


Cite this: *Nanoscale*, 2022, **14**, 5612

Super-resolution imaging of photogenerated charges on CdS/g-C₃N₄ heterojunctions and its correlation with photoactivity†

Shuyang Wu,^a Jinn-Kye Lee,^a  Pei Chong Lim,^a Rong Xu  and Zhengyang Zhang *^a

Construction of heterostructures is an effective way to improve photo-induced charge separation and photocatalytic performance. Among various structures, type II and direct Z-scheme heterojunctions with distinct charge separation mechanisms are the two typical representatives attracting much research attention. Here we prepared type II and Z-scheme CdS/g-C₃N₄ nanocomposites by thermal treatment and self-assembly chemisorption methods, respectively. High-resolution microscopy techniques including (scanning) transmission electron microscopy (TEM/STEM) and super-resolution fluorescence microscopy (SRM) were used to investigate the charge distribution and flow mechanism. The charge tracking results reveal that the nanocomposite prepared by thermal treatment has a type II heterostructure with charges flowing in the opposite direction, while the self-assembly sample possesses a Z-scheme structure. It was found that the type II system exhibited the lowest charge migration resistance and the best charge separation ability and stability of photoactivity, leading to the highest H₂ generation rate of 2410 μmol h⁻¹ g⁻¹. The SRM technique was applied for the first time to map the reactive sites of type II and Z-scheme structures at nanometer resolution. The photoactive species (*i.e.*, e⁻ and h⁺) were found to be preferentially distributed at the two end segments of CdS nanorods and the edge boundaries of g-C₃N₄. Therefore, our findings shed more light on the charge distribution and photocatalytic heterogeneity of composite materials at the nanoscale. Such results would provide guidance on optimizing nanocomposite properties and help to design better photocatalysts for efficient solar-to-chemical energy conversion.

Received 18th January 2022,

Accepted 14th March 2022

DOI: [10.1039/d2nr00316c](https://doi.org/10.1039/d2nr00316c)
rsc.li/nanoscale

1. Introduction

Photocatalytic water splitting has been considered as a promising way to produce renewable H₂ energy, since Fujishima and Honda's pioneer work on water splitting in 1972.¹ Among various semiconductor materials, functional 2D materials have attracted enormous research interests owing to their unique structural features. Compared with 0D and 1D materials, 2D nanosheets with large lateral size provide numerous active sites for surface adsorption of reactants and photoreactions.^{2,3} When forming heterostructures, the large surface, especially with functional groups would facilitate the anchoring and accommodation of guest components and help to construct

multifunctional 3D structures. Laminar g-C₃N₄ is one of the typical 2D nanomaterials, which has many advantages in photocatalytic applications.⁴ A layered structure bonded by weak van der Waals force largely facilitates morphology modification, which can be easily decorated with other materials. The high stability and earth-abundant resources for synthesis and a simple preparation route make g-C₃N₄ a promising catalyst for various applications.⁵ However, despite these merits, pristine g-C₃N₄ suffers from significant drawbacks of high charge recombination rates and limited utilization of visible light, which restrict its photocatalytic efficiency. Hence, many strategies such as a good morphology design, defect control, element doping, and coupling with other components have been applied to overcome these limitations.^{6,7}

Fabrication of heterojunctions has been proved an effective way to enhance the charge separation efficiency of g-C₃N₄-based photocatalysts.^{8,9} Numerous semiconductors have been investigated to form heterojunctions with g-C₃N₄, such as metal oxides (*e.g.*, ZnO, TiO₂ and Cu₂O),^{10,11} metal sulfides (*e.g.*, NiS and CdS),^{12,13} metal oxyhalides (*e.g.*, BiOCl and BiOBr)^{14,15} and other semiconductors (*e.g.*, perovskites and layered double

^aDivision of Chemistry and Biological Chemistry, School of Physical and Mathematical Sciences, Nanyang Technological University, 21 Nanyang Link, Singapore 637371. E-mail: zhang.zy@ntu.edu.sg

^bSchool of Chemical and Biomedical Engineering, Nanyang Technological University, 62 Nanyang Drive, Singapore 637459

†Electronic supplementary information (ESI) available. See DOI: [10.1039/d2nr00316c](https://doi.org/10.1039/d2nr00316c)



hydroxides).^{16,17} Based on the charge separation mechanism, these heterojunctions can be generally classified into two types: a type II heterostructure and an all-solid-state Z-scheme heterostructure. In the type II heterostructure, photogenerated electrons (e^-) and holes (h^+) kinetically flow in the opposite directions, leading to enhanced spatial charge separation.¹⁸ In contrast, the Z-scheme configuration maintains e^- and h^+ at relatively higher redox potentials, accompanied by a certain charge extinction with or without redox mediators.¹⁹ The Z-scheme system without charge mediators is called a direct Z-scheme, in which the two semiconductors have a direct interfacial contact. In this report, we construct CdS/g-C₃N₄ composites with these two heterostructures by coupling CdS nanorods and g-C₃N₄ nanosheets. CdS is chosen due to its small bandgap energy (~2.4 eV) which has light absorption up to 520 nm.²⁰ It has been widely used as a photosensitizer for wide-bandgap semiconductors like g-C₃N₄, which significantly extends the visible light response.²¹ Although it has been validated that CdS/g-C₃N₄ composites can be successfully constructed due to their well-matched band alignment,²² a direct observation of photo-reactions and mapping of photoactive species on CdS/g-C₃N₄ at the nanoscale have not been reported. Therefore, there is a great need to apply a spatially resolved technique to map the reactive sites at nanometer resolution and correlate the microscopic study with the bulk activity measurement.

Super-resolution fluorescence microscopy (SRM) such as point accumulation for imaging in nanoscale topography (PAINT) and stochastic optical reconstruction microscopy (STORM) have been applied as optical imaging techniques that can achieve nanoscale spatial resolution.^{23,24} These methods can quantify catalysis by characterizing the single-turnover events and localize the fluorescent molecules (*i.e.*, reactive sites) down to nanoscale resolution. Sambur *et al.* applied super-resolution imaging to map photoelectrocatalytic activities driven by electrons and holes in single TiO₂ nanorods. It was discovered that the site-selective deposition of an oxygen evolution catalyst could enhance the overall activity of a nanorod, which provided an effective strategy to rationally engineer photoelectrodes.²⁵ Ha *et al.* unveiled two different charge separation mechanisms on single Au–CdS hybrid catalysts using superlocalization fluorescence imaging.²⁶ It proposed a new strategy for engineering the charge flow direction by controlling the incident light wavelength. Hence, such results prove SRM an effective technique to study the catalytic kinetics and photo-excited charge distribution on a single nanocatalyst. However, previous works have not revealed such nature and photocatalytic properties on type II and Z-scheme heterostructures. Besides, the photoredox probing method has been applied to detect the e^- and h^+ locations by photo-deposition of metal or metal oxides,²⁷ but this method has the limitation of low spatially resolved resolution and cannot visualize single-turnover events. Therefore, it is highly desirable to apply the SRM technique to further explore the charge distribution and photocatalytic heterogeneity in composite materials at the nanoscale.

Herein, we prepared type II and Z-scheme CdS/g-C₃N₄ nanocomposites and investigated their photocatalytic performance

for H₂ generation under visible light. The charge separation mechanisms were systematically studied by photo-deposition and SRM methods, which revealed two distinct charge migration routes for the CdS/g-C₃N₄ composites. In addition, the SRM technique was applied for the first time to unveil the distribution of reactive sites on type II and Z-scheme systems at the nanoscale. It was found that the two end segments of CdS nanorods and the edge boundaries of g-C₃N₄ have higher reactivity than other parts probably due to the less saturated surface sites. The bulk photoactivity measurement indicates a superior photocatalytic activity and stability of the type II structure to the Z-scheme one owing to the improved charge separation and moving holes away from CdS to avoid photocorrosion. Through the combination of microscopic study and bulk property measurement, this work provides a deeper insight into the understanding of “structure–activity” correlations, which benefits the morphology design, interface engineering, and electronic band modulation and the control of charge and energy flows in nanocomposites.

2. Experimental section

2.1. Synthesis of the photocatalysts

Preparation of g-C₃N₄ nanosheets. g-C₃N₄ nanosheets were facilely synthesized by thermal polymerization of urea at an air atmosphere.²⁸ In a typical procedure, a 50 g portion of urea (Sigma, 99.0–100.5%) was placed in a crucible with a cover and then heated in the muffle furnace at 550 °C for 3 h with a ramping rate of 0.5 °C min^{−1}. After the thermal treatment and cooling down, the product was ground into powder without further treatment.

Preparation of CdS nanorods. CdS nanorods were synthesized through a solvothermal route.²⁹ Typically, 4.66 g of cadmium nitrate tetrahydrate (Sigma, ≥99.0%) and 3.45 g of thiourea (Sigma, ≥99.0%) were dispersed in 70 mL of ethylenediamine (Sigma, ≥99%). The solution was placed in an autoclave and then heated at 160 °C for 48 h. After cooling down, the yellow product was washed with ethanol and DI water several times to remove the organic residues.

