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Citation: Isimjan, Tayirjan Taylor, Rasul, Shahid, Nasser Aloufi, Maher, Khan, Mohd Adnan, Alhawaish, Ibrahim Khalid and Ahmed, Toseef (2019) Rational design of Pd-TiO₂/g-C₃N₄ heterojunction with enhanced photocatalytic activity through interfacial charge transfer. *Clean Energy*, 3 (1). pp. 59-68. ISSN 2515-4230

Published by: Oxford University Press

URL: <https://doi.org/10.1093/ce/zky021> <<https://doi.org/10.1093/ce/zky021>>

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RESEARCH ARTICLE

Rational design of Pd-TiO₂/g-C₃N₄ heterojunction with enhanced photocatalytic activity through interfacial charge transfer

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Abstract

A hybrid heterojunction-based photocatalyst is synthesized by an electrostatic self-assembly strategy including surface modification and controlled metal deposition. The interfacial contact was made by mixing negatively charged anatase TiO₂ nanoparticles with positively charged g-C₃N₄. Visible-light deposition of Pd nanoparticles largely on TiO₂ was made possible due to the charge transfer from C₃N₄ (excited by visible light) to the conduction band of TiO₂, reducing Pd ions on contact with its surface. In order to further test the efficiency of this cascade of electron transfer across the conduction bands of the two semiconductors, photocatalytic H₂ production from water was studied. Upon optimizing the ratio of the two semiconductors, increased H₂ production rates were observed and attributed to enhanced charge separation. Catalysts were studied by a variety of techniques in order to probe into their properties and link them to activity. The reaction rate, under visible-light excitation, of the best sample showed an 8-fold enhancement when compared to that of Pd-C₃N₄ in identical conditions and the highest apparent quantum yield of 31% was achieved by a 0.1%Pd/20%TiO₂/C₃N₄ sample in a 420- to 443-nm range.

Keywords: surface charge; heterojunction; photocatalyst; electrostatic self-assembly; hydrogen generation

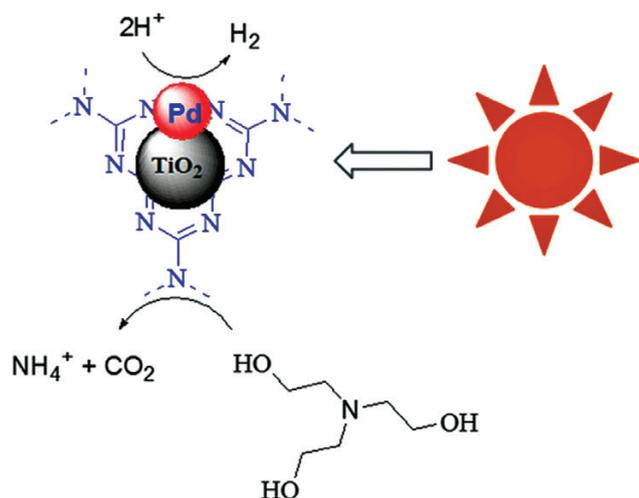
Introduction

While TiO₂ is one of the most promising photocatalysts, its application is hindered by its wide band gap energy, corresponding to a negligible fraction of the total solar energy on the Earth. To circumvent this, a good deal of effort has been devoted to developing visible-light-responsive TiO₂-based photo catalysts. These types of catalysts can be largely divided into two categories: either TiO₂ doped by

other elements [1–7] or TiO₂ coupled with smaller band gap semiconductors such as C₃N₄ [8–12], CdS [13, 14], MoS₂ [15, 16] and In₂S₃ [17] as photo-sensitizers. For the second approach, identifying a stable narrow bandgap semiconductor with a suitable band alignment with TiO₂ is a challenge. Among these semiconductors, graphitic carbon nitride (g-C₃N₄) is attractive because its conduction band edge is above that of TiO₂ and it possesses excellent chemical stability. Because g-C₃N₄, itself suffers from rapid charge

Received: 8 August, 2018; Accepted: 29 January, 2019

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carrier recombination rates, resulting in low photocatalytic activity [18], the quality of the interface with TiO_2 is crucial for the photocatalytic activity of the nanocomposites. For the coupling to be successful, the semiconductors need to fulfil at least the following two requirements: (i) an appropriate band structure that creates a potential offset between the two semiconductors, generating a band bending at the interface, which impels photo-generated electrons and holes to diffuse in opposite directions [19]; and (ii) the crystal structure and orientation of the two semiconductors need to match (at least in part) to minimize point and crystallographic defects [18]. To that end, it has been reported, based on detailed crystallographic studies, that a stable interface can be formed between the (22-40) plane of $\text{g-C}_3\text{N}_4$ and the (110) plane of $\text{TiO}_2(\text{B})$ (monoclinic, space group C2/m), because both planes have the same d-spacing (0.35 nm) and similar crystal orientation [20].

Overall, the two main strategies used to synthesize $\text{g-C}_3\text{N}_4$ - TiO_2 nanocomposites are direct mixing or simultaneous synthesis of $\text{C}_3\text{N}_4/\text{TiO}_2$ nanocomposites. For instance, Peng et al. [21] prepared $\text{g-C}_3\text{N}_4$ - Pt-TiO_2 (P25) composites by chemical adsorption and reported (at a mass ratio of 70:30 of C_3N_4 to TiO_2) a mild increase in the H_2 -generation rate under visible light with stable photocurrent, 1.5 times higher than that of pure $\text{g-C}_3\text{N}_4$. Liu et al. reported an *in situ* growth C_3N_4 -sensitized TiO_2 nanotube array-based photoanode that gave around a 7-fold improvement in the photocurrent under light irradiation (300–900 nm) when compared to TiO_2 NTs [22]. However, these methods result in self-aggregation, leading to phase separation between TiO_2 and C_3N_4 , and they consequently minimize the direct crystal contact between the two semiconductors to form a suitable interface for efficient charge transfer.

