Novel multilayer TiO$_2$ heterojunction decorated by low g-C$_3$N$_4$ content and its enhanced photocatalytic activity under UV, visible and solar light irradiation

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In this paper, we used a facile ball milling, microwave radiation and heating treatment method to achieve the surface modification of TiO$_2$ with low g-C$_3$N$_4$ concentration, and a multilayer heterojunction composite with TiO$_2$ as the main part and g-C$_3$N$_4$ as the modification agent was obtained. The obtained materials were analyzed by several characterizations to get information on their chemical composition, crystalline structure, vibrational features and optical properties. The photocatalytic performance was evaluated by degradation of rhodamine B (RhB) and methylene blue (MB) under UV, visible and direct solar light irradiation. Its photocatalytic activity was enhanced depended on the novel structure of g-C$_3$N$_4$/TiO$_2$ hybrid and the special Z-scheme electron-hole transfer model of multilayer heterointerfaces. The material preparation and structural features could be useful for the design and development of other photocatalysts with high photocatalytic activity.

Photocatalysis has been attracting great attention due to increasing energy demands and serious environmental pollution in recent years$^{1-7}$. As an important non-metallic semiconductor photocatalyst, graphitic carbon nitride (g-C$_3$N$_4$) has been widely used in photocatalytic sterilization, hydrogen production, pollutant degradation, and many other fields$^{8-11}$, due to its high stability, unique electronic structure, and narrow band gap (about 2.7 eV) that widens its light response to visible region of 450 nm$^{12}$. However, g-C$_3$N$_4$'s low yield and high e$^-$-h$^+$ pair recombination rate limit its practical application$^{13}$. On the other hand, titanium dioxide (TiO$_2$) is widely used in industry as a photocatalyst due to its high specific surface area, large pore volume, strong oxidizing power, low cost, and high chemical stability$^{14,15}$. But its' narrow light-response range and low quantum efficiency may restrict the practical applications$^{16}$. A new composite that combines g-C$_3$N$_4$ and TiO$_2$ might integrate their advantages, favor the separation of e$^-$-h$^+$ pairs, and improve the photocatalytic performance, which could become a new material with industrial application prospect.

Currently, most of the g-C$_3$N$_4$/TiO$_2$ composites are mainly prepared by hydrothermal method$^{16-18}$, calcination process$^{19-21}$, solvent evaporation method$^{22}$, solvothermal strategy$^{23}$, etc. Gu et al.$^{16}$ developed the flower-like TiO$_2$/g-C$_3$N$_4$ hybrid by surfactant-assisted hydrothermal treatment. Hao et al.$^{19}$ fabricated the g-C$_3$N$_4$/TiO$_2$ heterojunction photocatalysts that preserved a macroscopic network structure with a relatively regular array of macropores by a calcination method using tetrabutyl titanate and melamine as the feedstocks. The morphology of g-C$_3$N$_4$/TiO$_2$ hybrids is also varied via different preparing methods, such as flower-like$^{18}$, hollow nanobox$^{22}$, hollow core@shell structure$^{24}$, corn-shaped structure$^{17}$, multilayered structure$^{25}$, etc.

Table 1 lists several typical g-C$_3$N$_4$/TiO$_2$ hybrid photocatalysts that were prepared using different methods. The g-C$_3$N$_4$/TiO$_2$ hybrids with relatively high contents of g-C$_3$N$_4$ have been studied in recent years, however, little has been known about g-C$_3$N$_4$/TiO$_2$ hybrids with low contents (less than 10 wt%) of g-C$_3$N$_4$. What is the
structure and photocatalytic performance of g-C3N4/TiO2 hybrids with low content of g-C3N4. Could this type of photocatalysts form a heterostructure and enhance photocatalytic efficiency? Further exploration of the structure and photocatalytic properties of the g-C3N4/TiO2 hybrid with low g-C3N4 content (less than 10 wt%) is necessary.

Although the g-C3N4/TiO2 hybrids listed in Table 1 were reported better photocatalytic performance, technical difficulties in preparing g-C3N4/TiO2 hybrids still limit its utilization, including high temperature processing, complex chemical reactions, precursor hydrolysis and agglomerate, and difficult control in yield. Surface modification of photocatalytic materials is an important technical mean to improve the absorbability and photocatalytic properties of materials26,27. As the ball milling method and microwave-assisted heating method are both simple, effective, low-cost and green technique for industrial production to fabricate composite photocatalysts28,29, we combined these two methods to prepare a g-C3N4/TiO2 composite with low content (less than 10 wt%) g-C3N4. In addition, the structural properties and photocatalytic performance of the new g-C3N4/TiO2 composite was studied in this work, which have not been reported by others.

Materials and Methods

Materials. Melamine powder (Pur. >99.0%) used in the experiments was supplied by Aladdin Chemistry Co. Ltd. Nano TiO2 powder (anatase, 100 nm, Pur. >99.8%) was supplied by XuanCheng JingRui photocatalytic material Co. Ltd. RhB, MB, and other chemicals used in the study were purchased from Aladdin Chemistry Co. Ltd. and other China chemical reagent Ltd. All of the chemical reagents were of analytical purity and used without further purification. Deionized water was used throughout this study.

