a $\text{CeO}_2/\text{C}_3\text{N}_5$ nanocomposite.

Photocatalytic activity

In the presence of a hole scavenger in deionized water, the photocatalytic efficiency was evaluated in response to direct sunlight (Fig. 5).

The optimization of the $\text{CeO}_2/\text{C}_3\text{N}_5$ composite, exhibiting notable photocatalytic activity, has been carried out using pristine CeO_2 , C_3N_5 , 5 wt% $\text{CeO}_2/\text{C}_3\text{N}_5$, 10 wt% $\text{CeO}_2/\text{C}_3\text{N}_5$,

15 wt% $\text{CeO}_2/\text{C}_3\text{N}_5$, and 20 wt% $\text{CeO}_2/\text{C}_3\text{N}_5$ composites. The 10 wt% $\text{CeO}_2/\text{C}_3\text{N}_5$ composite produced an efficiency of $1256 \mu\text{mol g}^{-1} \text{h}^{-1}$ (Fig. 5a). This value is considerably greater than that of the pristine material ($125 \mu\text{mol g}^{-1} \text{h}^{-1}$) and other composites. With increasing CeO_2 , the rate of H_2 evolution initially increases and then decreases. Under visible-light irradiation, the H_2 evolution rate of the 10 wt% $\text{CeO}_2/\text{C}_3\text{N}_5$ heterojunction can reach its maximum value with the loading amount of CeO_2 . The values obtained are comparatively higher than those of the published articles in the field of photocatalytic hydrogen production, which is discussed in Table S1 (ESI[†]). The photocatalytic hydrogen evolution is carried out using different scavengers, including methanol, TEOA, glycerol and lactic acid, producing yields of 1290, 510, 1032, and $227 \mu\text{mol g}^{-1} \text{h}^{-1}$, respectively (Fig. 5b). In order to demonstrate the stability of the catalyst, the $\text{CeO}_2/\text{C}_3\text{N}_5$ sample was subjected to four consecutive cycles (Fig. 5c), and the outcomes show that the sample exhibits high stability and hydrogen production. XRD characterization studies confirm the structural integrity of the catalyst after repeated use (Fig. S8, ESI[†]).

Finally, the electronic charge transfer, advantages of C_3N_5 integrating with CeO_2 and formation of type II heterojunctions were understood based on the aforementioned results and prior experimental investigations. A potential mechanism for