As an alternative approach, electrostatic self-assembly is used to construct C_3N_4 -based heterojunction structures in which two semiconductors are attached through opposite charges. The nanocomposites synthesized through electrostatic interaction are constructed either by using intrinsic surface charge of the materials [11, 23–27] or by surface functionalization with opposite charges [28–31].

For example, Li et al. [32] fabricated 1D Ag@AgVO_3 nanowire/graphene/protonated $\text{g-C}_3\text{N}_4$ nanosheet heterojunctions via a simple electrostatic self-assembly process followed by photochemical reduction. The resulting catalyst showed much higher photo-activity on degrading methylene blue compared to that of $\text{g-C}_3\text{N}_4$. Similarly, Fan et al. [27] fabricated $\text{g-C}_3\text{N}_4/\text{Zn}_2\text{GeO}_4$ photocatalysts by utilizing opposite electrostatic interaction between two nanoparticles. The heterojunction system showed higher visible-light photocatalytic activity towards the degradation of methylene blue than those of pure $\text{g-C}_3\text{N}_4$ and pure Zn_2GeO_4 . In addition, Wu et al. reported an electrostatic self-assembly strategy to fabricate a 2D/2D hybrid photocatalyst ($\text{g-C}_3\text{N}_4/\text{rGO}$) by incorporating reduced graphene oxide (rGO) and protonated $\text{g-C}_3\text{N}_4$ in which they used the intrinsic negative charge of rGO and positive surface charge of PCN [33]. Also, Li et al. reported the synthesis of an oxygen reduction (ORR) catalyst prepared by mixing phosphorus-doped graphitic carbon nitride nanosheets with NH_2 -functionalized carbon black and they achieved similar performance with a standard ORR catalyst (Pt/Vulcan XC-72) [34].

These results proved that electrostatic interaction-based synthesis is a most efficient way to make nanocomposite because the high zeta potential not only separates similar semiconductor nanoparticles from each other but also increases the interaction between different semiconductors through opposite-charge repulsion to form a well-dispersed nanocomposite. It not possible to formulate $\text{TiO}_2/\text{C}_3\text{N}_4$ without surface modification. Nevertheless, there are no reports on electrostatic self-assembly of TiO_2 and $\text{g-C}_3\text{N}_4$.

In this work, we have designed, synthesized and tested a $\text{Pd-TiO}_2/\text{g-C}_3\text{N}_4$ hybrid system through surface-charge interaction wherein both TiO_2 and C_3N_4 exhibited high Z-potentials, therefore preventing self-aggregation while generating enough driving force to form a uniformly mixed heterojunction and then Pd was deposited preferentially on the TiO_2 through visible-light deposition. Finally, a uniformly dispersed $\text{Pd-TiO}_2/\text{C}_3\text{N}_4$ was synthesized. The high level of uniformity was confirmed by EDX mapping. Moreover, the photocatalytic H_2 -generation rate of the best-performing sample 0.1%Pd-20% $\text{TiO}_2/\text{g-C}_3\text{N}_4$ was found to be 14- and 8-fold higher than that of the prepared $\text{g-C}_3\text{N}_4$ and 0.1%Pd/ C_3N_4 , respectively. In the end, we proposed the related charge-transfer mechanism. We believe this unique two-step approach, including the electrostatic interaction and the directed metal deposition, can be used as a universal method to prepare various heterojunction systems.

1 Experiment

All reagents were of analytical grade, from Sigma-Aldrich and used without further purification. Detailed synthetic procedures are described below.

1.1 Synthesis of Pd-TiO₂/g-C₃N₄

1.1.1 Sulphonic acid-functionalized TiO₂ (TiO₂-SO₃Na)

A typical synthesis is described as follows. (i) To 3 g of TiO₂ (anatase) in 30 mL dry ethanol and 1 mL of H₃PO₄, 3 mL of (3-mercaptopropyl) trimethoxysilane was added and the reaction mixture was refluxed for 24 h. After this period, the mixture was filtered, washed with acetone and dried in air to obtain TiO₂-SH. (ii) The resulting TiO₂-SH was dispersed in 10 wt.% H₂O₂ in methanol (20 mL) and stirred for 24 h at room temperature. The prepared sample was then treated with 1 N H₂SO₄ at ambient temperature for complete protonation, filtered, re-dispersed in water, adjusted pH = 10. The resulting dispersion was filtered and washed with H₂O and acetone to obtain TiO₂-SO₃Na.

1.1.2 Positively charged g-C₃N₄

The g-C₃N₄ was prepared from urea at 675°C for 4 h with a heating rate of 0.5 °C/min. The yield was around 6%, consistently with other reported yields [35–37]; 1 g of as-prepared g-C₃N₄ was added to 200 mL of HCl (1 M) solution and ultra-sonicated for 1 h, stirred for 4 h, centrifuged, washed with DI water and then dried at 80°C for 12 h.

1.1.3 TiO₂-SO₃(-)/C₃N₄(+)

Positively charged g-C₃N₄(+) was dispersed in 70 mL of DI water and then added dropwise to certain amounts of TiO₂-SO₃Na dispersed in 30 mL of DI water. The resulting mixture was stirred at 25°C overnight, filtered and dried at 80°C for 3 h followed by calcination at 350°C for 4 h.

1.1.4 0.1%Pd-TiO₂/g-C₃N₄

To a 100-mL suspension of TiO₂/g-C₃N₄ was added 0.3 mL Pd (1 mg/mL). The resulting mixture was stirred under visible light (420–650 nm, 43 mW/cm²) for 4 h. The suspension was filtered, washed with ethanol and dried at 80°C in the air to give the final product.