Preparation of photocatalysts. A combination method of a facile ball-milling, microwave radiation and heating treatment was used in this work. The preparation procedure had four steps (Fig. 1): (1) TiO2 powder and a certain amount of melamine were added into a agate ball milling tank (PM2L, Nanjing, China) with a ball: powder ratio of 10: 1, and water was used as a dispersant. The mixtures were milled in the ball milling tank for 2–3 h at a speed of 400 rpm to promote the surface integration of g-C3N4 and TiO2 and to generate a homogeneous heterogeneous interface. (2) The well-mixed wet powder from the ball milling tank was then treated in a microwave oven at 700 W for 10–30 min to get uniform particle size and good compaction. (3) The collected powder was kept for 2–3 h to achieve molecular thermal equilibrium. (4) after that, the powder was heated at 520 °C for 2 h with a heating rate of 5 °C/min in an oven (SX2-2.5-10, Zhengzhou, China). After cooled to room temperature naturally, the final product was collected for further use. By varying the dosage of melamine (30, 40, 50 wt.%), a series of g-C3N4/TiO2 hybrids were obtained, labeled as TCN-1, TCN-2, TCN-3, respectively.

For comparison, pure TiO2 was also treated by the above four steps, labeled as TiO2-W. g-C3N4 was synthesized by directly heating melamine. Typically, a certain amount of melamine powder was poured into an alumina crucible and heated in a muffle furnace at 520 °C for 2 h with a heating rate of 5 °C/min. After cooling to room temperature, the product was collected and ground into powder.

Characterization. Thermogravimetric analysis (TGA) was performed on a Mettler Toledo TGA/DSC2 (Mettler-Toledo, Switzerland) at a heating rate of 5 °C/min from room temperature to 800 °C under air atmosphere. In order to determine the crystal phase composition and the crystallite size of the photocatalysts, the X-ray diffraction (XRD) patterns of the prepared samples were obtained using an X-ray diffractometer (Empyrean, Shimadzu, Japan) with Cu-K radiation (λ = 0.15405 nm, 60 kV, 60 mA). The samples were scanned in the range 2θ = 10°–80°. The morphology and EDX of the prepared samples were analyzed using a JSM-7500F (JEOL, Japan) scanning electron microscope (SEM). Transmission electron microscopy (TEM) and high-resolution transmission electron microscopy (HRTEM) images were obtained on a FEI Tecnai G20-Swin electron microscope (FEI, America) to further investigate the morphology and the structure of the catalysts. X-ray photoelectron spectroscopy (XPS) analysis was performed on an Escalab 250Xi system (Thermo Fisher, America). The thickness of the nanosheets and its multilayer heterointerfaces were determined by atomic force microscopy (AFM) (Bruker dimension icon, USA). UV–Vis diffuse reflectance spectra (DRS) measurements were carried out using

### Table 1. Optimal g-C3N4 mass ratio of several typical g-C3N4/TiO2 hybrids.

| Hybrids            | Preparation method | Target pollutants/products | Light source | Optimal g-C3N4 mass ratio (wt%) | Ref. |
|--------------------|--------------------|----------------------------|--------------|---------------------------------|------|
| g-C3N4/TiO2        | Single-pot microemulsion method | MO, MB, RhB                | 300 W Xe lamp > 420 nm | 17.1                             | 26   |
| g-C3N4/TiO2        | Annealing method   | RhB                        | 350 W Xe lamp > 420 nm | 28.3                             | 19   |
| g-C3N4/TiO2        | Annealing method   | AO7                        | 500 W tungsten halogen lamp > 420 nm | 12          | 20   |
| g-C3N4/TiO2        | Annealing method   | NO                         | 500 W Xe lamp > 420 nm | 15                               | 21   |
| g-C3N4/TiO2        | Annealing method   | MB                          | 500 W Xe lamp > 420 nm | 33                               | 25   |
| g-C3N4/TiO2        | Annealing method   | RhB                        | 500 W Xe lamp > 420 nm | 25.9                             | 30   |
| g-C3N4/TiO2        | Hydrothermal method | Propylene                  | 300 W Xe lamp > 420 nm | 30                               | 40   |
| g-C3N4/TiO2        | Hydrothermal method | RhB                        | 300 W Xe lamp > 420 nm | 60                               | 40   |
| g-C3N4/TiO2        | Hydrothermal method | Phenol                     | 500 W Xe lamp > 420 nm | 66.7                             | 40   |
| g-C3N4/TiO2        | Low-basicity solution chemistry method | MO | 300 W Xe lamp > 420 nm | 35                               | 61   |
| g-C3N4/TiO2        | Solegel method     | Hydrogen                   | 500 W Xe lamp > 420 nm | 67                               | 61   |
| g-C3N4/TiO2        | Ball milling method | Hydrogen                   | 450 W Xe lamp > 420 nm | 50                               | 62   |
a UV-3600 (Shimadzu, Japan) spectrophotometer equipped with an integrating sphere attachment. The analysis range was from 200 to 800 nm, and BaSO₄ was used as a reflectance standard. Photoluminescence (PL) emission spectra were obtained on a Fluorescence Spectrometer (F-7000, Hitachi, Japan). The excitation wavelength was 230–900 nm with the scanning speed of 2400 nm/min. The widths of excitation slit and emission slit were both 5 nm.