Preparation of CdS/g-C₃N₄ by thermal treatment (CS/CN-II). A CS/CN-II sample was prepared *via* a mixing and thermal treatment route.¹⁸ In a typical process, 300 mg of CdS nanorods and 200 mg of g-C₃N₄ (mass ratio of CdS/g-C₃N₄ = 3 : 2) were dispersed in a solution containing 15 mL of DI water and 15 mL of ethanol. Subsequently, the suspension was sonicated for 2 h and then dried in an oil bath at 60 °C with stirring. The dried powder was heated in a muffle furnace at 300 °C for 2 h with a ramping rate of 5 °C min^{−1}. A yellow product was finally obtained and denoted as CS/CN-II.

Preparation of CdS/g-C₃N₄ by self-assembly chemisorption (CS/CN-Z). A CS/CN-Z sample was synthesized by a self-assembly chemisorption method.¹⁹ Typically, 200 mg of g-C₃N₄ was dispersed in 80 mL of DI water and sonicated for 1 h. Then, 300 mg of CdS nanorods were added into the suspension (mass ratio of CdS/g-C₃N₄ = 3 : 2), followed by sonication treat-



ment for another 1 h. Subsequently, the suspension was stirred for 24 h at room temperature. After washing with DI water several times and dried at 60 °C, the final product was obtained and denoted as CS/CN-Z.

2.2. Materials characterization

The crystal structure was analyzed by X-ray diffraction (XRD) using a Bruker D2 Phaser diffractometer with Cu K α irradiation ($\lambda = 1.54184$ Å) as the incident beam at 30 kV and 10 mA. Field emission scanning electron microscopy (FESEM) images were obtained on a JEOL JSM 6701F microscope. Transmission electron microscopy (TEM) images were taken on a JEOL JEM-2100Plus microscope, and the high-angle annular dark-field (HAADF) scanning TEM (STEM) images were obtained on a JEOL JEM-ARM200F microscope. UV-vis diffuse reflection spectroscopy (DRS) spectra were recorded on a Shimadzu UV-2450 spectrophotometer using BaSO₄ as the reference. X-ray/ultraviolet photoelectron spectroscopy (XPS/UPS) was performed on a Shimadzu Kratos Axis Supra spectrometer equipped with an Al K α source (1486.7 eV) for XPS and a He-I photon source (21.21 eV) for UPS. The XPS binding energies were referenced to the C 1s peak at 284.7 eV. Fourier transform infrared spectroscopy (FTIR) was conducted on a PerkinElmer spectrophotometer (Spectrum 100). Inductively coupled plasma (ICP) measurement was performed on Prodigy High Dispersion ICP spectrometer. In a typical measurement, a Pt-loaded sample was first treated in H₂ at 300 °C for 4 h to fully convert PtO_x to Pt⁰. Subsequently, Pt⁰ was dissolved in aqua regia to form Pt⁴⁺ and then analyzed by an ICP instrument.

2.3. Photoelectrochemical measurement

The photocurrent was measured in a conventional three-electrode system composed of a working electrode (photocatalyst-deposited ITO glass), a reference electrode (Ag/AgCl) and a Pt counter electrode, which was controlled by an electrochemical workstation (CHI 660E). To prepare the working electrode, 5 mg of photocatalyst was dispersed in a 1 mL solution containing 980 μ L of ethanol and 20 μ L of 5 wt% Nafion. Then, after ultrasonication for 30 min, 40 μ L of the as-prepared solution was deposited on ITO glass with an area of 0.196 cm². Subsequently, the deposited electrode was dried in an 80 °C oven overnight. The light source was a 300 W Xe lamp (Newport) equipped with a cut-off filter ($\lambda > 400$ nm). The photocurrent was measured in a 0.5 M Na₂SO₄ solution over several ON/OFF irradiation cycles at 0.6 V bias voltage. EIS was conducted at a voltage of 10 mV applied to the electrode and in the frequency range of 0.1 Hz–10⁵ Hz under irradiation.

2.4. Photocatalytic measurement

Photocatalytic H₂ generation was conducted in a 300 mL Pyrex glass reactor connected to a gas circulation and evacuation system. In a typical measurement, 50 mg of photocatalyst and 500 μ L of 1 mg Pt mL⁻¹ H₂PtCl₆ solution (1 wt% Pt) were added into a 20 vol% lactic acid aqueous solution (100 mL). The suspension was subsequently ultrasonicated for 15 min. The light source was a 300 W Xe lamp (Newport) equipped with a cut-off

filter ($\lambda > 400$ nm). The temperature of the reactor was maintained at 18 °C by a cycle cooling water system. Before irradiation, the air in the reactor was fully removed by evacuating and purging Ar several times. The reactor was finally refilled with Ar to reach 30 Torr. The evolved H₂ was detected and measured using an online gas chromatograph (Agilent 6890N) equipped with a thermal conductivity detector. The stability of the photocatalysts was evaluated by four consecutive runs for 6 h for each run under the same conditions. The H₂ evolved after each run was fully removed by evacuating and purging the reactor with Ar several times.

2.5. Selective photo-deposition experiment

Photocatalysts with 1 wt% Pt deposited were obtained from the suspension after the photocatalytic activity measurement. The suspension was centrifuged, washed with DI water and then dried in an oven at 60°. For photo-deposition of 3 wt% PbO₂, the experiment was conducted under the same conditions, but using Pb(NO₃)₂ as the precursor and NaIO₃ as the electron scavenger. In a typical process, 50 mg of photocatalyst and a calculated amount of Pb(NO₃)₂ were added into an aqueous NaIO₃ solution (0.01 M, 100 mL). Then, the suspension was irradiated under a Xe lamp ($\lambda > 400$ nm) for 6 h. A 3 wt% PbO₂ loaded photocatalyst was obtained after washing with DI water and drying in an oven.

2.6. Super-resolution fluorescence imaging

The single-molecule fluorescence images were obtained on a total internal reflection fluorescence (TIRF) microscope (Nikon Eclipse Ti-U). In a typical procedure, 30 μ L of photocatalyst aqueous suspension (5 mg mL⁻¹) was drop-cast onto a cleaned glass coverslip and dried in an oven. Then the photocatalyst-deposited coverslip was gently rinsed with DI water to remove the unbound compounds and fully dried in an oven at 80 °C. The coverslip was mounted onto a cell holder and then transferred to the microscope stage. 1 mL of 50 nM non-fluorescent probe (*i.e.*, resazurin or Amplex Red) solution was added to the cell. The conventional brightfield images were collected without irradiation. To obtain the fluorescence images, circularly polarized 532 nm laser (20 mW, L6CC Oxxius) and 405 nm laser (2 mW) were used to excite the probe molecules and photocatalysts, respectively. The emission signal was filtered by a band-pass filter (ET605/70 m, Chroma) and then collected through an oil-immersion objective lens (Nikon Plan Apo λ 100 \times , NA 1.45) via a quad-edge laser dichroic mirror (Di03-R405/488/532/635-t1-25x36, Semrock). The fluorescence images were captured by an electron multiplying charge coupled device (EMCCD) camera (Andor iXon3) at 60 Hz. A movie composed of around 30 000 frames was collected for each run and finally analyzed by the software.

3. Results and discussion

3.1. Physicochemical properties of the as-prepared photocatalysts

XRD analysis was conducted to determine the crystal phase structures of CdS, g-C₃N₄, CS/CN-II and CS/CN-Z. As shown in



Fig. S1,[†] pure CdS exhibits the characteristic peaks at 24.8°, 26.5°, 28.2°, 36.6°, 43.7°, 47.8°, 51.9°, and 52.8°, which correspond to the (100), (002), (101), (102), (110), (103), (112) and (201) planes of the hexagonal wurtzite structure of CdS (JCPDS no. 41-1049), respectively. Pure g-C₃N₄ has two diffraction peaks at 13.1° and 27.4°, which belong to the (100) and (002) planes of graphite-like g-C₃N₄ (JCPDS no. 87-1526), respectively. The former with a weak intensity is ascribed to the in-plane packing of tri-s-triazine rings. The latter strong one is related to the interplanar stacking of conjugated aromatic rings.¹⁹ The binary composites, CS/CN-II and CS/CN-Z, show similar XRD patterns as that of CdS and no evident g-C₃N₄ characteristic peaks are observed in the composites. Such a result may arise from the less g-C₃N₄ percentage in the composites, lower crystallinity of g-C₃N₄ and peak overlapping with CdS. This phenomenon was also reported in the previous lit-

erature for CdS/g-C₃N₄ composites.^{13,30,31} However, the presence of g-C₃N₄ can be confirmed by other characterization methods such as FESEM, TEM, XPS and FTIR. Besides, the crystallinity of CS/CN-II is enhanced compared to that of CS/CN-Z due to calcination at elevated temperature (300 °C).