1.2 Characterization

UV–vis absorbance spectra of the powdered catalysts were collected over the wavelength range of 250–700 nm using a Thermo Fisher Scientific spectrophotometer equipped with praying mantis diffuse reflectance accessory. Absorbance (A) and reflectance (% R) of the samples were measured. The reflectance (% R) data were used to calculate the band gap of the samples using the Tauc plot (Kubelka–Munk function). XRD spectra were recorded using a Bruker D8 Advance X-ray diffractometer. Cu K α ($\lambda = 1.5406 \text{ \AA}$) radiation over the range of 2θ intervals between 20 and 90° with a step size of 0.010° and a step time of 0.2 s/step was used. XPS spectra of the samples were collected using a Thermo Scientific Escalab 250 XI XP spectrometer with Al K α X-ray source. The X-ray spot size was 650 × 650 μm^2 . The charge compensation was carried out using a standard flood gun. Before collecting

XPS data, samples were etched using Ar ions for 5 min at the ion energy of 1000 eV. The data were acquired using the following settings before and after etching: Pass Energy = 20 eV, Energy Step Size = 0.1 eV and Dwell Time = 100 ms. All the peaks were corrected with respect to the binding energy of adventitious C1s peak at 284.5 eV. All peaks were fitted using SMART background option and the Lorentzian/Gaussian ratio of 0.3. TEM analysis of the samples was performed using a TitanG2 80–300 CT from FEI Instruments that was equipped with a field-emission gun. The analysis was conducted by operating the microscope with a beam energy of 300 keV. TEM specimens were prepared by placing a small number of samples on holey carbon-coated copper (Cu) grids with a mesh size of 300. Several low- and high-resolution electron micrographs were acquired from various locations during the analysis. Fast-Fourier transform (FFT) analysis was applied to various regions of the High-resolution TEM (HRTEM) micrographs to investigate the different crystal structures. Scanning electron microscopy/energy-dispersive X-ray spectroscopy (SEM-EDS) analysis was done at 20 kV (Quanta 200, FEI, OR). Zeta-potential measurement was performed on a zeta potential and submicron particle size analyser equipped with delsa nano auto-titrator (Model DelsaNano C, Beckman Coulter). FT-IR spectra were obtained using the Thermo Nicolet 6700 FT-IR spectrometer. Photoluminescence (PL) spectra were measured at room temperature on an A10094 fluorescence spectrometer (Hamamatsu Photonics K. K., Japan) with an excitation wavelength of 320 nm and the suspension concentration was 0.06 mg/mL in deionized water.

1.3 Photocatalytic tests

Photocatalytic reactions were evaluated in a 137-mL-volume Pyrex glass reactor using 25 mg of catalyst and 30 mL of 10 vol.% triethanol amine (TEA) aqueous solution. The slurry was purged with N₂ gas to remove any O₂ by constant stirring before the reaction. A xenon lamp (Asahi spectra MAX-303) with a total flux of 42.5 mW/cm (of which the UV contribution is ~3 mW/cm) and visible up to 650 nm at a distance of 2 cm was used. Product analyses were performed by gas chromatography equipped with a thermal-conductivity detector connected to a Porapak Q packed column (2 m) at 45°C and N₂ was used as a carrier gas. The apparent quantum yields (AQY) were measured under a light intensity of 43 mW/cm² (420–650 nm). The illumination area was 10 cm².

1.4 Photocurrent measurements

The analysis was conducted in conventional two-electrode cells on a Biologic electrochemical workstation with the Pt as a counter electrode. The studied samples were ground roughly and 20 mg was dispersed in 1.0 mL of DMF. The mixture was ultrasonically treated to obtain a slurry, which

was spin-coated onto pre-treated FTO glass and the working electrode was further dried at 150°C for 2 h. 10% TEA in 1 M NaOH was used as the electrolyte for the photocurrent response measurements.

2 Results and discussion

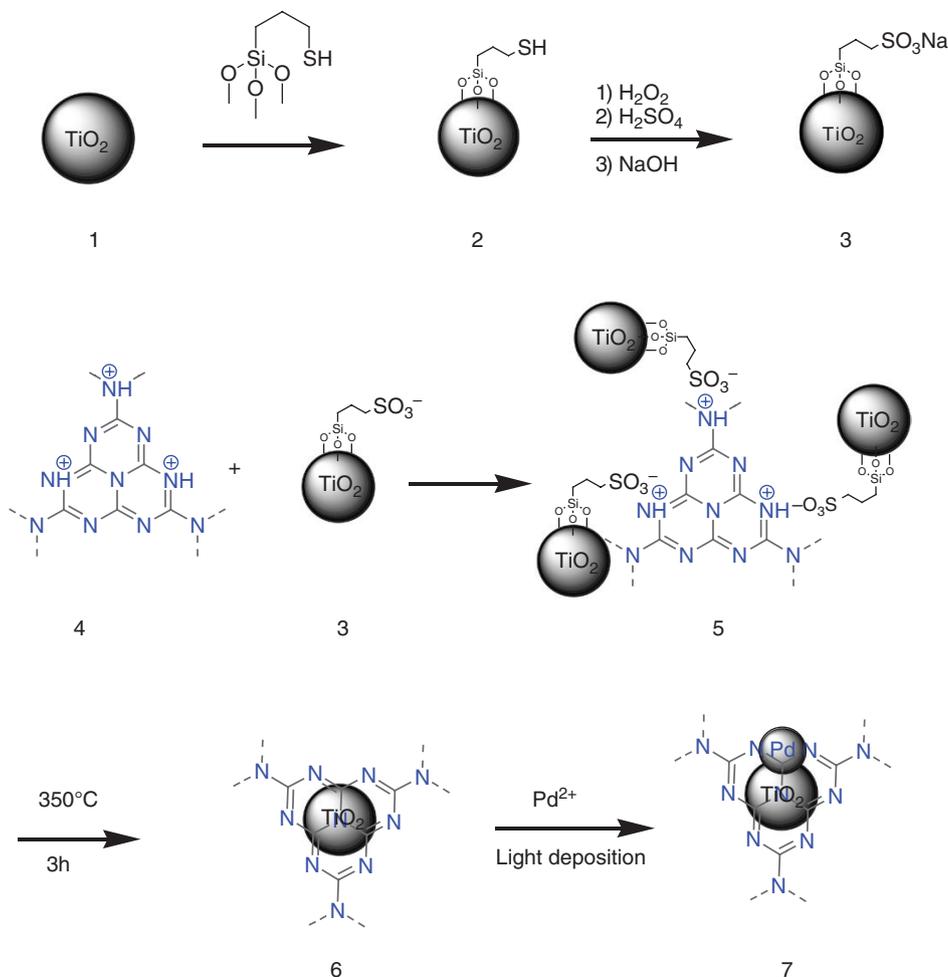
The surface-charge method used to prepare the $\text{TiO}_2/\text{g-C}_3\text{N}_4$ hybrid system is illustrated in Scheme 1. The surface of TiO_2 (1) was modified with 3-mercaptopropyl trimethoxysilane (MPTMS) to thiol-functionalized TiO_2 (2) followed by thiol oxidation to sulphonic acid (2), which was deprotonated by NaOH to form sulphonic ions functionalized TiO_2 (3). A negatively charged suspension of (3) was then uniformly mixed with a positively charged suspension of C_3N_4 (4) to produce the hybrid composite of $\text{TiO}_2\text{-SO}_3(-)/\text{g-C}_3\text{N}_4(+)$ (5), which was annealed at 350°C for 3 h in order to remove the surface ligand and form the targeted hybrid system $\text{TiO}_2/\text{C}_3\text{N}_4$ (6). Finally, Pd cations were light-deposited on (6) to give Pd- $\text{TiO}_2/\text{g-C}_3\text{N}_4$ (7).