Photocatalytic tests. The photocatalytic activities of as-obtained samples were measured by the degradation of RhB and MB under both UV and direct solar light irradiation. 50 mg of photocatalyst was suspended in 50 mL of an aqueous solution containing 20 mg/L organic dye. Before irradiation, the suspensions were stirred for 60 min in the dark to establish adsorption–desorption equilibrium. To study dye degradation under UV light, a high-pressure 300 W mercury lamp with the radiation of 365 nm was used as UV light source. And the average irradiance measured at the sample was about 500–800 W·m⁻² (measured by solar power meter, TES-1333R) and the reactor was directly exposed to sunlight for solar light degradation. A 300 W xenon lamp with a cutoff filter (λ > 420 nm) was used as visible light source to study the degradation efficiency under visible light. At given time intervals, 4 mL of the suspension was sampled to be centrifuged at 10000 rpm for 10 min to remove the particles. Then, the concentration of organic dyes in solution was analyzed using an Alpha-1506 UV–vis spectroscopy (Shanghai Lab-Spectrum Instruments Co. Ltd., China) at the wavelength of 554 nm and 664 nm for RhB and MB, respectively. The degradation for each sample was measured 3 times.

Results and Discussion
Characterization of prepared photocatalysts. Component analysis. Figure 2 shows TGA analysis results of TiO₂, g-C₃N₄ and TCN-2 photocatalyst. The TGA analysis curve of TCN-2 indicates the existence of g-C₃N₄ in the composite. And other reports also confirm that melamine can produce g-C₃N₄ at about 500 °C. The TCN-2 sample has two weight loss regions: the former weight loss occurred at around 100–300 °C due to desorption of surface bound water and the dehydrogenation and deamination of melamine to produce g-C₃N₄, the later weight loss from 300 °C to 500 °C was due to the combustion of g-C₃N₄ in the air atmosphere. And for the TCN-2 sample, the later weight loss is about 8.64% in the temperature range of 300–500 °C, indicating that the TCN-2 sample contains about 8.64% g-C₃N₄.

The XRD patterns of the samples are shown in Fig. 3. For comparison, the XRD patterns for pure TiO₂ and g-C₃N₄ are also given. For g-C₃N₄, the main diffraction peak at 2θ = 27.5° indicates its graphite-like structure. Another relative weak peak located at 13.1° may be due to the in-planar repeat period, and just like the hole-to-hole distance among the N-bridged tri-s-triazine units. It can be seen that the TiO₂ powder is in good agreement with anatase phase of TiO₂ indexed to the data in the JCPDS cards (21–1272). No obvious XRD peaks of g-C₃N₄ are observed in the TCN-2 hybrid may be due to the small amount of g-C₃N₄ whose peaks are not strong enough. These tests show that TiO₂ is the main part of TCN-2. Furthermore, no obvious XRD peaks of other new crystal phases are found in TCN-2, indicating that the hybridization of TiO₂ with g-C₃N₄ did not change the crystal structure of TiO₂ in our experiment. The peak of TCN-2 is slightly sharpened compared to pure TiO₂, indicating a better crystallinity than that before the combination.

The XPS was carried out to further analyze the surface chemical composition and chemical status of elements in TCN-2 (Fig. 4). The XPS spectra of C 1 s for TCN-2, with two peaks at 284.6 and 288.0 eV are shown in Fig. 4a. The former peak is attributed to C–C, which originate from the adventitious carbon. However, the latter one is ascribed to sp²-hybridized carbon in N-containing aromatic ring (N–C=N). The N 1 s XPS binding energy can be fitted into three peaks centered at 398.6, 399.5, 400.6 eV, which belong to sp²-hybridized nitrogen (C–N=C), the tertiary nitrogen N–(C), and the free amino groups (C–N–H), respectively. The Ti 2p spectrum has two peaks at binding energies of 458.6 (Ti 2p₃/2) and 464.3 eV (Ti 2p₁/2) (Fig. 4c). In Fig. 4d, the O 1 s spectrum is fitted with two peaks corresponding the Ti–O bond (529.8 eV) and O–H bond (531.6 eV), suggesting the presence of water molecule or a hydroxyl group on the surface of the TCN-2 composite microspheres. The XPS results indicate that g-C₃N₄ was generated through melamine during the reconstruction of TiO₂. And there
are some differences compared to the g-C₃N₄/TiO₂ with relatively high g-C₃N₄ content⁴⁰,⁴¹, which may be due to the changes of the binding force between atoms.

**Morphology analysis.** The SEM image of the prepared g-C₃N₄ shows the irregular non-nanoscale structure (Fig. 5a). The photograph of spherical TiO₂ particles is shown in Fig. 5b, and the average particle size is about 100 nm through the particle size distribution analysis. Figure 5c,d show the SEM and EDS images of TCN-2. Compared with Fig. 5b, TCN-2 is a lamellar structure (most obviously shown in the circled part of Fig. 5c), and it is more compact than pure TiO₂ in Fig. 5b. The structural differences between TiO₂ and TCN-2 may be the small amount of g-C₃N₄ that induced changes in the structure of TiO₂. EDS spectra shows the presence of elements C, N, Ti, and O. Their weight and atomic percentages are shown in the table given as the inset of Fig. 5d. It also shows the low content of g-C₃N₄ in accordance with the XRD and TGA results.