FESEM and TEM were used to study the morphology of the samples. In Fig. S2a and S2b,[†] pure CdS nanorods can be clearly identified. Pure g-C₃N₄ nanosheets have graphene-like laminar morphology, and are transparent because of their ultrathin nature (Fig. S2c and S2d[†]). The morphologies of the CdS/g-C₃N₄ composites are shown in Fig. 1. The FESEM and TEM images in Fig. 1a and b demonstrate that the g-C₃N₄ nanosheets and CdS nanorods are tightly adhered with each other in CS/CN-II. Therefore, the heterojunctions with closely contacted interfaces are formed between the two components, which could greatly improve the photogenerated charge separ-

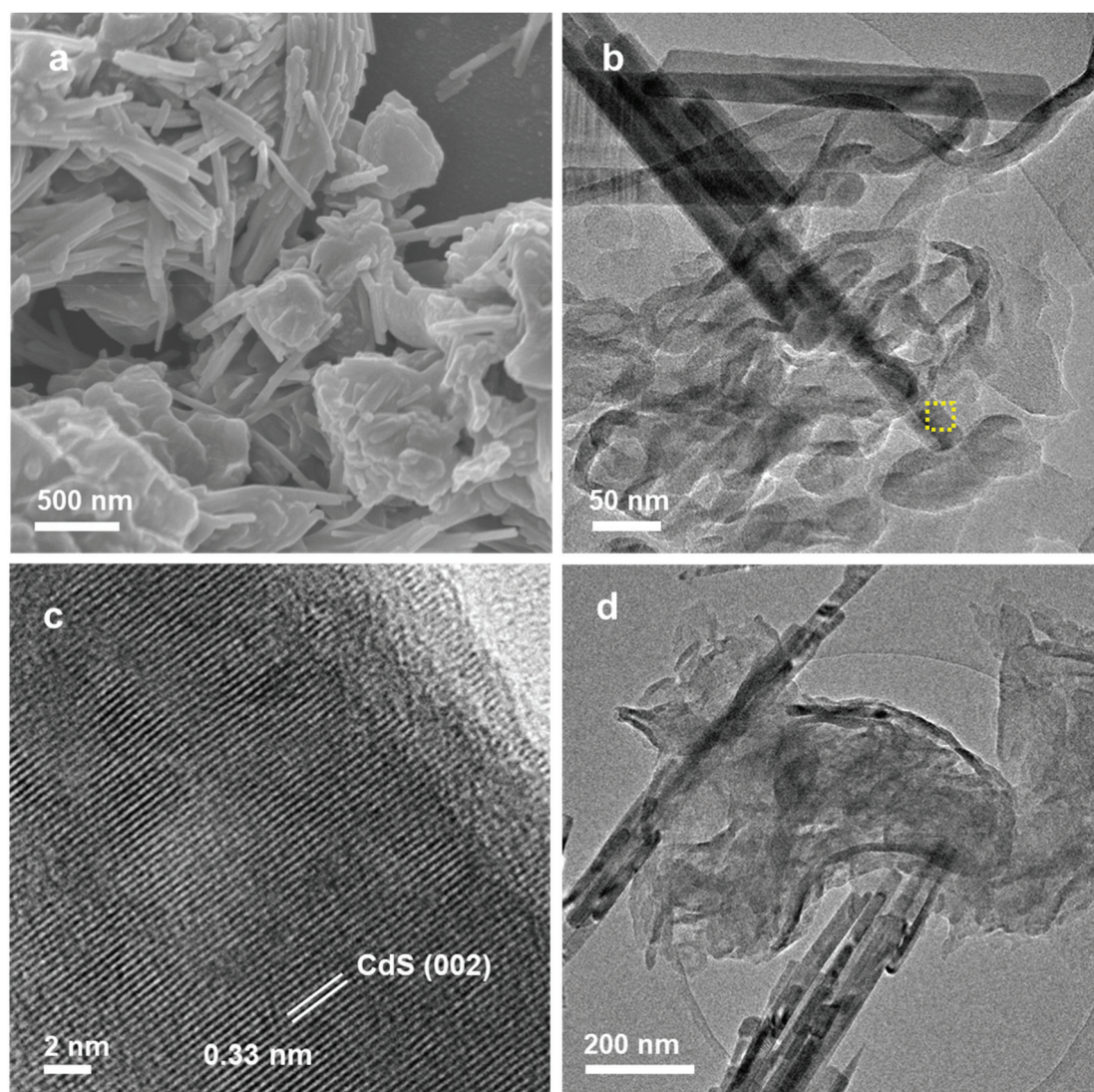


Fig. 1 Morphologies of CS/CN-II and CS/CN-Z. (a) FESEM image, (b) TEM image and (c) HRTEM image of CS/CN-II ((c) is the close-up of the marked areas (yellow dashed line) in (b)). (d) TEM image of CS/CN-Z.



ation. In Fig. 1c, the lattice spacing of 0.33 nm can be indexed to the (002) planes of hexagonal CdS, which is in accordance with the XRD results (Fig. S1†). In Fig. 1d, the CS/CN-Z sample exhibits a similar morphology to CS/CN-II. The heterostructure between the thin-layered g-C₃N₄ and CdS nanorods can be clearly identified. Hence, the FESEM and TEM results confirm that the synthesis of CS/CN-II by the calcination method and CS/CN-Z through the chemisorption route was successful.

The composition and chemical valence of the as-prepared samples were analyzed by XPS. The survey spectra in Fig. S3† confirm the presence of elements in the pure and composite materials. High-resolution XPS spectra were obtained for CS/CN-II. In the C 1s spectra (Fig. S4a†), the peak located at 284.7 eV for CS/CN-II is ascribed to the sp² C-C bonds in g-C₃N₄. The peak at 288.2 eV corresponds to the hybridized carbon in N-C=N.³² Fig. S4b† shows the N 1s spectra which are deconvoluted into three peaks. The peaks at 398.7 eV, 399.7 eV and 400.8 eV can be assigned to the sp²-bonded N in the triazine rings (C-N=C), tertiary nitrogen (N-(C)₃) and amino functional groups (C-N-H), respectively.³³ The characteristic bands for Cd 3d at 405.3 eV and 412.0 eV further confirm that the chemical valence state of the Cd element is Cd²⁺ (Fig. S4c†). The S 2p peaks at 161.5 eV and 162.8 eV belong to the S²⁻ ions in CdS nanorods (Fig. S4d†). Besides, it can be noticed that the XPS peaks of CS/CN-II have a slight

shift to the high-energy region compared to those of pure CdS and g-C₃N₄. This is attributed to the formation of hetero-junctions with a tight interface between the two components in CS/CN-II.¹⁸

The interaction of CdS nanorods with g-C₃N₄ nanosheets was also investigated by FTIR. In Fig. 2a, for pure CdS, the absorption bands located in the ranges of 700–820 cm⁻¹ and 1250–1660 cm⁻¹ arise from the Cd-S bonds.^{34,35} For pure g-C₃N₄, the broad band at 3150 cm⁻¹ belongs to the stretching mode of the N-H bonds. The characteristic bands between 1200 and 1750 cm⁻¹ are related to the skeletal stretching vibration of aromatic g-C₃N₄ heterocycles.³⁶ Additionally, the band located at 809 cm⁻¹ is ascribed to the out-of-plane bending vibration mode of g-C₃N₄ triazine rings.³¹ For the CdS/g-C₃N₄ composites, it is worth noting that this vibration mode has red shifts of 3 cm⁻¹ and 13 cm⁻¹ for CS/CN-Z and CS/CN-II, respectively (inset of Fig. 2a). The red shift of this characteristic band suggests that the bond strengths of C-N and C=N are weakened due to the strong interaction between CdS and g-C₃N₄.³⁷ As the CS/CN-Z sample was prepared through the chemisorption and self-assembly method by mechanically mixing at room temperature, a weak interaction was found between the two compounds, resulting in a small red shift of the characteristic band. In contrast, the CS/CN-II sample has a large red shift of 13 cm⁻¹. This is probably

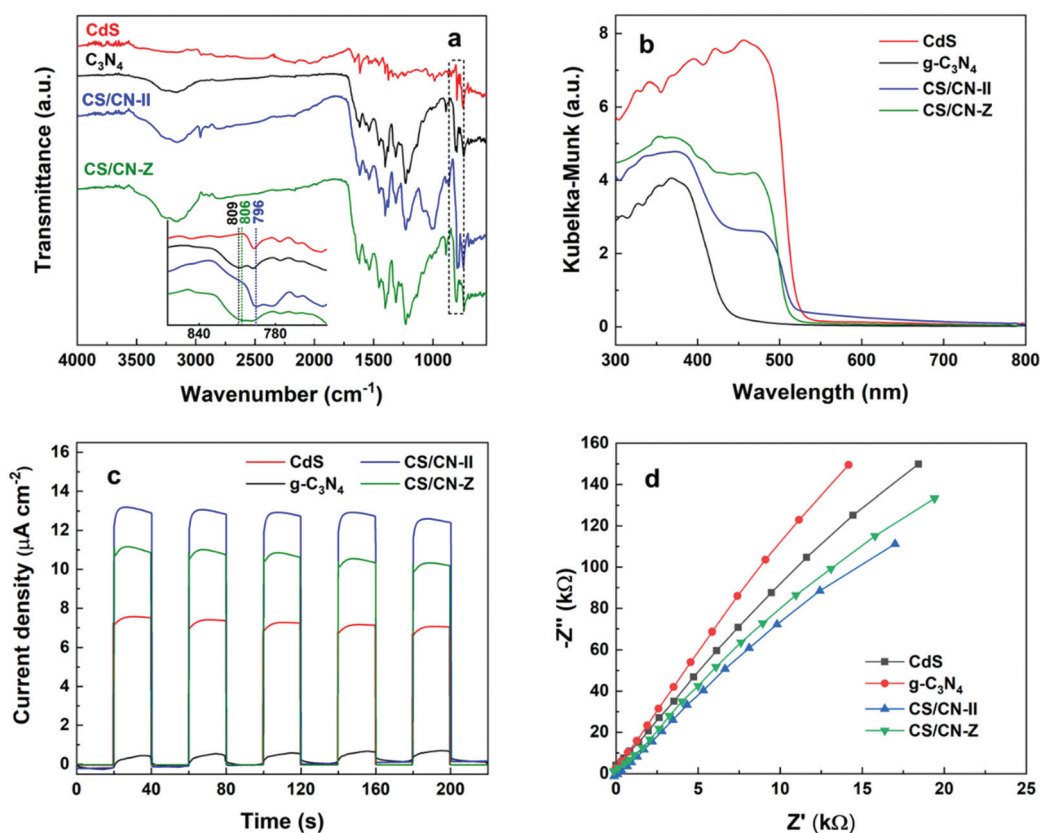


Fig. 2 Physicochemical and photophysical properties of the as-prepared samples. (a) FTIR spectra (inset: enlarged spectra in the range of 865–740 cm⁻¹), (b) UV-vis DRS spectra, (c) transient photocurrent density and (d) EIS spectra of CdS, g-C₃N₄, CS/CN-II and CS/CN-Z.



because the thermal treatment (300 °C) in the synthetic process could provide an intense binding strength between CdS and g-C₃N₄, which weakens the bonds between the C and N atoms. Therefore, both FTIR spectra and XPS results demonstrate that a closely contacted interface and an intense interaction between CdS and g-C₃N₄ were developed in CS/CN-II, which are stronger than those developed in CS/CN-Z.

