Decoration of TiO$_2$/g-C$_3$N$_4$ Z-scheme by carbon dots as a novel photocatalyst with improved visible-light photocatalytic performance for the degradation of enrofloxacin$^\dagger$

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A novel visible-light-driven carbon dot (CDs)/TiO$_2$/g-C$_3$N$_4$ photocatalyst was successfully synthesized by doping CDs in TiO$_2$ nanoparticles and the surface of g-C$_3$N$_4$ nanosheets via a facile hydrothermal process and was confirmed by characterization methods. UV-vis diffuse reflectance spectra (DRS) revealed that CDs/TiO$_2$/g-C$_3$N$_4$ showed obvious additional absorption in the 370–450 nm region. DMPO spin-trapping ESR spectra demonstrated the existence of O$_2^{-}$ and ·OH. The photocatalytic activity of the CDs (1.0 wt%)/TiO$_2$/g-C$_3$N$_4$ was remarkably enhanced as compared to that of the single components (TiO$_2$ and g-C$_3$N$_4$) and double component (TiO$_2$/g-C$_3$N$_4$) towards the degradation of enrofloxacin (ENX) under visible-light irradiation. About 91.6% of ENX was decomposed by CDs (1.0 wt%)/TiO$_2$/g-C$_3$N$_4$ in 1 h, which is nearly 3.95 times, 4.82 times, and 1.69 times that for TiO$_2$, g-C$_3$N$_4$, and TiO$_2$ (90.0 wt%)/g-C$_3$N$_4$, respectively. Scavenging experiments revealed that O$_2^{-}$ and ·OH played key roles during the photocatalytic degradation of ENX. This study provides a simple and convenient method to modify materials with enhanced photocatalytic performance, and the CDs/TiO$_2$/g-C$_3$N$_4$ catalyst is efficient, stable, and reusable for environmental practical applications.

**Introduction**

In the past decade, use of solar-energy-driven semiconductors as photocatalysts to remove aquatic contaminants became one of the most promising approaches.$^{1-3}$ As is known, semiconductor titanium dioxide (TiO$_2$) plays a significant role in the field of photocatalysis due to its nontoxicity, inexpensiveness, high activity, etc.$^4$ However, TiO$_2$ has a band gap of approximately 3.2 eV, which requires ultraviolet irradiation with wavelengths less than 385 nm for activation. That is, this excitation wavelength can activate electron transfer from the TiO$_2$ valence band (VB) to the conduction band (CB). Consequently, the photocatalyst utilizes less than 5% of the energy available in sunlight, which extremely restricts its application in practice.$^{5-7}$

More recently, a metal-free semiconductor, graphitic carbon nitride (g-C$_3$N$_4$), has been successfully synthesized through a facile thermal polycondensation method.$^{6,8}$ It is reported that g-C$_3$N$_4$ exhibits high photocatalytic activity due to its delocalized conjugated π structures, leading to relatively rapid photoinduced charge separation and slow charge recombination.$^{10}$ Furthermore, the low band gap (2.70 eV) capacitates g-C$_3$N$_4$ to efficiently harvest sunlight within the visible-light region. Although g-C$_3$N$_4$ possesses better photocatalytic activity as compared to TiO$_2$, it is far from enough to meet the demands in practical use. Various attempts including surface sensitization, chemical modification, and coupling with other semiconductor materials have been made to increase the photocatalytic efficiency of TiO$_2$ and g-C$_3$N$_4$.$^{9,11-12}$ Much interest has recently been concentrated on the fabrication of TiO$_2$/g-C$_3$N$_4$ heterostructures, which extend the absorption to the visible-light region.$^{13-15}$ However, to the best of our knowledge, h$^+$ from $E_{VB}$ of g-C$_3$N$_4$ is incapable of generating ·OH under visible light; in addition to the abovementioned shortcomings of TiO$_2$, the photocatalytic performance of TiO$_2$/g-C$_3$N$_4$ is still unsatisfactory.