The TEM image in Fig. 5e is in good agreement with the result of Fig. 5c, and it is further confirmed the multilayer heterointerfaces of TCN-2. In Fig. 5f,g, it is also easy to find the multilayer structure of TCN-2. In Fig. 5g, the two different lattice fringes of 0.205 nm and 0.246 nm correspond to the (200) and (004) plane of anatase TiO₂. And it could be found that the crystal surface of (200) deflected about 15°, which further confirms the interfacial misfit dislocation of TiO₂ films between the two different planes in Fig. 5g, so that there are much more mismatch in the multilayer heterointerfaces. Additionally, some parts of the lattices of TiO₂ are indistinct (Fig. 5g), which can be resulted from a thin layer of g-C₃N₄ covered with TiO₂. Thus, it is the stratified crossed structure of TiO₂ films induced by g-C₃N₄ (Figs 4g and 5f) that formed the multilayer hybrid in Fig. 5c. Figure 5h shows that the TCN-2 sample has other two lattice fringes: one is measured to be 0.352 nm matching the (101) plane of anatase TiO₂⁴³; the other one is measured to be 0.336 nm matching the (002) plane of g-C₃N₄⁴⁴. The observations further suggest that g-C₃N₄ distributed at the boundary of TiO₂ films, and it could be the small
amount of g-C$_3$N$_4$ that induced the multilayer heterointerfaces of the composite. The results of AFM (Fig. 6) showed that the average interlayer distance of the multilayer heterointerfaces is about 15 nm.

From the TGA, XRD and XPS analysis, it is clear that there was a combination of g-C$_3$N$_4$ and TiO$_2$. During the heat treatment, TiO$_2$ reconstructed its crystal structure, and the g-C$_3$N$_4$ fabricated from melamine combined with the surface of TiO$_2$ to form new interface. It is known that when the crystal surface grows with different substances, the chemical bonding direction of the surface chemisorption will be different, leading to the deflection of the crystal self-assembly growth$^{45-47}$. During the self-assembly reconfiguration of TiO$_2$, the new interface structure of g-C$_3$N$_4$ and TiO$_2$ was formed. And the distortion of the interface leaded to the deflection of the crystal, thus reducing the interface energy to achieve the overall thermal stability, which further caused the deflection of about 15° for the crystal surface of (200). And the interfacial misfit dislocation of TiO$_2$ films induced by g-C$_3$N$_4$ resulted in the generation of multilayer heterointerfaces. This is the major reason that caused the morphology of TCN-2 different from other structures. The above investigations strongly confirmed that the as-prepared composite including both g-C$_3$N$_4$ and TiO$_2$ forms a heterostructure rather than a physical mixture. Therefore, the formation of a heterojunction between g-C$_3$N$_4$ and TiO$_2$ would be a system to achieve enhanced electron-hole separation.

The structure of the TCN in our work is somewhat similar to those studied with high g-C$_3$N$_4$ concentrations$^{25,41,48,49}$, which suggests that changing the amount of g-C$_3$N$_4$ to a small range can also form a heterogenous structure. Furthermore, the composite in our work showed some new features, such as the multilayer structure of the hybrid and the 15 nm average interlayer distance (Fig. 6). The above TGA, XRD and XPS results showed that the reconstructed TiO$_2$ crystals were accompanied with melamine thermal polymerization, and the low concentration of g-C$_3$N$_4$ diffused to the crystal interfaces of the reconstructed TiO$_2$ at 520°C, resulting in the formation of g-C$_3$N$_4$ with (C-N=), (N-C=) and N-(C)$_3$ bonding characteristics.

**Optical properties analysis.** The UV-vis diffuse reflectance spectra of the photocatalysts are shown in Fig. 7a. And the band gap energy of the photocatalysts was calculated by the following equation$^{50}$: $\alpha$ = $A(h\nu - E_g)^{n/2}$. In this equation, $\alpha$, $h$, $v$, $A$, and $E_g$ are absorption coefficient, Planck constant, light frequency, proportionality, and band gap energy, respectively; $n$ keys the properties of the transition in a Semiconductor ($n$ = 1 for direct transition, and $n$ = 4 for indirect transition). The values of $n$ for g-C$_3$N$_4$ and TiO$_2$ are both$^{41,52}$. The calculated results are shown in Fig. 7b, and the absorption onset of g-C$_3$N$_4$ was found at about 466 nm, corresponding to band gap energy of 2.66 eV. This result is consistent with other groups’ work$^{53}$. Meanwhile, the band gap of TiO$_2$ was estimated to be about 3.18 eV with the absorption edge of 389 nm. After combined with g-C$_3$N$_4$, the absorbance of multilayer TCN-2 is slightly extended to the visible region as the presence of g-C$_3$N$_4$. Combined with the analysis of Fig. 5, it could be concluded that the existence of g-C$_3$N$_4$ had impact on the band gap of TiO$_2$. 

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**Figure 4.** High-resolution XPS spectra of (a) C 1 s, (b) N 1 s, (c) Ti 2p, and (d) O 1 s for TCN-2.
The band edge positions of CB and VB of semiconductor could be determined by a simple approach. The valance band edge (E$_{VB}$) and conduction band edge (E$_{CB}$) of a semiconductor at the point of zero charge (pHzPC) can be predicted by the following two equation:\[ E_{VB} = \chi - E_g \] and \[ E_{CB} = -E_{VB} - E_g \]. The $\chi$ values for g-C$_3$N$_4$ and TiO$_2$ are 4.73 eV and 5.81 eV\cite{19}, respectively. The band gap energies of photocatalysts are shown in Table 2. Compared to the g-C$_3$N$_4$/TiO$_2$ hybrids with high g-C$_3$N$_4$ content\cite{49,54}, the band gap did not change obviously, which is due to the small amount of g-C$_3$N$_4$ did not influence the band gap of the composite significantly.