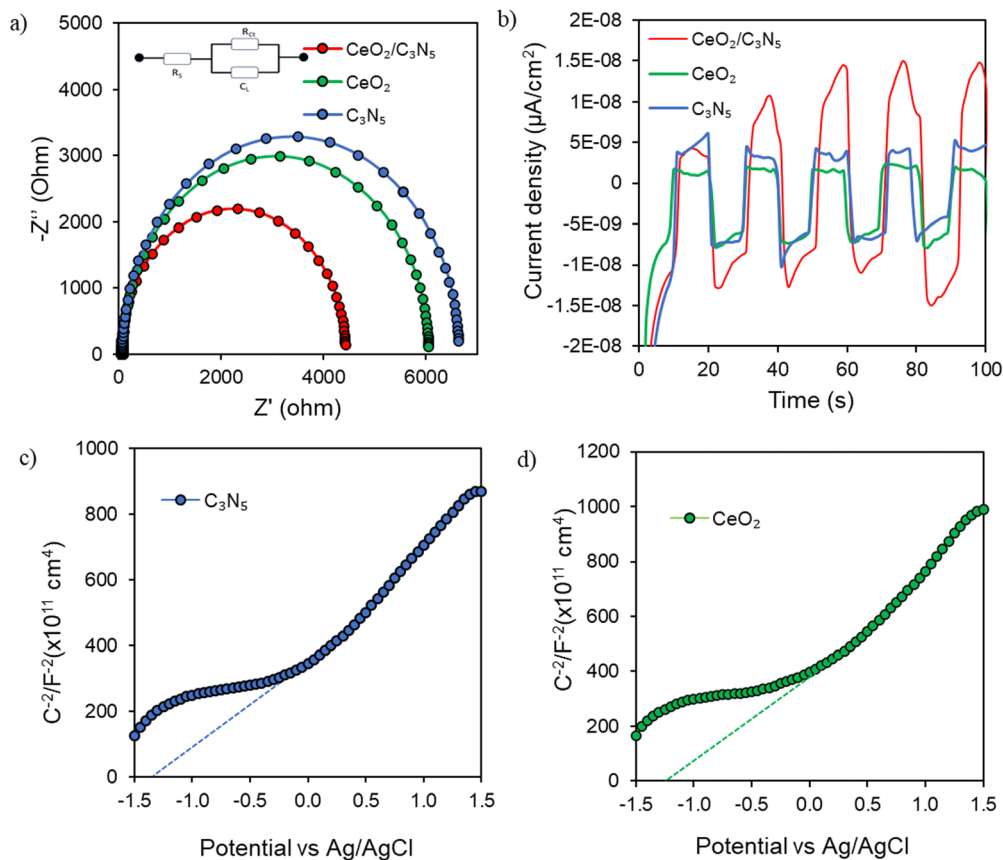


Fig. 4 (a) Impedance spectra plots, (b) photocurrent response, and (c) and (d) Mott–Schottky analysis of the samples.

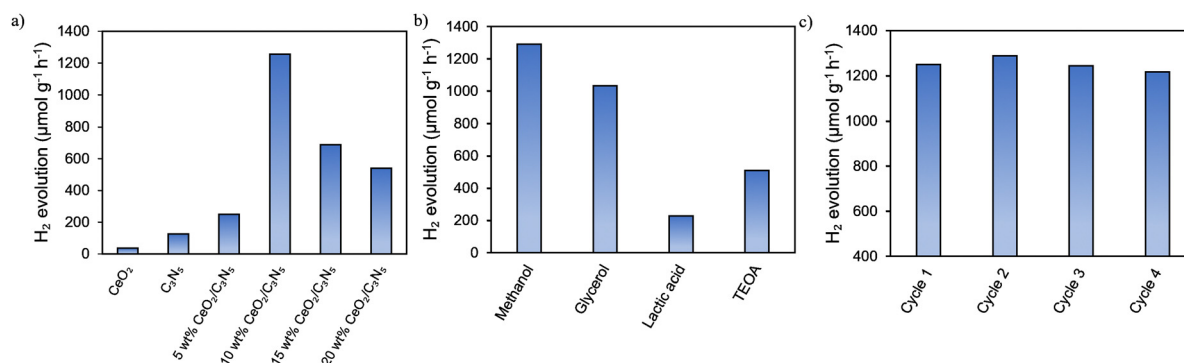


Fig. 5 (a) Photocatalytic H_2 evolution rates of the $\text{CeO}_2/\text{C}_3\text{N}_5$ composite. (b) Photocatalytic H_2 evolution rate with different scavengers. (c) Stability study.

the 10 wt% $\text{CeO}_2/\text{C}_3\text{N}_5$ composite is suggested in (Fig. 6) based on the observed results. e^- is excited from the valence band (VB) to the conduction band (CB) of C_3N_5 and CeO_2 by high-energy photons under light irradiation. Using the Mott Schottky analysis, the flat band positions of C_3N_5 and CeO_2 with respect to the Ag/AgCl electrode were -1.42 eV and -1.08 eV, respectively, in contrast to the redox potential of H^+/H_2 . The relationship between the flat band and the conduction band potential (CB) in n-type semiconductors is well-established. The conduction band edges were -0.81 eV and -0.47 eV, as calculated from the Tauc plot analysis (eqn (S2), ESI[†]). Therefore, a type II heterojunction

mechanism was suggested for the $\text{CeO}_2/\text{C}_3\text{N}_5$ composite in response to the band levels of C_3N_5 and CeO_2 . The conduction band accumulates excited electrons (e^-) in response to photon irradiation, while the valence band of C_3N_5 and CeO_2 accumulates holes (h^+). Due to variations in energy levels, the accumulated electrons migrate from the CB of C_3N_5 to the CB of CeO_2 . At the same time, the holes from the valence band of CeO_2 are transferred to the valence band of C_3N_5 . At the CB of CeO_2 , the reduction of H^+ to hydrogen occurs concurrently, while the oxidation of the hole scavenger occurs at the VB of C_3N_5 . Consequently, the synergistic effect of C_3N_5 and CeO_2 can