Carbon dots (CDs), with an appealing group of zero-dimensional nanostructures, have attracted extensive research interest in the photocatalytic field due to their low-cost, aqueous dispersibility, chemical stability, unique photoinduced electron transfer, and electron reservoir properties.$^{16}$ Hence, photocatalysts with the incorporation of CDs have achieved excellent effects in the past few years. For example, Guo et al. synthesized a carbon quantum dot/carbon nitride hybrid photocatalyst to improve the photocatalytic degradation of methyl orange.$^{17}$ Wu et al. used CDs as a solid-state electron mediator for the
preparation of BiVO₄/CD/CdS Z-scheme photocatalyst, which showed excellent photocatalytic activity and stability for overall water splitting under visible light. Recently, Yu et al. reported a novel CD/ZnFe₂O₄ composite photocatalyst with enhanced photocatalytic performance for the removal of NOₓ, demonstrating that CDs acted as an electron reservoir and transporter as well as a powerful energy-transfer component. Due to the outstanding photocatalytic properties, in this study, CDs were first used for the decoration of the TiO₂/g-C₃N₄ Z-scheme through a facile hydrothermal-polymerized method.

Pharmaceuticals and personal care products (PPCPs), as emerging contaminants in recent years, have received unprecedented concern because of their potential hazard to the ecological environment and human health. According to the previous study, PPCPs have been confirmed to be ubiquitous throughout the world, and even worse, the drinking water treatment plants are also included. Consequently, it is extremely urgent to develop efficient methods for the elimination of PPCPs in aquatic environments.

In this study, the visible-light-driven CD/TiO₂/g-C₃N₄ ternary composites were first synthesized and used as photocatalysts for the degradation of PPCPs. Herein, enrofloxacin (ENX), largely used as veterinary bactericides, was taken as targeted PPCPs under the effects of the as-prepared materials. The structural and optical properties of the composites were investigated in detail by TEM, XRD, XPS, PL, etc. In addition, the influence of the TiO₂ : g-C₃N₄ molar ratio and doping amount of CDs on the photocatalytic performance of these composite photocatalysts was studied in an attempt to illuminate the potential mechanisms. The CD/TiO₂/g-C₃N₄ ternary composites were efficient towards the decomposition of refractory pollutants because of the synergic effect of accelerated charge separation and an extended light harvesting region. It was anticipated that this study could be instrumental in studying the removal of PPCPs and fabrication of novel ternary composite photocatalysts.

### Experimental

#### Materials

Enrofloxacin (ENX, 98% purity) was acquired from TCI Reagent Co. Ltd. (China). Degussa P25 (commercial TiO₂) was purchased from Sinopharm Chemical Reagent Co. Ltd. (Shanghai, China). Dicyandiamide was commercially purchased from Aladdin (China). Acetonitrile and methanol were obtained from CNW Technologies GmbH (Germany, HPLC-grade). Other reagents (e.g., citric acid, urea, acetic acid, sulfuric acid, sodium hydroxide, benzoquinone, isopropanol, potassium iodide, and sodium azide) were of analytical grade and used without further purification. Deionized (DI) water from a Milli-Q apparatus (Germany) was used in the whole experiment.

#### Synthetic procedures

**Preparation of the CDs.** C-Dots were prepared via the hydrothermal method and modified according to a previous report. Briefly, a weighed quantity (3.0 g) of citric acid and (3.0 g) urea was mixed with 10 mL DI water, which was placed into a Teflon autoclave for 5 h at 180 °C. After being cooled to room temperature, the obtained brown water solution was centrifuged at 10 000 rpm for 15 min to remove large particles. Finally, the remaining solution was transferred to the vacuum drier for 3 h at 80 °C to obtain a brownish black solid (C-dots).

**Preparation of g-C₃N₄.** g-C₃N₄ was prepared by directly heating dicyandiamide. In brief, 5.0 g of dicyandiamide was placed in an alumina crucible with a cover and then heated to 550 °C for 3 h at a heating rate of 2.8 °C min⁻¹. After being cooled down to room temperature, the obtained bulk g-C₃N₄ was milled into powders for further use.