To monitor the charge of the materials, we performed zeta-potential measurements of the protonated $\text{C}_3\text{N}_4(+)$,

the negatively charged $\text{TiO}_2\text{-SO}_3(-)$ and the heterojunction made of $\text{TiO}_2\text{-SO}_3(-)$ (20 wt%)/ $\text{C}_3\text{N}_4(+)$. The zeta potential of $\text{C}_3\text{N}_4(+)$ was found to be equal to +19.6 mV and that of $\text{TiO}_2\text{-SO}_3(-)$ was equal to -21.3 mV. The potential difference between the two (40.9 mV) is much higher than that reported in the literature [27]. The high zeta potentials prevent self-aggregation of each component in water and generate enough driving force to form a uniformly dispersed suspension. Cancellation of charges was noticed upon the heterojunction formation where the overall Z-potential of the heterojunction system was +2.5 mV. Therefore, a uniformly dispersed $\text{TiO}_2/\text{C}_3\text{N}_4$ was achieved through this process.

To further investigate the dispersion of both components, scanning electron microscopy-energy-dispersive X-ray spectroscopy (SEM-EDS) was used to conduct the elemental mapping of Pd(0.1 wt%)- TiO_2 (20 wt%)/ C_3N_4 , Fig. 1. It is clear that the Ti atoms (of TiO_2) were well dispersed over the g- C_3N_4 semiconductor.

XRD patterns of the g- C_3N_4 mixed with different mass ratios from 5 wt% to 30 wt% of TiO_2 are presented in Fig. 2a. Most of the characteristic peaks for both components are visible in all studied ratios. The two main characteristic



Scheme 1. Schematic diagram for the synthesis of $\text{TiO}_2/\text{C}_3\text{N}_4$ via chemical modifications and electrostatic self-assembly strategy

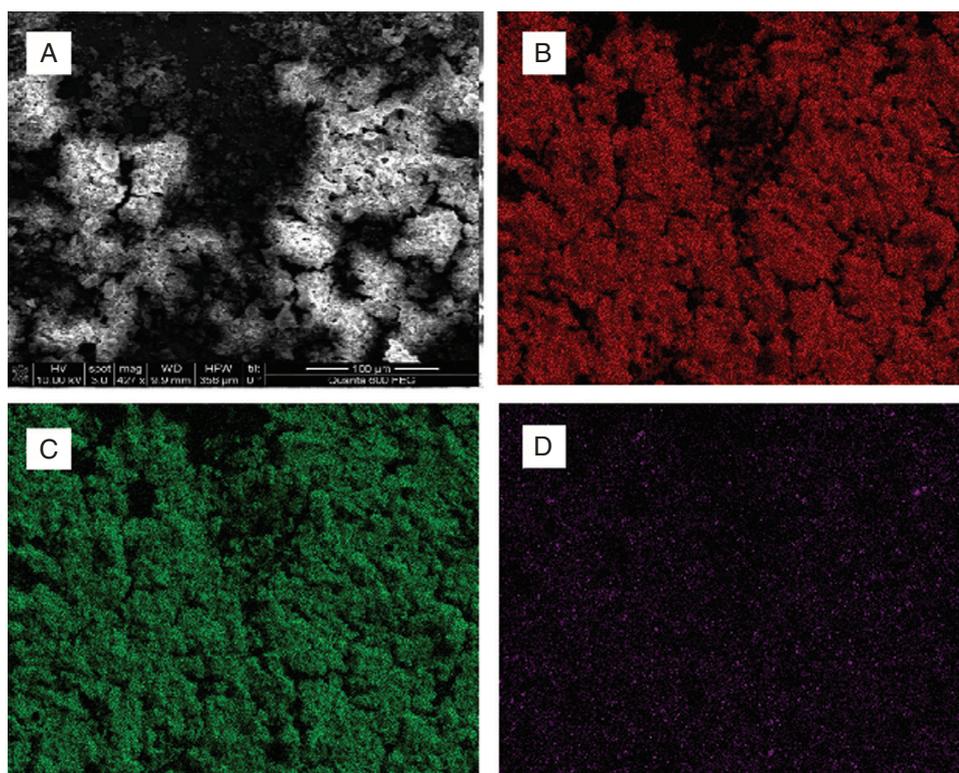


Fig. 1 SEM-EDX mapping areas. (a) Mapping area, (b) carbon, (c) nitrogen and (d) titanium. The scale in Fig. 1a is for 100 μm .