**Evaluation of photocatalytic activity.** The degradation of RhB and MB by prepared photocatalysts was studied under UV and solar light illumination. The photocatalytic efficiencies of RhB and MB were calculated from the following expression: \[ \eta = \left( \frac{C_0 - C_t}{C_0} \right) \times 100\% \], where $\eta$ is the photocatalytic efficiency; $C_0$ is the concentration of reactant before illumination; $C_t$ is the concentration of reactant after illumination time $t$. The pseudo-first-order model is expressed by the equation:\[ \ln \left( \frac{C_0}{C_t} \right) = k t \], where $C_0$ and $C$ are the concentrations of dye in solution at time 0 and $t$ respectively, and $k$ is the pseudo-first-order rate constant.

As shown in Fig. 8a, after 30 min UV illumination, the photocatalytic efficiencies of TCN-2 for RhB and MB were 89.44% and 95.48%, respectively, which were higher than that of g-C$_3$N$_4$ (72.09% for RhB; 65.06% for MB) and pure TiO$_2$ (73.42% for RhB; 89.77% for MB). The photocatalytic efficiencies of TiO$_2$–W for RhB and MB were 77.24% and 91.73%, respectively, showing a slight increase in the degradation efficiency compared to pure TiO$_2$. As the photocatalytic degradation process of organic dyes follows first-order kinetics\cite{55}, the first-order kinetics of RhB and MB degradation by the different photocatalysts are illustrated in Fig. 8b. The rate constants of TiO$_2$–W for RhB and MB were calculated to be 0.0841 min$^{-1}$ and 0.107 min$^{-1}$, respectively. Moreover, the three TCN samples also showed an improved photocatalytic activity under solar light irradiation (Fig. 9). The degradation efficiencies of all TCN samples reached more than 90% in 60 min, which were much better than pure
TiO₂, TiO₂-W and g-C₃N₄. And the degradation rates of TCNs for RhB and MB were more than 3 times that of pure TiO₂, TiO₂-W and g-C₃N₄. In Fig. 10, under visible light (λ > 420 nm) irradiation for 4 h, the degradation efficiencies of TCN-2 sample reached 78.27% and 82.62% for RhB and MB, respectively, which were also better than pure TiO₂, TiO₂-W and g-C₃N₄. And the degradation rates of TCN-2 for RhB and MB were more than 2 times that of pure TiO₂, TiO₂-W and g-C₃N₄. And there was no significant difference in the degradation efficiency between TiO₂-W and TiO₂. This further indicates that the low-content g-C₃N₄ doping as a surface modifying agent can improve the photocatalytic activity of TiO₂. Although the slight change of g-C₃N₄ contents in the three TCN samples had a little difference in the photocatalytic performance under solar light, which may be due to the small change of hybrid’s band gap caused by the low g-C₃N₄ contents, the degradation efficiencies of the three samples could also further illustrate that it is feasible to improve the photocatalytic performance of TiO₂ with low g-C₃N₄ content. Thus, this method would be beneficial to TiO₂ optimization and could avoid the shortage of low yield of g-C₃N₄. The application of the ball-milling, microwave radiation and heating treatment method can not only effectively achieve good degradation efficiency but also reduce the cost of photocatalytic materials.

Reusability is important for photocatalysts in practical applications. In the photocatalytic cycling test, the TCN-2 sample was recycled by centrifugation and then dried in an oven at 60°C for 24 h. Then, the sample was directly applied into the next recycle without any further disposal. From Fig. 11, it can be found that the

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**Table 2.** The χ, Eₓ, Eₘ, Eₚ and Eₜ for TiO₂ and g-C₃N₄ at the point of zero charge. χ, Eₓ, Eₘ, Eₚ and Eₜ denote absolute electronegativity, calculated CB edge, calculated VB position and band gap energy, respectively.

| Sample  | χ (eV) | Eₓ (eV) | Eₘ (eV) | Eₚ (eV) | Eₜ (eV) |
|---------|--------|---------|---------|---------|---------|
| TiO₂    | 5.81   | −0.28   | 2.90    | 3.18    |
| g-C₃N₄  | 4.73   | −1.10   | 1.56    | 2.66    |

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Figure 7. (a) UV–vis diffuse reflectance spectra of the photocatalysts and (b) their corresponding Tauc’s plots.
photocatalytic activity of the TCN-2 hybrid was stable, and it changed little after used 3 times. Thus the compound is expected to be applied in practical photocatalysis.