The optical absorption spectra obtained by UV-vis DRS are shown in Fig. 2b. Pure g-C₃N₄ and CdS exhibit the absorption edges at 435 and 525 nm, respectively, due to their intrinsic electronic properties. For composite materials, the incorporation of CdS plays a key role in sensitizing g-C₃N₄ and improving its optical response to visible light. Hence, the absorptions of CS/CN-II and CS/CN-Z are notably extended to the visible light region due to synergistic absorption. Compared to the physically mixed CdS/g-C₃N₄ sample with an absorption threshold of 507 nm, the absorption edges of CS/CN-Z and CS/CN-II are red-shifted to 512 and 521 nm, respectively which are close to that of CdS (Fig. S5†). This proves the formation of heterojunctions at the interface of CdS and g-C₃N₄ and the more intense interaction leads to a larger band edge shift. The bandgap energies (E_g) of CdS and g-C₃N₄ were obtained by fitting the DRS data using the Tauc/Davis–Mott model.³⁸ As revealed by the Tauc plot in Fig. S6,† the bandgap energies of CdS and g-C₃N₄ were found to be 2.36 and 2.85 eV, respectively. The bandgap energies of the composite materials are between those of CdS and g-C₃N₄, and closer to that of CdS due to the photosensitization by CdS. Moreover, the valence band (VB) potential (E_{VB}) was measured by UPS. In Fig. S7a and S7b,† the VB top positions of CdS and g-C₃N₄ were measured to be 1.83 and 1.64 eV, respectively *versus* the Fermi levels of the samples. Therefore, based on the formula $E_g = E_{VB} - E_{CB}$, the conduction band (CB) potentials (E_{CB}) of CdS and g-C₃N₄ are found to be −0.53 and −1.21 eV. Besides, the E_{CB} and E_{VB} values of CdS and g-C₃N₄ can be also determined by the following equations:³⁹

$$E'_{CB} = \chi - E^e - 1/2E_g \quad (1)$$

$$E'_{VB} = E_g + E'_{CB} \quad (2)$$

where χ is the electronegativity of the semiconductor, which is the geometric average of the absolute electronegativity of the constituent atoms and E^e is the energy of free electrons on the hydrogen scale (~4.5 eV). According to $E_g = 2.36$ eV for CdS, the calculated E'_{CB} and E'_{VB} values for CdS based on eqn (1) and (2) are −0.50 and 1.86 eV, respectively. Similarly, for g-C₃N₄ with $E_g = 2.85$ eV, the E'_{CB} and E'_{VB} values for g-C₃N₄ are found to be −1.22 and 1.63 eV, respectively. Hence, the experimental values of E_{CB} and E_{VB} for CdS and g-C₃N₄ are very close to the calculated results using the equations (E'_{CB} and E'_{VB}), which proves the good accuracy of the electronic band measurement.

The transient photocurrent was measured to study the charge separation properties of photocatalysts. As shown in Fig. 2c, all the samples exhibit a reproducible photocurrent during each ON/OFF irradiation cycle. The composite

materials show a higher photocurrent density than those of pure CdS and g-C₃N₄. This indicates that the formation of heterojunctions can significantly improve the charge separation. CS/CN-II possesses the highest photocurrent density, which is about 23% higher than that of CS/CN-Z. This result demonstrates that the type II heterostructure of CdS/g-C₃N₄ has a superior charge separation ability to the Z-scheme one. Furthermore, the electron impedance spectroscopy (EIS) study provides the information of charge migration kinetics at the photoelectrode/electrolyte interface. Generally, the smaller arc radius of EIS spectra indicates the lower charge migration resistance and faster charge transfer rate at the interface.⁴⁰ In Fig. 2d, the CS/CN-II sample has the smallest arc radius among the samples, suggesting the lowest interfacial charge transfer resistance. The large radius of g-C₃N₄ can be attributed to the low electrical conductivity of the C–N network. The results of photocurrent and EIS spectra exhibit a similar trend, which signifies that the construction of heterojunctions can efficiently facilitate the charge separation and transportation. The type II composite shows a superior photoelectrochemical performance to the Z-scheme system. Such properties would eventually have a significant impact on their photocatalytic activities.

3.2. Photocatalytic activity

The photocatalytic activity was evaluated in a closed gas circulation system under visible light illumination ($\lambda > 400$ nm). To optimize the conditions of photoreactions, H₂ generation was first measured in aqueous solutions containing different scavengers such as methanol, lactic acid, triethanolamine (TEOA) and Na₂S/Na₂SO₃ solutions. In Fig. S8,† the photocatalyst in a 20 vol% lactic acid solution (pH = 3) exhibits the highest H₂ evolution, suggesting that lactic acid is the optimal sacrificial reagent for the as-prepared samples. This is consistent with the conclusions reported by previous works, in which lactic acid was used as a hole scavenger for CdS-based photocatalysts.^{36,41–43} As shown in Fig. 3a, CS/CN-II exhibits the highest H₂ generation rate of 2410 $\mu\text{mol h}^{-1} \text{g}^{-1}$ with 1 wt% Pt loaded as a cocatalyst. The evolution rate is 33% higher than that of CS/CN-Z and about 11 times that of pure g-C₃N₄. It is not surprising to note that the trend of H₂ production of the samples is in good agreement with those of photocurrent and EIS (Fig. 2c and d). The excellent charge separation and rapid charge migration in CS/CN-II lead to a remarkable H₂ generation activity. The performance is better than or comparable to some of the best CdS/g-C₃N₄ photocatalysts reported for visible-light-driven H₂ generation (Table S1†). Pure CdS exhibits much higher photocatalytic performance than that of g-C₃N₄ due to the extended visible light absorption (Fig. 2b). The stabilities of H₂ generation on CS/CN-II, CS/CN-Z and CdS were tested for four consecutive runs under the same conditions. In Fig. 3b, the photocatalytic activity of CS/CN-II decreased 12% after four runs, indicating a good stability of CS/CN-II. In contrast, after 24 h reaction, the performances of CS/CN-Z and CdS decreased by 29% and 44%, respectively (Fig. 3c and d). The decrease in the activity of CdS-based



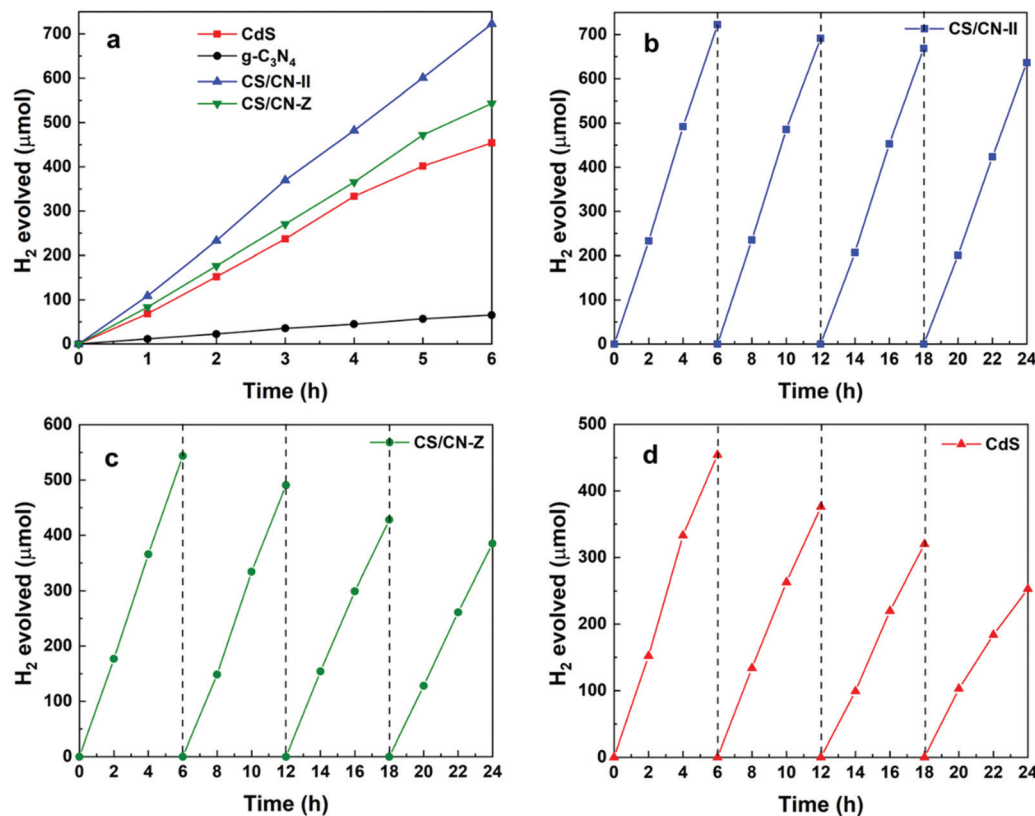


Fig. 3 Photocatalytic activity and stability of the samples. (a) Photocatalytic H₂ generation for 6 h over the as-prepared samples (50 mg of photocatalysts, 1 wt% Pt, 20 vol% lactic acid, 300 W Xe lamp, >400 nm). Stability tests of (b) CS/CN-II, (c) CS/CN-Z and (d) CdS under the same conditions.