peaks of inter-layer packing (100) and inter-planar stacking (002) of $\text{g-C}_3\text{N}_4$ show at $2\theta = \sim 13$ and $\sim 27^\circ$, respectively, and are observed [38] in addition to the (101) diffraction of the anatase TiO_2 peak. Fig. 2b presents the corresponding diffuse reflectance absorption spectra of the bare 0.1%Pd- $\text{g-C}_3\text{N}_4$, TiO_2 and Pd/ TiO_2 / C_3N_4 series with various TiO_2 and $\text{g-C}_3\text{N}_4$ mass ratios. The absorption edge of TiO_2 is not visible from the spectrum. The main absorption edge of $\text{g-C}_3\text{N}_4$ occurs around 440 nm and the band gap was estimated to be around 2.8 eV from the Tuac plots (Fig. 2b, insertion).

Morphologies of 0.1% Pd-20% TiO_2 / C_3N_4 were investigated by TEM/HRTEM (Fig. 3). Fig. 3a shows that the particle sizes of Pd and TiO_2 are around 5 and 15 nm, respectively. Lattice fringes are shown in Fig. 3c and 3d in which the d-spacing of 0.2 and 0.19 nm correspond to the (200) planes of anatase TiO_2 and Pd, respectively [39]. The corresponding d-spacings from FFT are given in Fig. 3e and f. A further important observation made for Pd/ TiO_2 / C_3N_4 (Fig. 3b) was the inter-particle heterojunctions between TiO_2 and C_3N_4 as well as TiO_2 and Pd. Although the inter-particle heterojunction between C_3N_4 and Pd was also detected, the Pd density on TiO_2 is much higher than that of C_3N_4 , as shown in the STEM (Fig. 3i) and the detailed calculation is shown in Fig. S1 in the online Supplementary Data. Moreover, nanoscale EDXs are performed at the points where TiO_2 and C_3N_4 are separated (Fig. 3g). The Pd signal was only detected on TiO_2 (Fig. 3h). These results clearly indicate that the electron generated by visible light flows from the conduction band of C_3N_4 to that of TiO_2 , since the Pd was light-deposited onto the surface of the TiO_2 / C_3N_4 .

This is also indirect evidence of the proposed mechanism in Fig. 5d.

2.1 XPS spectra of 0.1% Pd-20% TiO_2 / $\text{g-C}_3\text{N}_4$ (Fig. 4 and Fig. S2 in the online Supplementary Data)

CPS C1, N1s are consistent with those reported for $\text{g-C}_3\text{N}_4$ by us [40] and others previously [21, 38, 41–44]. Two points are, however, worth mentioning. First, Pd metal, despite its low loading, was detected (around 0.1 wt%) and similar loading was also measured by ICP (Fig. S3 in the online Supplementary Data): binding energy attributed to Pd $3d_{5/2}$ and $3d_{3/2}$ at 334.4 and 339.4 eV [45]. The presence of S was clear; this was due to the ligand used to link TiO_2 to $\text{g-C}_3\text{N}_4$; binding energy at 168 eV attributed to S $2p_{3/2}$ [46] for TiO_2 - SO_3Na , indicating that the SH from the previous step was oxidized into $-\text{SO}_3^-$ to form negative charges on the TiO_2 surface, supporting Scheme 1.

FT-IR results of the hybrid system are shown in Fig. S4 in the online Supplementary Data, the broadband from 2900 to 3500/cm corresponding to N-H and O-H stretching vibrations; the latter is due to absorbed hydroxyl species [47]. The peaks at 1200–1600/cm correspond to the stretching mode of CN heterocycles as well as the bending modes of adsorbed water and surface hydroxyls [48]. PL measurements were conducted on the best-performing catalyst of 0.1%Pd-20% TiO_2 / C_3N_4 (Fig. 5c) and its precursors, in order to probe into the charge-recombination process [49] at room temperature with an excitation wavelength of 320 nm. There is a significant decrease in the PL intensity

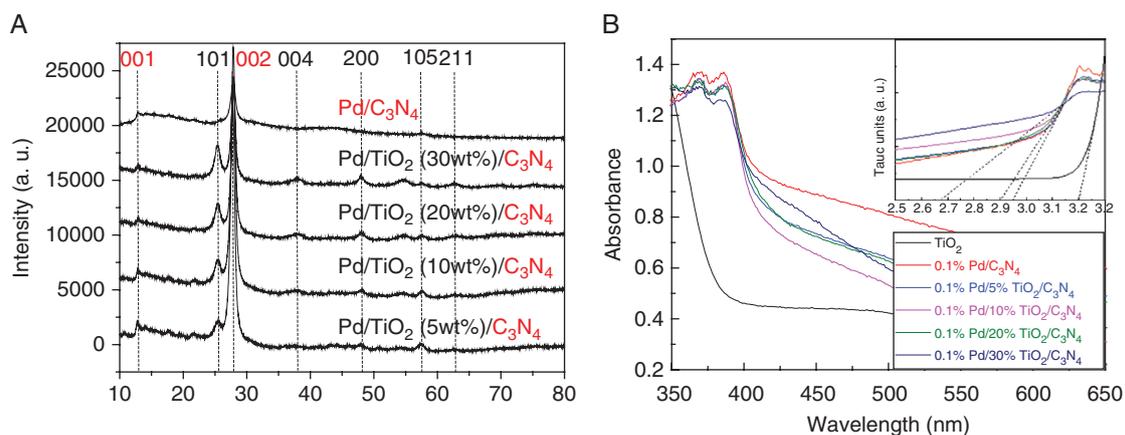


Fig. 2 (a) XRD patterns of 0.1%Pd-g-C₃N₄ and Pd/TiO₂/g-C₃N₄ composites with different mass ratios of TiO₂ and g-C₃N₄. (b) UV-vis diffuse reflectance absorption spectra of the patterns of 0.1%Pd-g-C₃N₄ and Pd-TiO₂/g-C₃N₄ composites with different mass ratios of TiO₂ and g-C₃N₄.