**Preparation of the TiO₂/g-C₃N₄ binaries.** The TiO₂/g-C₃N₄ composites were prepared using a typical method. Briefly, an appropriate amount of TiO₂ was mixed with 20 mL of methanol in a beaker and sonicated until it was completely dispersed. After this, a calculated amount of g-C₃N₄ was mixed with the TiO₂ suspension and stirred in a fume hood for 15 h. Subsequently, the obtained materials were transferred to a muffle furnace and heated to 550 °C at a heating rate of 2.8 °C min⁻¹, and the temperature was maintained at 550 °C for 3 h. Through this method, a series of TiO₂ (wt%)/g-C₃N₄ photocatalysts (wt = 10.0, 30.0, 60.0, and 90.0) were synthesized and labelled as TCN-x (x = 10, 30, 60, and 90, respectively).

**Decoration of the CDs onto TiO₂/g-C₃N₄ composites.** The CDs/TiO₂/g-C₃N₄ ternary photocatalysts with different doping amounts of CDs (0.5, 1, 2, and 3 wt%) were synthesized as follows: first, the optimal mass ratio of the as-prepared TiO₂/g-C₃N₄ that exhibited the best performance for the photocatalytic degradation of ENX was placed in an alumina crucible with a cover. After this, a series of different mass ratios of CDs were added to the alumina crucible and heated to 550 °C for 3 h at a heating rate of 2.8 °C min⁻¹. After being cooled down to room temperature, the final CD/TiO₂/g-C₃N₄ composite photocatalysts were obtained and denoted as CTCN-0.5, CTCN-1, CTCN-2, and CTCN-3.

**Characterization**

Thermogravimetric (TG) analysis was conducted using STA-449C Jupiter (NETZSCH Corporation, Germany). The interlayer structure of the samples was characterized by a transmission electron microscope (TEM, JEM-2100HR). The crystal structure and phase purity of the obtained materials were tested by X-ray diffraction (XRD, BRUKER D8 ADVANCE) equipped with Cu Kα radiation (λ = 0.154178 nm). The UV-vis diffuse reflection spectra (DRS) of the as-prepared samples were analyzed via a UV2450 UV-vis spectrophotometer (Shimadzu) using BaSO₄ as a reflectance standard. X-ray photoelectron spectroscopy (XPS) was used to analyze the ionic characteristics, which was conducted using a Thermo VG ESCALAB 250 spectrometer with Al Kα radiation at 1486.6 eV. Fourier transform infrared (FT-IR) spectra were obtained using a Nicolet 6700 spectrophotometer (Thermo Fisher).

Photoluminescence (PL) spectra were obtained using a FluoroMax-4 fluorescence spectrophotometer (HORIBA Jobin Yvon).
Photocatalytic experiments

Photocatalytic activity tests of the photocatalysts were carried out in a XPA-7 rotary photochemical reactor (Fig. 1, Nanjing Xujiang machine plant). A 350 W xenon lamp with a 420 nm cut-off filter was used as the visible-light irradiation source. For each experiment, 50 mL of a 4 mg L$^{-1}$ ENX aqueous solution with 1.0 g L$^{-1}$ catalysts was introduced into a 50 mL quartz tube, which was adjusted to a neutral pH via 1% NaOH and H$_2$SO$_4$ solution. Before the light irradiation, the reaction solution was stirred in the dark for 30 min to reach adsorption equilibrium for ENX on the photocatalyst. The samples were filtered through 0.22 mm millipore filters to remove the nanophotocatalyst. ENX was analyzed by high-performance liquid chromatography (HPLC), and the chromatographic conditions of ENX are listed as follows: column: Zorbax Eclipse XDB-C18, 4.6 × 150 mm, 5 μm; temperature: 40 °C; mobile phase: methanol/Milli-Q water containing 0.2% formic acid (65:25 v/v); flow rates: 1 mL min$^{-1}$; injection volume: 20 μL; and detection wavelength: 278 nm.