catalysts is mainly attributed to the accumulation of photo-generated h^+ on CdS.⁴⁴ S^{2-} ions are very susceptible to oxidation by h^+ during the photoreaction. CS/CN-II has a good stability because h^+ migrate from the CdS VB to the g-C₃N₄ VB, which will be discussed in the later parts. Hence, photocorrosion can be effectively suppressed when h^+ transport away from the sulfide. For CS/CN-Z and pure CdS, the photo-oxidation reaction mainly occurs at CdS which makes the photocatalyst very unstable, leading to a severe photocorrosion and activity decrease.

3.3. Photo-deposition of Pt/PbO₂ on CS/CN-II and CS/CN-Z

To study the charge separation properties of CdS/g-C₃N₄ composites, photoreduction and photo-oxidation probing reactions were performed to monitor the photogenerated e^- and h^+ distributions. HAADF STEM and high-resolution TEM (HRTEM) images were obtained for CS/CN-II loaded with 1 wt% Pt. As shown in Fig. 4a and b and S9a,† the Pt nanoparticles evolved from the photoreduction of Pt⁴⁺ by electrons were located on CdS nanorods. The lattice spacing of 0.225 nm is assigned to the (111) plane of metallic Pt and the planar space of 0.33 nm corresponds to CdS (002) planes (Fig. S9a†). This confirms that Pt nanoparticles are loaded on the CdS surface due to the accumulated electrons. Correspondingly, to probe the distribution of photo-excited

h^+ , photo-deposition of PbO₂ was also conducted. PbO₂ is formed by photo-oxidation of Pb²⁺ with IO₃⁻ used as an electron scavenger. In Fig. 4c and S9b,† it can be seen that the PbO₂ clusters are selectively deposited on g-C₃N₄ nanosheets, especially at the edge positions. This suggests that holes preferentially accumulate at the edge of g-C₃N₄ rather than the basal planes, leading to a higher reactivity of edges than that of basal planes. The lattice spacing of 0.28 nm in the HRTEM image can be ascribed to the (101) planes of PbO₂ (inset of Fig. S9b†). The results of probing reactions demonstrate that the photo-induced e^- and h^+ are spatially separated on CS/CN-II. Electrons transfer to the CB of CdS which is ready for the photoreduction reactions (*i.e.*, formation of metallic Pt or H₂ generation). Meanwhile, the holes flow to the VB of g-C₃N₄ accounting for the photo-oxidation reactions (*i.e.*, oxidation of Pb²⁺ or sacrificial reagents). These results confirm the formation of a type II heterostructure in CS/CN-II. Furthermore, the probing reactions were also performed on CS/CN-Z. As shown in Fig. S9c,† Pt nanoparticles with (111) facets were deposited on the g-C₃N₄ nanosheets. The HAADF image in Fig. 4d clearly shows that numerous tiny Pt nanoparticles were selectively distributed on g-C₃N₄ instead of CdS. More Pt nanoparticles are found at the edge of g-C₃N₄ with fewer identified at the basal planes. This demonstrates that the edge of g-C₃N₄ could facilitate localizing the elec-



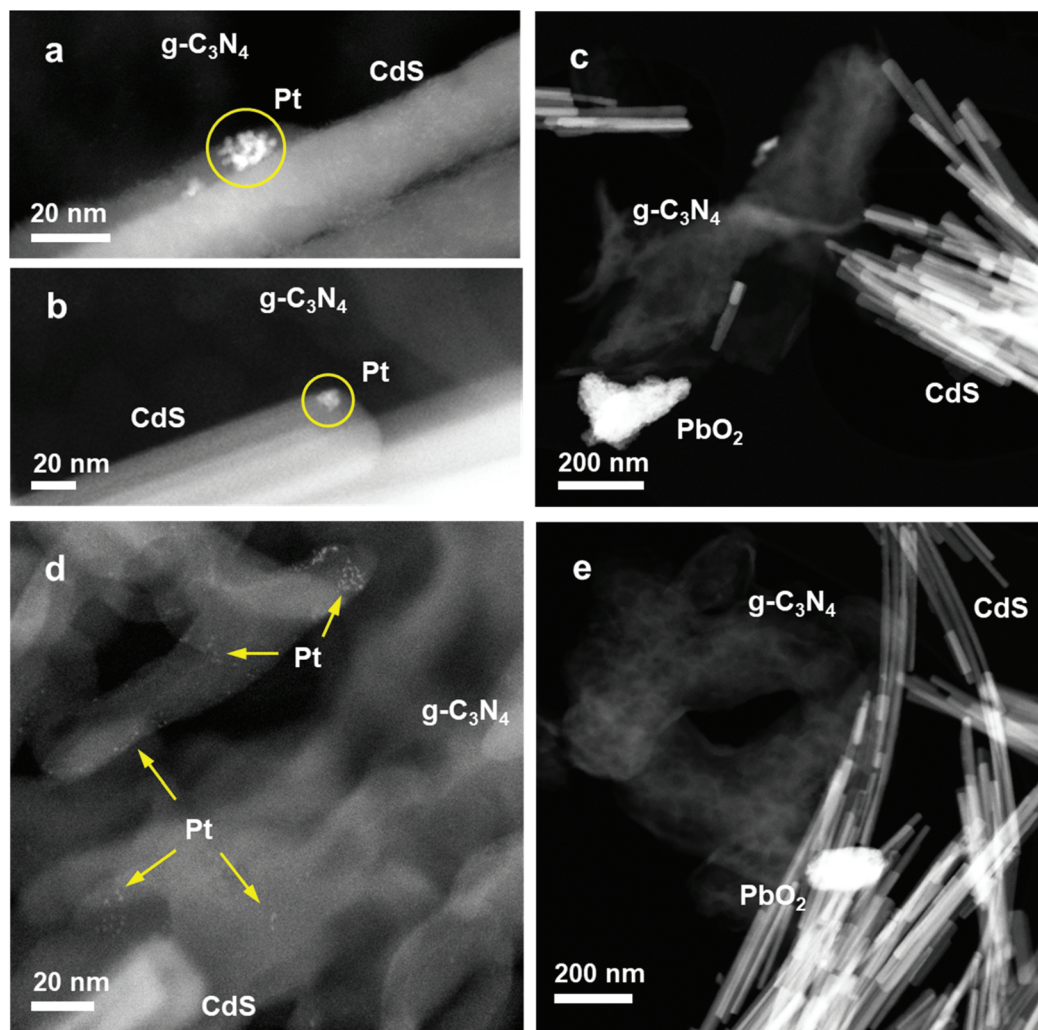


Fig. 4 Charge distribution on CS/CN-II and CS/CN-Z. HAADF images of CS/CN-II with (a and b) 1 wt% Pt and (c) 3 wt% PbO₂ loaded, and CS/CN-Z with (d) 1 wt% Pt and (e) 3 wt% PbO₂ loaded.

trons and anchoring Pt nanoparticles, which act as active sites for photoreduction reactions. Moreover, the PbO₂ clusters with (101) facets were located on the CdS nanorods (Fig. 4e and S9d†). Therefore, it can be concluded that the electrons are distributed at the CB of g-C₃N₄ and the holes are located at the VB of CdS for CS/CN-Z. Charge extinction may exist in CS/CN-Z through recombination of e⁻ at the CdS CB and h⁺ at the g-C₃N₄ VB. Although spatial charge separation is achieved by CS/CN-Z, it has a different charge separation mechanism and transfer route compared to CS/CN-II.