Proposed reaction mechanisms. PL emission spectra. PL emission spectra can be employed to investigate the migration, transfer and separation efficiency of the photoinduced carriers in the photocatalytic reactions, due to the intensity of PL emission spectra could indicate the recombination speed of photoexcited electron-hole pairs. In Fig. 12, in the case of the pure g-C₃N₄ sample, the strong emission peak was at about 466 nm, which was consistent with the UV–vis DRS analysis. This strong peak is ascribed to the band-band PL phenomenon with the energy of light approximately corresponding to Huang’s work. The PL spectrum of pure TiO₂ was in the wavelength range between 350 and 550 nm, which is attributed to the emission of bandgap transition and excitonic PL that resulted from the surface oxygen vacancies and defects. When compared to the pure TiO₂ and g-C₃N₄, the emission spectrum of TCN-2 hybrid decreased sharply, which indicates that the e⁻–h⁺ pair recombination rate is much lower in the multilayer heterointerfaces, demonstrating the fact that g-C₃N₄ layers with two-dimensional π-conjugation structure could serve as an effective carrier-transfer channel and suppress the direct recombination of e⁻–h⁺ pairs in TiO₂. As the PL intensity of TCN-2 was much weaker than that of the pure TiO₂ and g-C₃N₄, it is not difficult to understand that the TCN-2 sample exhibits an enhanced photocatalytic performance on the decomposition of RhB and MB.

Roles of reactive species. It is generally accepted that the •O₂⁻, •OH and h⁺ are the major reactive species in the photocatalytic oxidation process. In our work, the benzoquinone (BQ) was applied to reduce the superoxide radical (•O₂⁻), the isopropanol (IPA) was employed to impair the hydroxyl radical (•OH), and the ammonium oxalate (AO) was used to remove hole (h⁺). The results are shown in Fig. 13. It could be found that, whether it is the degradation of RhB or MB, it is remarkably suppressed when the BQ and IPA was added into the reaction.
solutions, while the impact caused by AO was insignificant. Therefore, \( \bullet \text{O}_2^- \) and \( \bullet \text{OH} \) are the major reactive species for TCN samples in the photocatalytic reaction system and the influencing degree is \( \bullet \text{O}_2^- > \bullet \text{OH} > h^+ \).

**Photocatalytic mechanisms.** Based on the results of the photocatalytic activities over catalysts and the trapping experiments of the photogenerated carriers, the separation processes of photoexcited electron-hole can be described in Fig. 14. From Table 2, the CB bottom of g-C\(_3\)N\(_4\) (−1.10 eV) is more negative than that of TiO\(_2\) (−0.28 eV), nevertheless the VB top of TiO\(_2\) (2.90 eV) is more positive than that of g-C\(_3\)N\(_4\) (1.56 eV). Thus, these accumulated electrons in the CB of TiO\(_2\) cannot reduce O\(_2\) to yield \( \bullet \text{O}_2^- \)\((\text{O}_2/\bullet \text{O}_2^- = -0.33 \text{ eV}^{58})\), and the holes in the VB of g-C\(_3\)N\(_4\) cannot oxidize OH\(^-\) to give \( \bullet \text{OH}(\text{OH}^-/\bullet \text{OH} = 2.4 \text{ eV}^{13})\). Therefore, the photocatalyst transfer scheme of charge carriers is a special Z-scheme multilayer heterointerface model rather than a traditional one\(^{21,59,60}\). Under this mechanism, the fast combination is achieved between the photoexcited holes in the VB of g-C\(_3\)N\(_4\) and photoexcited electrons in the CB of TiO\(_2\). At the same time, the electrons in the CB of g-C\(_3\)N\(_4\) with more negative potential reduce the molecular O\(_2\) to yield \( \bullet \text{O}_2^- \); and the holes in the VB of TiO\(_2\) with more positive potential oxidize OH\(^-\) to produce abundant \( \bullet \text{OH} \) radicals. Furthermore, in the special crystal interfaces of the multilayer g-C\(_3\)N\(_4\)/TiO\(_2\) composite, the charge distribution has a superposition effect based on a special Z-scheme electron-hole transfer model. And from Fig. 6, several active surfaces and interfaces can be formed at the multilayer step interfaces, and the \( \bullet \text{O}_2^- \) and OH\(^-\) can be produced at different layers respectively, thus leading to the better photocatalytic efficiency.

**Conclusions**
A novel multilayered g-C\(_3\)N\(_4\)/TiO\(_2\) composite with low g-C\(_3\)N\(_4\) content was prepared via a simple ball milling, microwave radiation and heating treatment method. This method, with low temperature and simple process to produce heterojunction materials, achieved the surface modification of TiO\(_2\) by a small amount of g-C\(_3\)N\(_4\) and feasible for industrial production. The prepared g-C\(_3\)N\(_4\)/TiO\(_2\) hybrid decorated by low g-C\(_3\)N\(_4\) content has the...
Figure 10. The degradation efficiency (a) and the kinetics (b) of RhB and MB by different photocatalysts under visible light (λ > 420 nm) irradiation in 4 h.

Figure 11. The recycling tests of TCN-2 hybrid under solar light irradiation.
following three features: 1) the multilayer structure shows an average interlayer distance of about 15 nm; 2) the low content of g-C3N4 fabricated from melamine spreads over the crystal surface boundary of TiO2 and results in the deflection of TiO2 films; 3) it is the deflection of different TiO2 crystal surfaces that further formed the multilayer heterointerfaces, leading to an improved photocatalytic activity through the degradation of RhB and MB. Furthermore, through our surface modification method, the small amount of g-C3N4 (less than 10 wt%) with the C-N=C, N-C=N and N-(C)3 bonding characteristics clustered at the crystal interfaces of the multilayer heterostructure, promoting the formation of multilayer steps with uniform thickness (15 nm). At such surface...
interface with special crystal interfaces and step defects, the charge distribution has a superposition effect based on a special Z-scheme electron-hole transfer model, and the $\cdot \text{O}_2^-$ \text{g-C}_3\text{N}_4$ and $\text{OH}^-$ \text{TiO}_2 can be produced at different layers respectively. Thus, the photocatalytic property of the material is effectively improved. As different preparing methods and the changes of \text{g-C}_3\text{N}_4 content could lead to different properties of the heterostructures, we will continue to further explore and study in this filed.