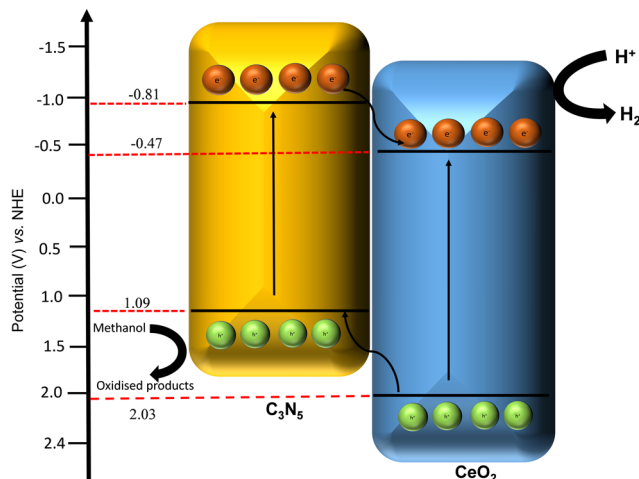


Fig. 6 Plausible mechanism of the $\text{CeO}_2/\text{C}_3\text{N}_5$ catalyst.

effectively separate photoelectron-hole pairs, enticing a greater number of carriers to participate in the photodegradation process, thereby improving the photocatalytic activity.

Conclusion

In summary, the CeO_2 doped C_3N_5 composite is the product of a straightforward synthesis method, which led to the successful development of CeO_2 -incorporated C_3N_5 . The primary goal of this research is to investigate the impact of organic-inorganic hybrid photocatalysts by optimizing the catalyst to enhance the reaction performance. According to the results of the morphological investigation, the composition and morphological transformation of the synthesized samples are significantly affected by CeO_2 dispersed on C_3N_5 . Moreover, the optical band gap decreased, leading to an increased absorption of the visible spectrum. In addition, photoelectrochemical methods have consistently exhibited an effective ability to prevent the recombination of photogenerated electron-hole pairs. Notably, the photocatalytic hydrogen evolution for the 10 wt% CeO_2 doped C_3N_5 catalyst is $1256 \mu\text{mol g}^{-1} \text{h}^{-1}$, which is 10 times greater than that of C_3N_5 alone ($125 \mu\text{mol g}^{-1} \text{h}^{-1}$). Due to the efficient transfer of photogenerated holes, the recombination rate of excitons has been significantly reduced with the simultaneous enhancement of the rate of photocatalytic H_2 production. This study offers a unique perspective that aids in creating innovative photocatalysts for efficient solar energy conversion into fuel.

Data availability

The data supporting this article have been included within the manuscript and as part of the ESI.†

Conflicts of interest

The authors declare no conflict of interest.

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References

- 1 S. Chu and A. J. Majumdar, *Nature*, 2012, **488**, 294–303.
- 2 T. Hisatomi and K. J. Domen, *Nat. Catal.*, 2019, **2**, 387–399.
- 3 L. Wang, S. Duan, P. Jin, H. She, J. Huang, Z. Lei, T. Zhang and Q. J. Wang, *Appl. Catal., B*, 2018, **239**, 599–608.
- 4 A. Naldoni, M. Altomare, G. Zoppellaro, N. Liu, S. Kment, R. Zboril and P. J. Schmuki, *ACS Catal.*, 2018, **9**, 345–364.
- 5 X. Yin, X. Zhao and W. J. Zhang, *Energy*, 2018, **158**, 1204–1212.
- 6 M. I. Alhamid, Y. Daud, A. Surachman, A. Sugiyono, H. Aditya, T. J. R. Mahlia and S. E. Reviews, *Renewable Sustainable Energy Rev.*, 2016, **53**, 733–740.
- 7 C. Xia, H. Wang, J. K. Kim and J. J. Wang, *Adv. Funct. Mater.*, 2021, **31**, 2008247.
- 8 W. Zhou, W. Li, J.-Q. Wang, Y. Qu, Y. Yang, Y. Xie, K. Zhang, L. Wang, H. Fu and C. S. Zhao, *J. Am. Chem. Soc.*, 2014, **136**, 9280–9283.
- 9 D. Ma, D. Sun, Y. Zou, S. Mao, Y. Lv, Y. Wang, J. Li and J.-W. Shi, *J. Colloid Interface Sci.*, 2019, **549**, 179–188.
- 10 A. Augustin, C. Chuaicham, M. Shanmugam, B. Vellaichamy, S. Rajendran, T. K. Hoang, K. Sasaki and K. J. Sekar, *Nano-scale Adv.*, 2022, **4**, 2561–2582.
- 11 L. Yu, G. Zhang, C. Liu, H. Lan, H. Liu and J. J. A. C. Qu, *ACS Catal.*, 2018, **8**, 1090–1096.
- 12 Y. Zhang, D. Ma, J. Li, C. Zhi, Y. Zhang, L. Liang, S. Mao and J.-W. J. C. C. R. Shi, *Coord. Chem. Rev.*, 2024, **517**, 215995.
- 13 M. Shanmugam, C. Chuaicham, A. Augustin, K. Sasaki, P. J. Sagayaraj and C. Sekar, *New J. Chem.*, 2022, **46**, 15776–15794.
- 14 M. S. Yesupatham, A. Augustin, N. Agamendran, B. H. Honnappa, M. Shanmugam, P. J. Sagayaraj, T. Ganesan, N. C. S. Selvam and K. J. Sekar, *Sustainable Energy Fuels*, 2023, **7**, 4727–4757.
- 15 D. Ma, J. Chen, J. Li, X. Ji and J.-W. Shi, *J. Mater. Chem. A*, 2024, **12**, 12293–12324.
- 16 C. Goodeve and J. J. Kitchener, *Trans. Faraday Soc.*, 1938, **34**, 570–579.
- 17 Y. Zou, S. Li, D. Zheng, J. Feng, S. Wang, Y. Hou and G. J. Zhang, *Sci. China Chem.*, 2024, 1–9.
- 18 A. Fujishima and K. J. n Honda, *Nature*, 1972, **238**, 37–38.
- 19 C. Hu, Y.-H. Lin, M. Yoshida and S. J. Ashimura, *ACS Appl. Mater.*, 2021, **13**, 24907–24915.
- 20 D. Ma, Z. Zhang, Y. Zou, J. Chen and J.-W. J. Shi, *Coord. Chem. Rev.*, 2024, **500**, 215489.
- 21 B. Debnath, S. Singh, S. M. Hossain, S. Krishnamurthy, V. Polshettiwar and S. J. L. Ogale, *Langmuir*, 2022, **38**, 3139–3148.