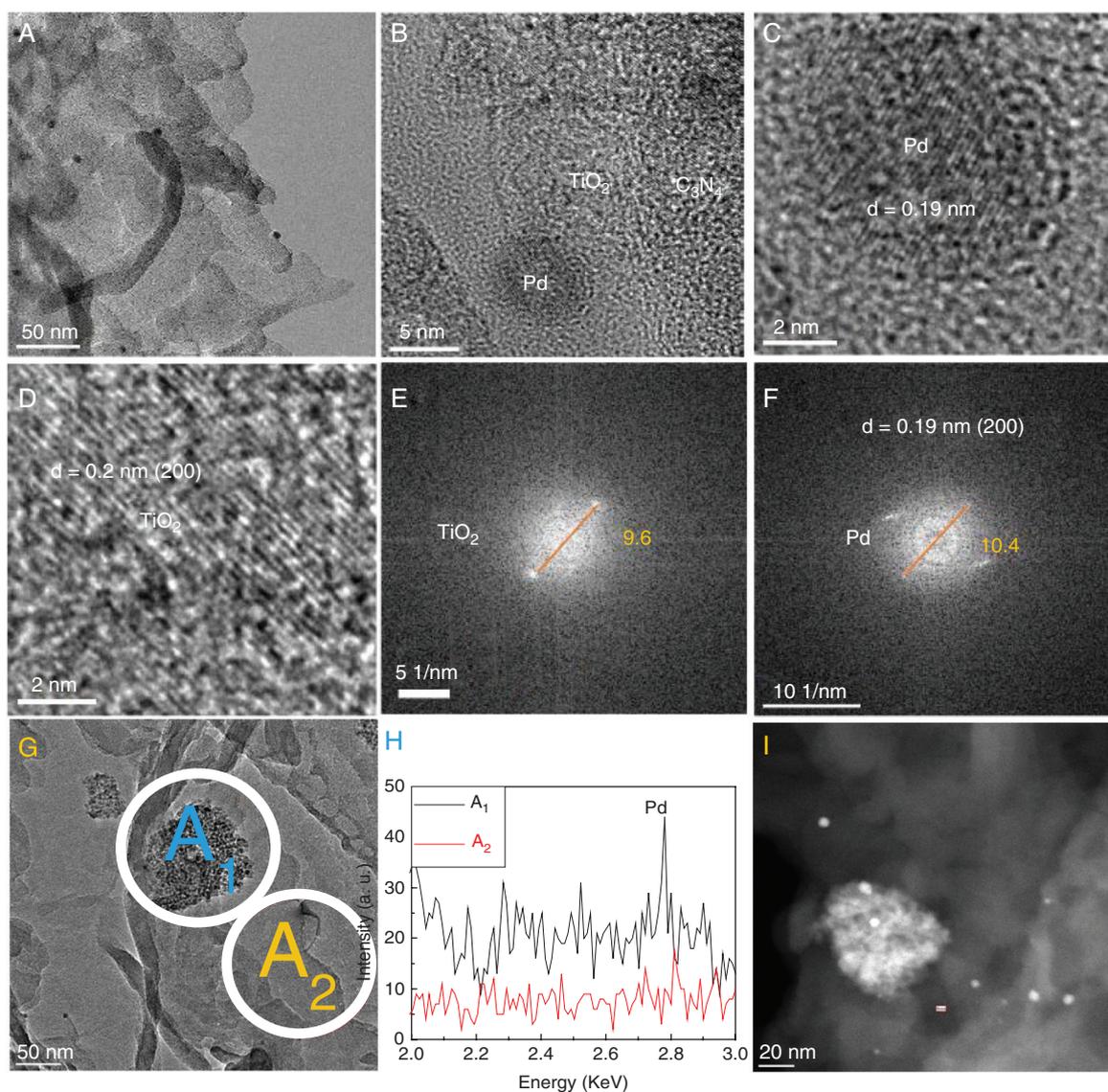


Fig. 3 HR-TEM images of 0.1 wt% Pd/TiO₂/C₃N₄: (a) low magnification, (b) high magnification, (c) lattice fringes of Pd, (d) lattice spacing of TiO₂, (e) FFT diffraction pattern of TiO₂ ($d = 0.2$ nm), (f) FFT diffraction pattern of a Pd particle ($d = 0.19$ nm), (g) TEM images of the areas (circled in orange) that performed EDX analysis, (h) EDX results of the circled areas in (g) and STEM images, (i) STEM image.