XPS analysis was conducted to analyze the chemical valence of Pt and Pb after the photoreactions. Since metallic Pt⁰ is regarded as the active site for photocatalytic H₂ generation,⁴⁵ its percentage in the Pt species will largely affect the photocatalytic activity. As shown in Fig. S10a and S10b,† the two main bands located at 71.1 and 74.3 eV are ascribed to the Pt 4f_{7/2} and Pt 4f_{5/2} of Pt⁰, respectively. The spectra demonstrate that most of the Pt species loaded on CS/CN-II and CS/CN-Z exist in the form of Pt⁰. The shoulder peaks at 72.4 and 75.6

eV which belong to the Pt 4f of Pt²⁺ species indicate a small portion of Pt⁴⁺ was partially reduced to Pt²⁺. In Table S2,† based on the XPS and ICP results, the actual percentage of Pt⁰ loading in CS/CN-II is 0.69 wt%, which is 23% higher than that of CS/CN-Z (0.56 wt%). This may result in a higher photocatalytic performance of CS/CN-II due to the more Pt⁰ reactive sites. Fig. S10c and S10d† show the Pb species loaded on CS/CN-II and CS/CN-Z after the photo-deposition. The two main peaks at 137.4 and 142.3 eV correspond to the Pb 4f_{7/2} and Pb 4f_{5/2} of Pb⁴⁺, respectively. It suggests that most of Pb²⁺ was oxidized to Pb⁴⁺ on both samples. Less amount of Pb²⁺ exists in the samples which is evidenced by the shoulder peaks at 138.7 and 143.5 eV. Higher percentages of Pt⁰ and Pb⁴⁺ species on CS/CN-II indicate a superior charge separation efficiency, which is consistent with the photocurrent result (Fig. 2c). More photo-excited e⁻ and h⁺ are spatially separated and available for photoredox reactions in CS/CN-II. Such properties accounts for the excellent photocatalytic H₂ generation activity of CS/CN-II.



3.4. SRM study of charge distribution at single-turnover resolution

To study the charge distribution and photocatalytic heterogeneity at single-turnover resolution, super-resolution fluorescence microscopy was used to visualize the photoreactions (Fig. 5a). Fluorogenic reactions were designed to obtain the mappings of photo-excited e^- and h^+ , in which the non-fluorescent reactants are catalyzed to highly fluorescent products. The photoreduction of non-fluorescent resazurin to fluorescent resorufin can be applied to localize the electrons.²⁵ Correspondingly, photogenerated h^+ can be detected by an indirect mechanism, in which non-fluorescent Amplex Red can be oxidized to resorufin by $\cdot OH$ produced from the reaction between holes and adsorbed H_2O .²⁶ Hence, each fluorescence burst observed in the SRM image originates from a single photocatalytic turnover which leads to the formation of a fluorescent resorufin molecule at a reactive site (Fig. 5b). As shown in the intensity trajectory in Fig. 5c, τ_{off} represents the waiting time before the formation of each resorufin molecule, while τ_{on} signifies the residence time of resorufin on the catalyst surface before dissociation. Thus, the one-step ON/OFF change of fluorescence signals with low background in Fig. 5c suggests the stochastic single-turnover events. By recording the locations of millions of catalytic events, all the single-molecule frames containing fluorescent bursts were integrated and fitted by point spread function. Consequently, SRM images can be reconstructed with a spatial resolution of 16 nm (Fig. S11†). Fig. S12† shows the brightfield images, SRM images and density maps of pure CdS nanorods with resazurin. The SRM images provide the information of super-localization of photoactive species and reactive sites with nanometer-scale precision. The density maps processed from SRM images clearly demonstrate the density of these species, catalytic reactivity and heterogeneity at different locations. Under laser excitation, the fluorescent bursts of resorufin are mainly located at the end segments of CdS nanorods and detected less at the central parts, as shown at the positions 1, 2, 3, and 4 in Fig. S12.† This demonstrates that the separated electrons

are predominantly distributed at the ends of CdS nanorods, indicating a higher reactivity at this region. Such phenomena are likely due to the low-coordination environment and disordered structure at the end segments, which can be observed from the positions 1, 2, and 3 in Fig. S2b.† The regions such as defects, terrace, kinks, steps and corner sites are believed to have a higher catalytic performance owing to the high coordinative unsaturation.^{46–48} Fig. S13† shows the morphologies of CdS before and after illumination with Amplex Red. The CdS nanorods suffer from severe photocorrosion when h^+ are localized and react with Amplex Red. Hence, the morphology of CdS totally changed after irradiation (Fig. S13b†) and no SRM images can be obtained. For pure $g-C_3N_4$, it was found that the fluorescent bursts of resazurin photoreduction mostly take place at the edges of nanosheets, suggesting the photogenerated e^- migrated and concentrated at this position (Fig. S14†). Moreover, the fluorescent spots generated from the photo-oxidation of Amplex Red mainly emerged at the central parts of $g-C_3N_4$ nanosheets (Fig. S15†). This indicates that the h^+ are mostly distributed at the basal planes of nanosheets. Therefore, spatial charge separation can be also achieved and observed on pure $g-C_3N_4$.

The SRM images were also obtained for CdS/ $g-C_3N_4$ nanocomposites to study the charge flow and distribution. In Fig. 6a–c, the fluorescent bursts on CS/CN-II from resazurin photoreduction mainly occurred at CdS nanorods, suggesting the diffusion direction and preferred location of photo-induced electrons. More electrons are found at the end segments of CdS than the central parts, which is also observed for pure CdS (Fig. S12†). Correspondingly, the h^+ are found to be distributed at the edges of $g-C_3N_4$ as shown in Fig. 6d–f. It has been reported that the charge (*i.e.*, e^- and h^+) density at the edge of $g-C_3N_4$ /N-doped graphene is higher than that of basal planes owing to the higher local curvature of edges than that of basal planes.⁴⁹ A similar phenomenon was also observed for other 2D layered materials using the SRM method.⁵⁰ With electrons flowing to CdS in the composite, the holes would preferentially locate at the edge position of $g-C_3N_4$ in CS/CN-II. The reason why holes are located at the basal planes of pure

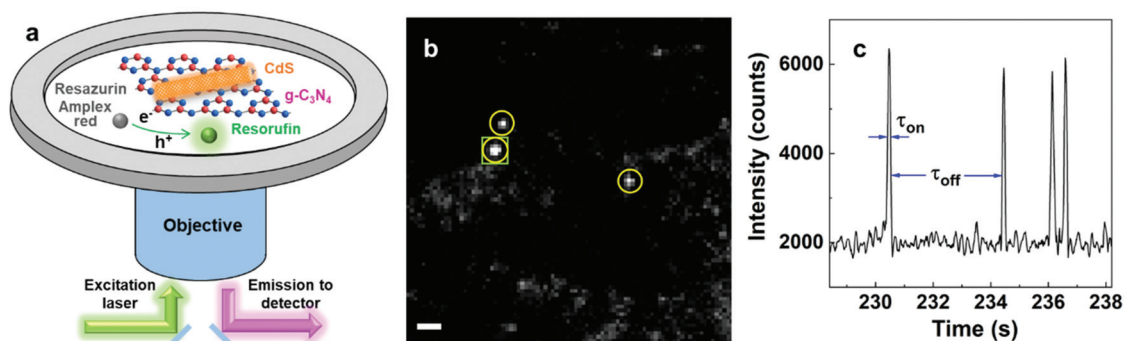


Fig. 5 Super-resolution imaging of charge locations using fluorescent probes. (a) Schematic of total internal reflection fluorescence microscopy. (b) A single-frame image of fluorescent resorufin molecules (in yellow circles) on CdS nanorods. Scale bar: 1 μm . (c) Typical fluorescence intensity trajectory for an area of 500 nm \times 500 nm in the marked green square in (b) (bin time: 30 ms).



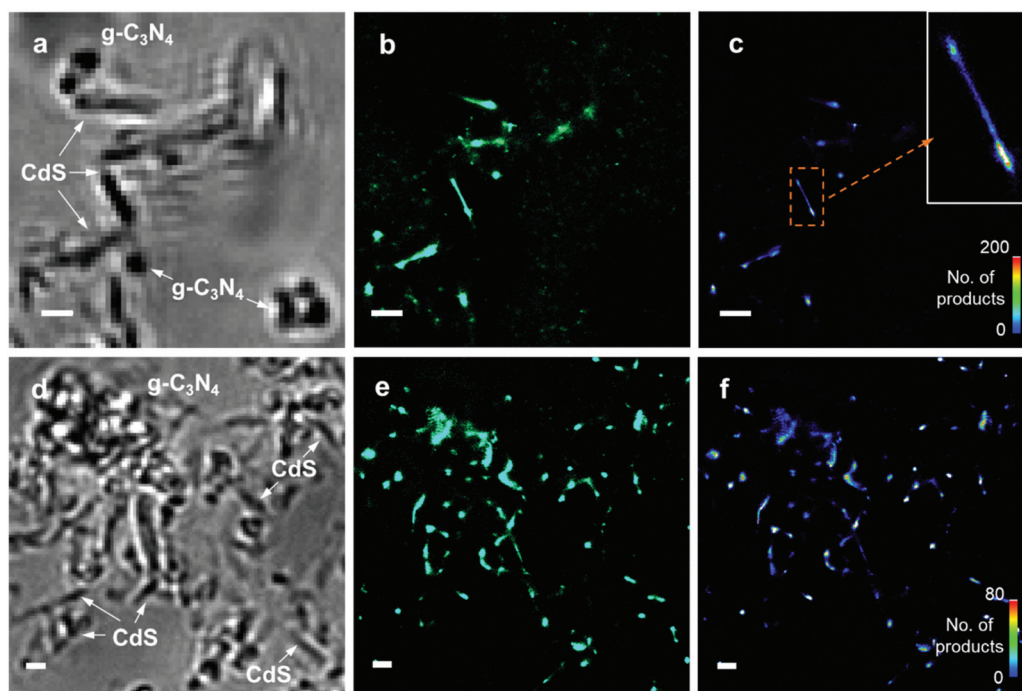


Fig. 6 Single-molecule SRM of CS/CN-II at single-turnover resolution. Conventional brightfield images, SRM images and density maps of CS/CN-II with (a–c) resazurin (inset: the close-up of the marked area (orange dashed line) in (c), bin size: 10 nm × 10 nm) and (d–f) Amplex Red (bin size: 25 nm × 25 nm). Scale bar: 1 μ m.