- 22 H. Wang, M. Li, Q. Lu, Y. Cen, Y. Zhang and S. J. Yao, *ACS sustainable Chem.*, 2018, **7**, 625–631.
- 23 P. Kumar, E. Vahidzadeh, U. K. Thakur, P. Kar, K. M. Alam, A. Goswami, N. Mahdi, K. Cui, G. M. Bernard and V. K. Michaelis, *J. Am. Chem. Soc.*, 2019, **141**, 5415–5436.
- 24 Q. Wang, S. Li, D. Zheng, S. Wang, Y. Hou and G. J. Zhang, *ACS Appl. Energy Mater.*, 2024, **7**, 6090–6095.
- 25 Q. Wang, G. Zhang, W. Xing, Z. Pan, D. Zheng, S. Wang, Y. Hou and X. J. Wang, *Angew. Chem.*, 2023, **135**, e202307930.
- 26 B. K. Raja, M. Govindaraj, M. K. Muthukumaran and S. J. Ramar, *New J. Chem.*, 2024, **48**, 15324–15337.
- 27 D. Gao, Y. Zhang, Z. Zhou, F. Cai, X. Zhao, W. Huang, Y. Li, J. Zhu, P. Liu and F. J. Yang, *J. Am. Chem. Soc.*, 2017, **139**, 5652–5655.
- 28 Y. Wang, Z. Chen, P. Han, Y. Du, Z. Gu, X. Xu and G. J. A. C. Zheng, *ACS Catal.*, 2018, **8**, 7113–7119.
- 29 J. Tian, Y. Sang, Z. Zhao, W. Zhou, D. Wang, X. Kang, H. Liu, J. Wang, S. Chen and H. J. Cai, *Small*, 2013, **9**, 3864–3872.
- 30 W. Zou, Y. Shao, Y. Pu, Y. Luo, J. Sun, K. Ma, C. Tang, F. Gao and L. J. Dong, *Appl. Catal., B*, 2017, **218**, 51–59.
- 31 M. A. Sha, G. Mohanan, L. Elias, T. Bhagya and S. J. Shibli, *Mater. Chem. Phys.*, 2023, **294**, 127019.
- 32 M.-Y. Liu, J.-Y. Wang, D. Lian, L. Xian and L. J. Zhang, *J. Fuel Chem.*, 2022, **50**, 243–249.
- 33 B. Wu, T. Sun, N. Liu, L. Lu, R. Zhang, W. Shi and P. J. Cheng, *ACS Appl. Mater. Interfaces*, 2022, **14**, 26742–26751.
- 34 J. Song, F. Wu, Y. Lu, X. Zhang and Z. J. Li, *ACS Appl. Nano Mater.*, 2021, **4**, 4800–4809.