References
1. Chae, S. Y. et al. Insight into Charge Separation in \text{WO}_3/\text{BiVO}_4 Heterojunction for Solar Water Splitting. \textit{ACS Appl. Mater. Inter.} \textbf{9}, 19780–19790 (2017).
2. Huo, R. et al. Self-assembly synthesis of \text{BiVO}_4/\text{Polydopamine}/\text{g-C}_3\text{N}_4 with enhanced visible light photocatalytic performance. \textit{Mater. Res. Bull.} \textbf{98}, 225–230 (2018).
3. Reshak, A. H. Sulfide oxide X\text{ZnSO} \text{g-C}_3\text{N}_4 preparing methods and the changes of \text{g-C}_3\text{N}_4 content could lead to different properties of the heterostructures, we will continue to further explore and study in this filed.

Figure 14. Proposed mechanism for \text{g-C}_3\text{N}_4/\text{TiO}_2 photocatalysis under direct solar light irradiation.
23. Huang, Z. A. et al. Effect of contact interface between TiO₂ and g-C₃N₄ on the photoreactivity of g-C₃N₄/TiO₂ photocatalyst: (001) vs (101) facets of TiO₂. Appl. Catal. B: Environ. 164, 420–427 (2015).
24. Guo, N. et al. Novel mesoporous TiO₂@g-C₃N₄ hollow core/shell heterojunction with enhanced photocatalytic activity for water treatment and H₂ production under simulated sunlight. J. Hazard. Mater. 353, 80–88 (2018).
25. Fu, M. et al. Growth of g-C₃N₄ Layer on Commercial TiO₂ for Enhanced Visible Light Photocatalytic Activity. J. Nanomater. 2014, 1–8 (2014).
26. Neoh, K. G. et al. Surface modification strategies for combating catheter-related complications: recent advances and challenges. J. Mater. Chem. B 5, 2045–2067 (2017).
27. Méndez-Medrano, M. G. et al. Surface Modification of TiO₂ with Ag Nanoparticles and CuO Nanoclusters for Application in Photocatalysis. J. Phys. Chem. C 120, 5143–5154 (2016).
28. James, S. L. et al. Mechanochemistry: opportunities for new and cleaner synthesis. Chem. Soc. Rev. 41, 413–47 (2012).
29. Majid, F. et al. Microwave-assisted sol–gel synthesis of Bi₂O₃ nanoparticles. J. Sol-Gel Sci. Technol. 74, 310–319 (2014).
30. Sun, Z. et al. Facile synthesis of two clay minerals supported graphitic carbon nitride composites as highly efficient visible-light-driven photocatalysts. J. Colloid Interf. Sci. 511, 268–276 (2018).
31. Zada, A. et al. Improved visible-light activities for degrading pollutants on TiO₂/g-C₃N₄ nanocomposites by decorating SPR Au nanoparticles and 2,4-dichlorophenol decomposition path. J. Hazard. Mater. 342, 715–723 (2018).
32. Zhou, J. et al. Photocatalytic enhancement of hybrid g-C₃N₄/TiO₂ prepared via ball milling method. Phys. Chem. Chem. Phys. 17, 3647–52 (2015).
33. Yu, J. et al. Enhanced photocatalytic performance of direct Z-scheme g-C₃N₄–TiO₂ photocatalysts for the decomposition of formaldehyde in air. Phys. Chem. Chem. Phys. 15, 16883–90 (2013).
34. Li, Z. et al. Highly efficient hydrogen evolution over Co(OH)₂ nanoparticles modified g-C₃N₄ co-sensitized by Eosin Y and Rose Bengal under Visible Light Irradiation. Appl. Catal. B: Environ. 188, 56–64 (2016).
35. Liu, Y. et al. Enhanced visible-light photocatalytic activity of Z-scheme graphitic carbon nitride/oxygen vacancy-rich zinc oxide hybrid photocatalysts. Chinese J. Catal. 36, 2135–2144 (2015).
36. Li, X. et al. Hydrogenated Defects in Graphitic Carbon Nitride Nanosheets for Improved Photocatalytic Hydrogen Evolution. J. Phys. Chem. C 119, 14958–14964 (2015).
37. Xuang, P.-Y. et al. g-C₃N₄ decorated ZnO nanorod arrays for enhanced photoelectrocatalytic performance. Appl. Surf. Sci. 358, 296–303 (2015).
38. Chen, Y. et al. Construction of heterostructured g-C₃N₄/Ag/TiO₂ microsheres with enhanced photocatalysis performance under visible-light irradiation. ACS Appl. Mater. Inter. 6, 14040–14 (2014).
39. Kim, J. et al. Individually carbon-coated and electrostatic-force-derived graphene-oxde-wrapped lithium titanium oxide nanofibers as anode material for lithium-ion batteries. Electrochim. Acta 199, 35–44 (2016).
40. Zhou, D. et al. Facile Construction of g-C₃N₄ Nanosheets/TiO₂ Nanotube Arrays as Z-Scheme Photocatalyst with Enhanced Visible-Light Performance. ChemCatChem 8, 3064–3073 (2016).
41. Zang, Y. et al. Hybridization of brookite TiO₂ with g-C₃N₄ a visible-light-driven photocatalyst for As⁵⁺ oxidation, MO degradation and water splitting for hydrogen evolution. J. Mater. Chem. A, 15774–15780 (2014).