$g\text{-C}_3\text{N}_4$ is possibly because electrons are preferable to occupy the high curvature sites at edges without the heterojunction structure.⁵¹ Future studies are needed to further clarify the detailed mechanism of charge distribution and catalytic events on pure 2D nanosheets. Fig. 7 shows the conventional images, SRM images and density maps of CS/CN-Z with resazurin. Under laser excitation, the fluorescent spots which represent the locations of photo-induced e^- are distributed at the edges of $g\text{-C}_3\text{N}_4$ nanosheets. This result suggests that the electrons are accumulated at $g\text{-C}_3\text{N}_4$ in CS/CN-Z. However, the photo-oxidation of Amplex Red on CS/CN-Z leads to the photocorrosion of CdS in the composite, as indicated in the conventional images of CS/CN-Z before and after laser irradiation

(Fig. S16[†]). This indicates that the h^+ are located at CdS, which proves the Z-scheme structure of CS/CN-Z. Therefore, the super-resolution mapping results clearly reveal the surface reactive sites at the nanoscale and elucidate the two charge separation mechanisms of CS/CN-II and CS/CN-Z.

Based on the charge tracking results of photo-deposition and super-resolution fluorescence imaging, two charge transfer routes were confirmed for CS/CN-II and CS/CN-Z. In Scheme 1a, the band alignment of CdS and $g\text{-C}_3\text{N}_4$ in CS/CN-II makes the charges migrate in the opposite direction, leading to an efficient spatial charge separation. Charge extinction can be significantly suppressed through this configuration which contributes to the excellent photocatalytic performance and

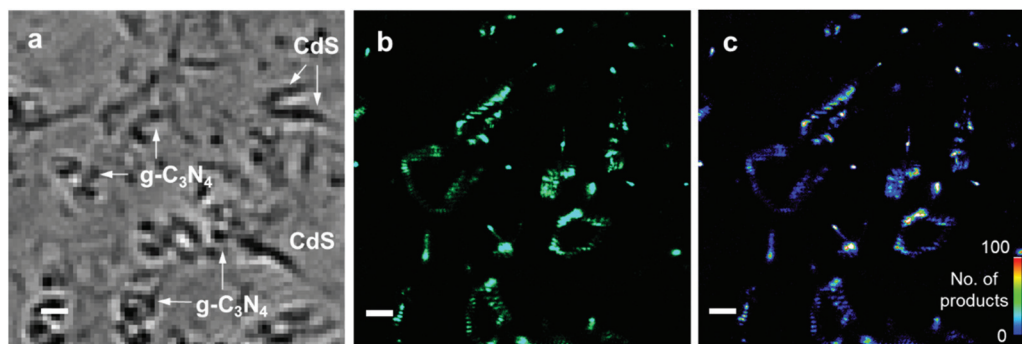
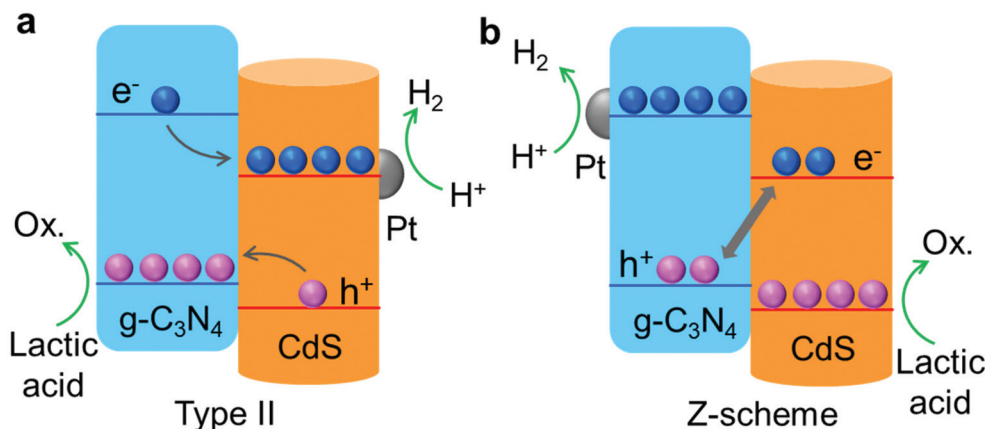


Fig. 7 Single-molecule SRM of CS/CN-Z at single-turnover resolution. (a) Conventional brightfield images, (b) SRM images and (c) density maps (bin size: 25 nm × 25 nm) of CS/CN-Z with resazurin. Scale bar: 1 μ m.





Scheme 1 Schematic of charge transfer and photocatalytic reactions on (a) CS/CN-II and (b) CS/CN-Z.

stability of CS/CN-II. In contrast, CS/CN-Z exhibits a different charge transfer route with charge recombination present between the CdS CB and the g-C₃N₄ VB (Scheme 1b). Hence, the charge separation efficiency is expected to be lower than that of CS/CN-II. Besides, as shown in Fig. S7c,† CS/CN-II exhibits a VB potential of 1.71 eV, which is closer to the g-C₃N₄ VB (1.64 eV). This implies that the h⁺ in the composite rapidly transfer to the VB of g-C₃N₄. In contrast, the VB potential of CS/CN-Z (1.77 eV) is found to be closer to that of CdS (1.83 eV), indicating the accumulation of h⁺ on the CdS VB (Fig. S7d†). Despite the stronger oxidative VB potential of CS/CN-Z compared to CS/CN-II, the photocatalytic performance is inferior to CS/CN-II (Fig. 3a). This is because the poor charge separation efficiency and serious photocorrosion in CS/CN-Z act as the main factors limiting the photocatalytic efficiency and stability. Although CS/CN-II and CS/CN-Z have the same content of CdS and g-C₃N₄, two distinct heterostructures were formed due to the different synthetic routes. As revealed by the FTIR and XPS results (Fig. 2a and S4†), an intense interaction between CdS and g-C₃N₄ was developed in CS/CN-II because of thermal calcination at 300 °C. The closely contacted interface is a key prerequisite and can promote the kinetic flow of charges between the two components. In the case of CS/CN-Z, which was prepared by the self-assembly method at room temperature, the charges do not prefer to transfer kinetically and tend to stay at their own electronic band due to the weak interaction. Furthermore, it was found that CdS in CS/CN-II at the interface position is more crystalline than that in CS/CN-Z. As shown in the HRTEM images in Fig. S17a,† the highly crystalline CdS at the interface was observed for CS/CN-II due to calcination. However, some disordered areas which contain bulk defects were identified for CS/CN-Z (Fig. S17b†). The bulk defects could act as deep trapping sites for photogenerated e⁻ and h⁺ and retard their mobility, leading to the formation of charge recombination centers at these sites. In contrast, a well-ordered crystalline structure of CdS at the interface of CS/CN-II facilitates rapid charge transportation to the catalyst surface for redox reactions. The notably enhanced crystallinity of CS/CN-II can be also verified by the XRD results (Fig. S1†).

Therefore, the more intense interaction and the enhanced crystalline structure in CS/CN-II due to thermal calcination account for the efficient interfacial charge transfer and separation. Hence, based on the above discussion, our work demonstrates a direct imaging method to visualize and map the photoactive species in composites at the nanoscale. It provides an in-depth understanding of charge distribution and energy flows in two distinct heterostructures and correlates these properties with the photocatalytic activity. We expect that this study would benefit the rational design of type II and Z-scheme composites, intended for efficient photocatalytic applications.

4. Conclusion

In conclusion, we applied the techniques of HRTEM, HAADF STEM and SRM to investigate the charge separation mechanisms of type II and Z-scheme CdS/g-C₃N₄ composites. Different charge transfer routes were observed for the two heterostructures. In particular, the SRM method was used for the first time to map the photoactive sites on type II and Z-scheme heterojunctions at the nanoscale. It was found that the photo-excited electrons are predominantly located at the two end segments of CdS nanorods while the photoactive species are accumulated at the edges of g-C₃N₄. As revealed by the photoactivity measurement, type II heterostructure of CdS/g-C₃N₄ exhibits a superior photocatalytic performance to the Z-scheme one, due to the higher charge separation efficiency and less photocorrosion. Therefore, this work is essential for the following reasons. First, through combining the microscopic study with the bulk activity measurement, our findings provide a deep insight into the understanding of “structure–activity” correlations in nanocomposites. Second, it enables us to potentially control the energy and charge flows and engineer the reactive sites in these materials. Third, such investigation would benefit the electronic band modulation and interface engineering of heterogeneous catalysts, leading to an enhancement of photocatalytic activity under visible light.



Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We acknowledge the financial support from the Ministry of Education, Singapore, under its Academic Research Fund Tier 1 (No. RG10/20), and the Singapore Agency for Science, Technology and Research (A*STAR) AME YIRG grant (No. A2084c0065). We also would like to thank Haojing Wang for her help in photoelectrochemical measurements.