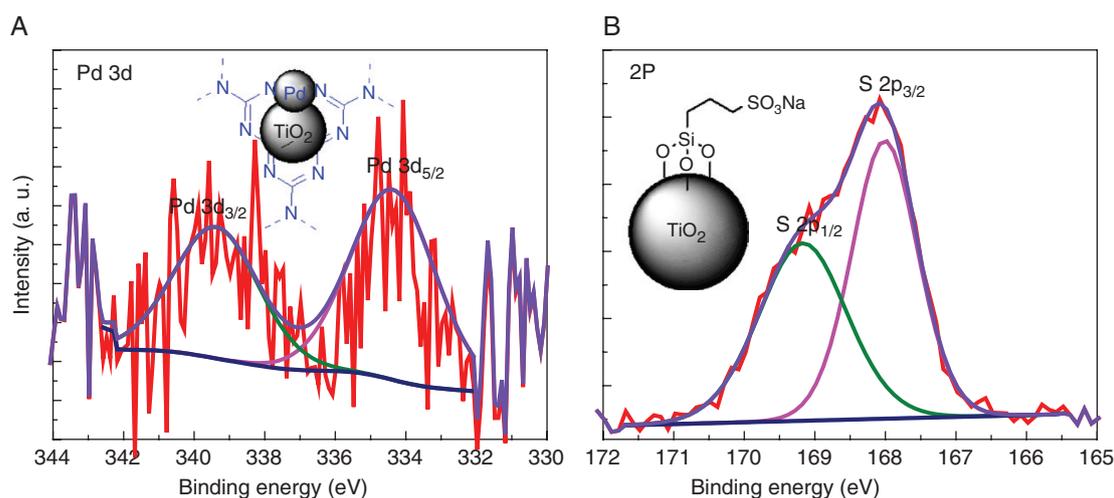


Fig. 4 XPS spectra of 0.1% Pd-TiO₂/C₃N₄ and TiO₂-SO₃Na. (a) Pd 3d, (b) S 2p of TiO₂-SO₃Na.

of 0.1% Pd-20%TiO₂/C₃N₄ compared to those of 20%TiO₂/C₃N₄ and pristine g-C₃N₄. The weak intensity of the PL signal can be attributed to a lower recombination rate of the photo-generated charge carriers [50–52], which is also consistent with other studies over metal-supported TiO₂ [53]. Therefore, one can draw the conclusion that introducing TiO₂ onto the surface of g-C₃N₄ could effectively decrease the electron–hole recombination rates. The addition of Pd, even in such a small amount, has resulted in further decreasing the PL signal, which is again further evidence of electron transfer from the conduction band (CB) of the hybrid semiconductor to Pd metal.

The photocatalytic H₂ production rates over the Pd-TiO₂/g-C₃N₄ composites containing different mass ratios are shown in Fig. 5a and Fig. S5 in the online Supplementary Data. No H₂ production is seen in the absence of either a photocatalyst or light irradiation. Moreover, H₂ rates of as-prepared g-C₃N₄ were very low; it doubled for 0.1% Pd-g-C₃N₄. Because TiO₂ anatase has a band gap of about 390 nm, it is not active under visible-light excitation and was not tested. H₂ production rates increased considerably for Pd/ x%TiO₂/g-C₃N₄ hybrid catalysts with a maximum observed at ~20% in the investigated range, corresponding the maximum AQY of 31% (420–443 nm), whereas further increasing the content of TiO₂ in the composite leads to decrease the H₂ production rate. The H₂ production rate enhancement along with the increase in TiO₂ may be due to the synergistic effect between TiO₂ and g-C₃N₄. The synergic effect could be caused by the electron transfer from HUMO of g-C₃N₄ to the CB of TiO₂ owing to the band positions as shown in Fig. 5d, resulting in a slow charge recombination [21]. Moreover, Pd as an H₂ co-catalyst and meanwhile acting as an electron drain hinders the charge recombination; therefore, the H₂-generation rate was further improved. On the other hand, the decrease in photo-activity upon increasing the TiO₂ mass ratio may be attributed to the fact that there are not enough active sites of g-C₃N₄ to be exposed for

photo-oxidation because of the excess surface coverage by TiO₂. Furthermore, the excess TiO₂ may cause aggregation of TiO₂ nanoparticles that also creates the bulk resistance for electron transfer. Interestingly, the H₂-generation rate of Pd/TiO₂/g-C₃N₄ is extremely low—almost at zero under UV and UV-vis, which means the electron–hole recombination was enhanced on the interface when both semiconductors were activated at the same time. Others also observed the same phenomena [21]. The stability test was conducted on the best-performing catalyst for up to 20 h and almost no degradation was detected (Fig. S6 in the online Supplementary Data).

A fast and steady photocurrent response was observed for each working electrode (Fig. 5b). The highest photocurrent density was achieved on Pd/ TiO₂(20 wt%)/g-C₃N₄, which is more than 10 times that of Pd/C₃N₄ and the great enhancement in the photocurrent can be ascribed to the improved charge separation and accelerated charge transfer by the selective positioning of Pd on TiO₂.

The reported value of the band positions of the highest occupied molecular orbital and the lowest unoccupied molecular orbital (LUMO) of C₃N₄ are –1.3 and 1.4 eV, respectively [54]. The CB and valence band band edge potentials of anatase TiO₂ are reported as –0.5 and 2.7 eV correspondingly [55]. The proposed charge-transfer mechanism is shown in Fig. 5d. Where, the electron–hole pairs are generated by g-C₃N₄ under visible-light illumination, the excited electrons migrate to Pd nanoparticles through the CB of TiO₂ and are then quenched by 2H⁺ to form H₂ molecules. Subsequently, a better electron–hole separation can be achieved through these multiple electron-transfer processes. Accordingly, the hole that was kept on LUMO of g-C₃N₄ will be quenched eventually by TEOA [56]. Furthermore, Fig. S8 in the online Supplementary Data shows a schematic diagram of the work function of Pd as well as the band configurations of TiO₂ [57] and g-C₃N₄ [58, 59] before and after contact. As shown in Fig. 8S in the online Supplementary Data, the electrons from the CB of g-C₃N₄ flow to the Pd through the