42. Ye, L. et al. Synthesis of anatase TiO₂ nanocrystals with [011], [001] or [010] single facets of 90% level exposure and liquid-phase photocatalytic reduction and oxidation activity orders. J. Mater. Chem. A 1, 10532 (2013).
43. Gun, Y. et al. Joint Effects of Photocative TiO₂ and Fluoride-Doping on SnO₂ Inverse Opal Nanochitectures for Solar Water Splitting. ACS Appl. Mater. Inter. 7, 20929–303 (2015).
44. Li, G. et al. Enhanced visible-light driven photocatalytic inactivation of Escherichia coli using g-C₃N₄/TiO₂ hybrid photocatalyst synthesized using a hydrothermal-calcination approach. Water Res. 86, 17–24 (2015).
45. Chūn, Y. et al. Ab initio total energy study of ZnO adsorption on a graphene (0001) surface. Phys. Rev. B 70, 8 (2004).
46. Huang, P. et al. A DFT study of STO adsorption on GaN (0001) surface. Chem. Phys. Lett. 640, 119–123 (2015).
47. Yang, C. et al. Modelling and simulation of reaction mechanisms in early growth of STO thin films from ab initio calculations. Comp. Mater. Sci. 49, 845–849 (2010).
48. Fu, M. et al. A Cost-Effective Solid-State Approach to Synthesize g-C₃N₄ Coated TiO₂ Nanocomposites with Enhanced Visible Light Photocatalytic Activity. Int. J. Photoenergy 2013, 1–7 (2013).
49. Tong, Z. et al. Biomimetic fabrication of g-C₃N₄/TiO₂ nanosheets with enhanced photocatalytic activity toward organic pollutant degradation. Chem. Eng. J 260, 117–125 (2014).
50. Zhang, J. et al. Design of a direct Z-scheme photocatalyst: preparation and characterization of Bi₂O₃/g-C₃N₄ with high visible light activity. J. Hazard. Mater. 280, 713–22 (2014).
51. Chen, S. et al. Study on the separation mechanisms of photogenerated electrons and holes for composite photocatalysts g-C₃N₄–WO₃, Appl. Catal. B: Environ. 150–151, 564–573 (2014).
52. Sridharan, K. et al. Novel visible light active graphitic C₃N₄–TiO₂ composite photocatalyst: Synergistic synthesis, growth and photocatalytic treatment of hazardous pollutants. Appl. Catal. B: Environ. 142–143, 718–728 (2013).
53. Giannakopoulou, T. et al. Tailoring the energy band gap and edges’ potentials of g-C₃N₄/TiO₂ composite photocatalysts for NOx removal. Chem. Eng. J 310, 571–580 (2017).
54. Zhao, S. et al. g-C₃N₄/TiO₂ hybrid photocatalyst with wide absorption wavelength range and effective photogenerated charge separation. Sep. Purif. Technol. 99, 50–54 (2012).
55. Bansal, P. et al. Comparative study of catalytic activity of ZrO₂ nanoparticles for sonocatalytic and photocatalytic degradation of cationic and anionic dyes. Chem. Eng. J 280, 475–485 (2015).
56. Yu, J. et al. Enhanced photocatalytic activity of mesoporous TiO₂ aggregates by embedding carbon nanotubes as electron-transfer channel. Phys. Chem. Chem. Phys. 13, 3491–501 (2011).
57. Li, W. et al. Evidence for the Active Species Involved in the Photodegradation Process of Methyl Orange on TiO₂. J. Phys. Chem. C 116, 3552–3560 (2012).
58. Gao, S. et al. Photocatalytic removal of tetraphemobisphenol A by magnetically separable flower-like BiOBr/BiOI/Fe₂O₃ hybrid nanocomposites under visible-light irradiation. J. Hazard. Mater. 331, 1–12 (2017).
59. Wei, Z. et al. Photocatalytic degradation of phenol-containing wastewater by TiO₂/g-C₃N₄ hybrid heterostructure thin film. Appl. Catal. B: Environ. 201, 600–606 (2017).
60. Wei, X. et al. Facile in situ synthesis of plasmonic nanoparticle-decorated g-C₃N₄/TiO₂ heterojunction nanofibers and comparison study of their photocatalytic effects for efficient photocatalytic H₂ evolution. Nanoscale 8, 11034–43 (2016).
61. Wang, J. et al. A Synthesis of g-C₃N₄/TiO₂ with enhanced photocatalytic activity for H₂ evolution by a simple method. Int. J. Hydrogen Energy 39, 6354–6363 (2014).
62. Yan, H. et al. TiO₂–g-C₃N₄ composite materials for photocatalytic H₂ evolution under visible light irradiation. J. Alloys Compd. 509, L26–L29 (2011).
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Author Contributions
Y.W. wrote the main manuscript text. Y.W. and W.P. performed the experiments, analyzed the data. J.Y., J.T. and C.Y. contributed substantially to revisions. All authors reviewed the manuscript.

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