References

- 1 A. Fujishima and K. Honda, *Nature*, 1972, **238**, 37–38.
- 2 Y. Liu, X. Zeng, X. Hu, J. Hu and X. Zhang, *J. Chem. Technol. Biotechnol.*, 2019, **94**, 22–37.
- 3 C. Tan, X. Cao, X.-J. Wu, Q. He, J. Yang, X. Zhang, J. Chen, W. Zhao, S. Han and G.-H. Nam, *Chem. Rev.*, 2017, **117**, 6225–6331.
- 4 Y. Zhu, Y. Feng, S. Chen, M. Ding and J. Yao, *J. Mater. Chem. A*, 2020, **8**, 25626–25648.
- 5 X. Zhang, X. Yuan, L. Jiang, J. Zhang, H. Yu, H. Wang and G. Zeng, *Chem. Eng. J.*, 2020, **390**, 124475.
- 6 Y. Lu, H. Wang, P. Yu, Y. Yuan, R. Shahbazian-Yassar, Y. Sheng, S. Wu, W. Tu, G. Liu and M. Kraft, *Nano Energy*, 2020, **77**, 105158.
- 7 W. Tu, Y. Xu, J. Wang, B. Zhang, T. Zhou, S. Yin, S. Wu, C. Li, Y. Huang, Y. Zhou, Z. Zou, M. Kraft and R. Xu, *ACS Sustainable Chem. Eng.*, 2017, **5**, 7260–7268.
- 8 J. Liang, X. Yang, Y. Wang, P. He, H. Fu, Y. Zhao, Q. Zou and X. An, *J. Mater. Chem. A*, 2021, **9**, 12898–12922.
- 9 Y. Ni, R. Wang, W. Zhang, S. Shi, W. Zhu, M. Liu, C. Yang, X. Xie and J. Wang, *Chem. Eng. J.*, 2021, **404**, 126528.
- 10 N. Feng, H. Lin, F. Deng and J. Ye, *J. Phys. Chem. C*, 2021, **125**, 12012–12018.
- 11 X. Zhao, Y. Fan, W. Zhang, X. Zhang, D. Han, L. Niu and A. Ivaska, *ACS Catal.*, 2020, **10**, 6367–6376.
- 12 Y. Luo, J. Qin, G. Yang, S. Luo, Z. Zhao, M. Chen and J. Ma, *Chem. Eng. J.*, 2021, **410**, 128394.
- 13 N. Güy, *Appl. Surf. Sci.*, 2020, **522**, 146442.
- 14 Y. Chen, F. Wang, Y. Cao, F. Zhang, Y. Zou, Z. Huang, L. Ye and Y. Zhou, *ACS Appl. Energy Mater.*, 2020, **3**, 4610–4618.
- 15 Z. Shi, Y. Zhang, X. Shen, G. Duoerkun, B. Zhu, L. Zhang, M. Li and Z. Chen, *Chem. Eng. J.*, 2020, **386**, 124010.
- 16 Y. Wang, H. Huang, Z. Zhang, C. Wang, Y. Yang, Q. Li and D. Xu, *Appl. Catal., B*, 2021, **282**, 119570.
- 17 M. Zhang, C. Lai, B. Li, F. Xu, D. Huang, S. Liu, L. Qin, X. Liu, H. Yi and Y. Fu, *Chem. Eng. J.*, 2021, **422**, 130120.
- 18 X. Zhou, Y. Fang, X. Cai, S. Zhang, S. Yang, H. Wang, X. Zhong and Y. Fang, *ACS Appl. Mater. Interfaces*, 2020, **12**, 20579–20588.
- 19 G. Li, B. Wang, J. Zhang, R. Wang and H. Liu, *Appl. Surf. Sci.*, 2019, **478**, 1056–1064.
- 20 D. P. Kumar, S. Hong, D. A. Reddy and T. K. Kim, *J. Mater. Chem. A*, 2016, **4**, 18551–18558.
- 21 J. Pan, P. Wang, P. Wang, Q. Yu, J. Wang, C. Song, Y. Zheng and C. Li, *Chem. Eng. J.*, 2021, **405**, 126622.
- 22 N.-N. Vu, S. Kaliaguine and T.-O. Do, *ACS Appl. Energy Mater.*, 2020, **3**, 6422–6433.
- 23 M. J. Rust, M. Bates and X. Zhuang, *Nat. Methods*, 2006, **3**, 793–796.
- 24 A. Sharonov and R. M. Hochstrasser, *Proc. Natl. Acad. Sci. U. S. A.*, 2006, **103**, 18911–18916.
- 25 J. B. Sambur, T.-Y. Chen, E. Choudhary, G. Chen, E. J. Nissen, E. M. Thomas, N. Zou and P. Chen, *Nature*, 2016, **530**, 77–80.
- 26 J. W. Ha, T. P. A. Ruberu, R. Han, B. Dong, J. Vela and N. Fang, *J. Am. Chem. Soc.*, 2014, **136**, 1398–1408.
- 27 M. Shi, G. Li, J. Li, X. Jin, X. Tao, B. Zeng, E. A. Pidko, R. Li and C. Li, *Angew. Chem., Int. Ed.*, 2020, **59**, 6590–6595.
- 28 J. Liu, T. Zhang, Z. Wang, G. Dawson and W. Chen, *J. Mater. Chem.*, 2011, **21**, 14398–14401.
- 29 L. J. Zhang, T. F. Xie, D. J. Wang, S. Li, L. L. Wang, L. P. Chen and Y. C. Lu, *Int. J. Hydrogen Energy*, 2013, **38**, 11811–11817.
- 30 Y. Li, X. Wei, H. Li, R. Wang, J. Feng, H. Yun and A. Zhou, *RSC Adv.*, 2015, **5**, 14074–14080.
- 31 M. Lu, Z. Pei, S. Weng, W. Feng, Z. Fang, Z. Zheng, M. Huang and P. Liu, *Phys. Chem. Chem. Phys.*, 2014, **16**, 21280–21288.
- 32 P. Wang, T. Wu, C. Wang, J. Hou, J. Qian and Y. Ao, *ACS Sustainable Chem. Eng.*, 2017, **5**, 7670–7677.
- 33 H. He, J. Cao, M. Guo, H. Lin, J. Zhang, Y. Chen and S. Chen, *Appl. Catal., B*, 2019, **249**, 246–256.
- 34 L. Ge, F. Zuo, J. Liu, Q. Ma, C. Wang, D. Sun, L. Bartels and P. Feng, *J. Phys. Chem. C*, 2012, **116**, 13708–13714.
- 35 B. Chong, L. Chen, D. Han, L. Wang, L. Feng, Q. Li, C. Li and W. Wang, *Chin. J. Catal.*, 2019, **40**, 959–968.
- 36 Z. Wang, Z. Wang, X. Zhu, C. Ai, Y. Zeng, W. Shi, X. Zhang, H. Zhang, H. Si and J. Li, *Small*, 2021, 2102699.
- 37 X. Liu, Y. Liu, W. Zhang, Q. Zhong and X. Ma, *Mater. Sci. Semicond. Process.*, 2020, **105**, 104734.
- 38 Y. Xu, Y. Li, P. Wang, X. Wang and H. Yu, *Appl. Surf. Sci.*, 2018, **430**, 176–183.
- 39 Y. Xu and M. A. Schoonen, *Am. Mineral.*, 2000, **85**, 543–556.
- 40 S. Wu, W. Wang, W. Tu, S. Yin, Y. Sheng, M. Y. Manuputty, M. Kraft and R. Xu, *ACS Sustainable Chem. Eng.*, 2018, **6**, 14470–14479.
- 41 W. Li, C. Feng, S. Dai, J. Yue, F. Hua and H. Hou, *Appl. Catal., B*, 2015, **168**, 465–471.
- 42 L. Qian, Y. Hou, Z. Yu, M. Li, F. Li, L. Sun, W. Luo and G. Pan, *Mol. Catal.*, 2018, **458**, 43–51.
- 43 S. Yin, J. Han, Y. Zou, T. Zhou and R. Xu, *Nanoscale*, 2016, **8**, 14438–14447.
- 44 X. Ning and G. Lu, *Nanoscale*, 2020, **12**, 1213–1223.
- 45 S. Wu, M. Y. Manuputty, Y. Sheng, H. Wang, Y. Yan, M. Kraft and R. Xu, *Small Methods*, 2021, **5**, 2000928.
- 46 M. J. K. Ow, J. J. Ng, J. X. Yong, B. Y. L. Quek, E. K. Yeow and Z. Zhang, *ACS Appl. Nano Mater.*, 2020, **3**, 3163–3167.



- 47 L. Liu and A. Corma, *Chem. Rev.*, 2018, **118**, 4981–5079.
- 48 X. Zhou, N. M. Andoy, G. Liu, E. Choudhary, K.-S. Han, H. Shen and P. Chen, *Nat. Nanotechnol.*, 2012, **7**, 237–241.
- 49 C. Xu, S. Wu, G. Xiong, X. Guo, H. Yang, J. Yan, K. Cen, Z. Bo and K. K. Ostrikov, *Appl. Catal., B*, 2021, **280**, 119461.
- 50 T.-X. Huang, B. Dong, S. L. Filbrun, A. A. Okmi, X. Cheng, M. Yang, N. Mansour, S. Lei and N. Fang, *Sci. Adv.*, 2021, **7**, eabj4452.
- 51 J. Yang, Y. Liu, D. Zhang, X. Wang, R. Li and Y. Li, *Nano Res.*, 2015, **8**, 3054–3065.