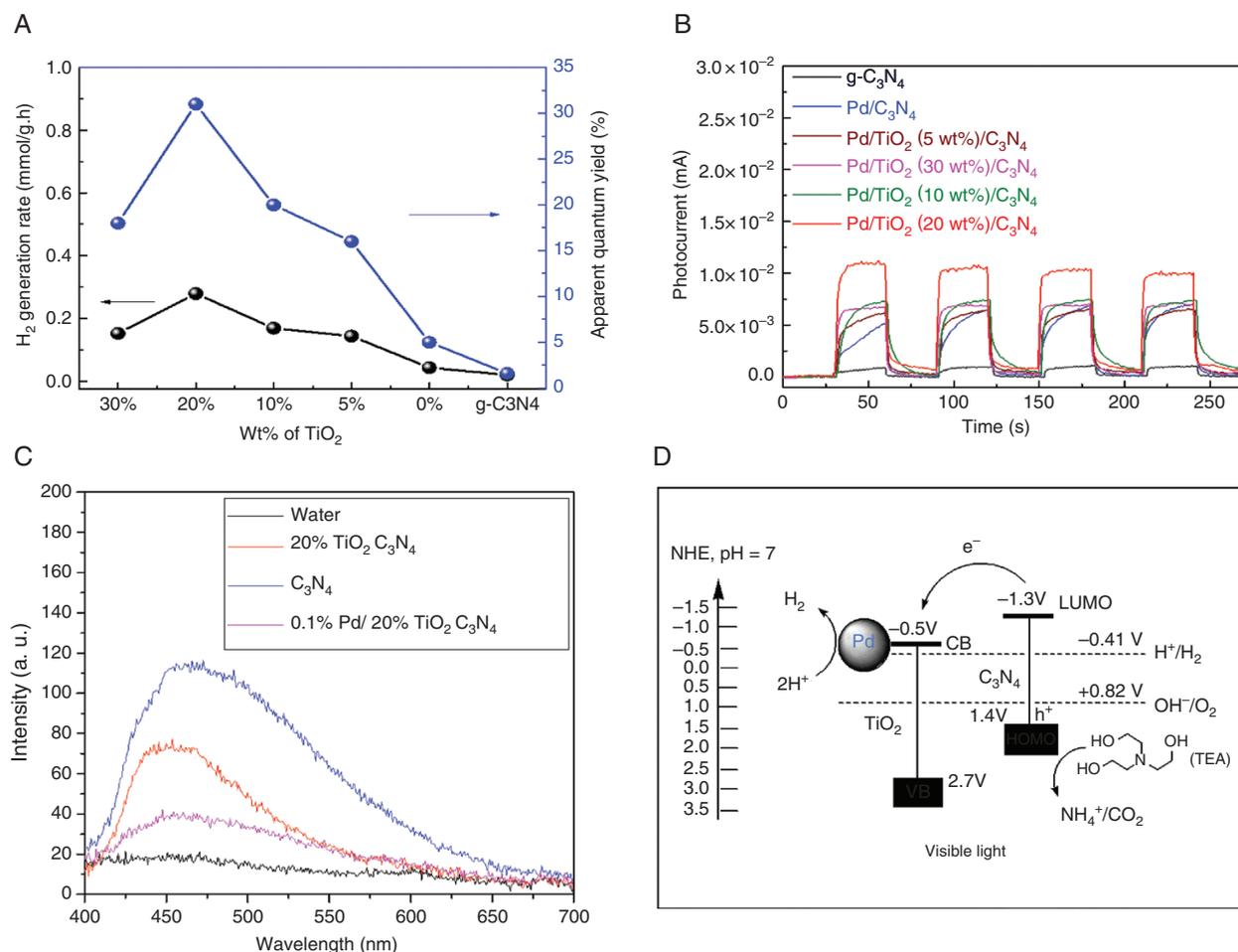


Fig. 5 (a) Photocatalytic H₂ evolution rates of g-C₃N₄, 0.1wt%Pd-g-C₃N₄ and 0.1wt%Pd/TiO₂/g-C₃N₄ composites with different TiO₂/C₃N₄ mass ratios. 20 mg catalyst, 30 mL 10 vol% TAE aqueous solution and 300 W Xe lamp with 23% intensity (43 mW/cm²) for the visible-light ($\lambda \geq 420$ nm) irradiation and the apparent quantum yield (AQY) (420–650 nm), (b) photocurrent responses of the catalysts under Xe lamp irradiation with 400-nm cut-off filter, (c) photoluminescence of pristine g-C₃N₄, 20%TiO₂/C₃N₄ and 0.1% Pd/20%TiO₂/C₃N₄ and (d) proposed charge-transfer mechanism between g-C₃N₄ and TiO₂ through the heterojunction interface under visible-light irradiation.

CB of TiO₂ when the TiO₂ and g-C₃N₄ acquire an equalized Fermi level with the work function of Pd [57]. Furthermore, Pd/TiO₂/g-C₃N₄ nanocomposite inhibits the electron-hole pair recombination ensuing from the enhanced photocatalytic activities of the nanocomposite.

3 Conclusions

In summary, we designed a novel approach to preparing a Pd/TiO₂/g-C₃N₄ hybrid system via four-step processing such as surface modification, mixing, calcination and light-assisted deposition. The synergistic effect of TiO₂ and g-C₃N₄ promoted photo-generated charge separation. Moreover, the Pd was preferentially deposited on the TiO₂ of TiO₂/g-C₃N₄ through light deposition, which remarkably improved the charge transfer between TiO₂ and g-C₃N₄, hindered electron-hole recombination and enhanced the photocatalytic H₂-generation rate. The results show that the photocatalytic H₂-generation rate of optimized sample 0.1%Pd/20% TiO₂/g-C₃N₄ is 14 and 8 times higher than

that of as-prepared g-C₃N₄ and 0.1%Pd/C₃N₄. Most importantly, this work focused on introducing a novel approach using electrostatic interaction and light deposition to fabricate a model nanocomposite of Pd/TiO₂/C₃N₄. In addition, the photocatalytic activities of the Pd/TiO₂/C₃N₄ were measured to further evaluate this approach. The results show that this two-step method is a better way of making nanocomposites. This study provides a novel way of preparing new, stable and visible-light-responsive photocatalysts.

Acknowledgements

The SABIC-CRD financially supported this work. We are also grateful to Dr Hicham Idriss from SABIC for valuable suggestions.

Conflict of interest statement. None declared.

Supplementary data

Supplementary data is available at *Clean Energy* online.

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