The quenching experiments were employed to prove the existence of reactive species in the ENX photodegradation process. Herein, 0.0001 mol L$^{-1}$ benzoquinone (BQ, O$_2$$^{•-}$ scavenger), 0.01 mol L$^{-1}$ isopropanol (IPA, ·OH scavenger), 0.01 mol L$^{-1}$ potassium iodide (KI, ·OH and h$^+$ scavenger), and 0.075 mol L$^{-1}$ sodium azide (NaN$_3$, 1O$_2$ scavenger) were used as quenchers in the experiments.$^{30-33}$

Results and discussion

Characterization

To determine the real contents of g-C$_3$N$_4$ in TiO$_2$/g-C$_3$N$_4$, thermogravimetric analysis was performed from 30 °C to 800 °C at a heating rate of 10 °C min$^{-1}$ under air conditions. It can be observed from Fig. 2 that the burning of g-C$_3$N$_4$ occurs in the temperature range of 500–650 °C, and the pure TiO$_2$ presents almost no weight loss, which is in accordance with the previous report.$^{34}$ As shown in Fig. 2, the real contents of g-C$_3$N$_4$ can be easily calculated from the weight remained after combustion.

Fig. 3 shows the TEM and HRTEM images of the obtained materials. It can be clearly observed that the g-C$_3$N$_4$ displays a graphite-like layered structure with the agglomeration of TiO$_2$ on the surface (Fig. 3A and B). CDs, as the decoration of TiO$_2$/g-C$_3$N$_4$ heterojunction, are embedded in TiO$_2$ nanoparticles and deposited onto the surface of g-C$_3$N$_4$ nanosheets (Fig. 3C and D). The HRTEM images of the CDs/TiO$_2$/g-C$_3$N$_4$ samples (Fig. 3E) clearly reveal an interplanar spacing of 0.35 nm, which authenticates the existence of TiO$_2$ (anatase 101).$^{35}$ HRTEM images of CDs and the inset in Fig. 3F show CDs (average size: 8 nm).
with a lattice spacing of 0.22 nm corresponding to the (100) facet of graphite.\textsuperscript{36}

XRD was carried out to reveal the phase composition of the samples (Fig. 4). As shown in Fig. 4A, no impurity diffraction peaks are observed in the XRD patterns of TiO\textsubscript{2} and g-C\textsubscript{3}N\textsubscript{4}, which can be indexed to anatase (JCPDS 21-1272), rutile phases (JCPDS 21-1276) of TiO\textsubscript{2}, and hexagonal phase of g-C\textsubscript{3}N\textsubscript{4} (JCPDS 87-1526). TCN-90 samples show relatively strong peaks at the position of R(100), indicating the introduction of g-C\textsubscript{3}N\textsubscript{4}.\textsuperscript{37} After decoration of CDs, the characteristic diffraction peaks of TiO\textsubscript{2}/g-C\textsubscript{3}N\textsubscript{4} remain unchanged with the increasing mass contents of CDs. It can be concluded that CDs have a negligible effect on the crystalline structure of the TiO\textsubscript{2}/g-C\textsubscript{3}N\textsubscript{4} Z-scheme, decorated via a facile hydrothermal process. For others, no new diffraction peaks appear due to the low doping content of CDs.

To study the surface chemical composition and chemical status of CD/TiO\textsubscript{2}/g-C\textsubscript{3}N\textsubscript{4}, XPS analysis was carried out, and the results are shown in Fig. 5. It can be concluded that the ternary composites primarily consist of C, N, Ti, and O elements. High-resolution spectra of C 1s at 288.88 eV and 284.78 eV are shown in Fig. 5B. The former can be attributed to the N–C\textsubscript{N} bonds, whereas the latter is assigned to the C–C coordination including sp\textsuperscript{2}-hybridized C atoms present in g-C\textsubscript{3}N\textsubscript{4}.\textsuperscript{37} High-resolution spectra of C 1s of TCN-90 are provided as a contrast. It can be observed that C–C at 288.88 eV of CTCN-1 occupies relatively larger percentage than TCN-90 due to the doping of CDs. The N 1s high-resolution spectrum in Fig. 5C shows three individual peaks with the binding energies of 400.98 eV, 399.98 eV, and 398.78 eV, which can correspond to C–N–H, N–(C)\textsubscript{3}, and C–N–C, respectively.\textsuperscript{38} From Fig. 5D, due to the phenomena of spin–orbit separation, two peaks at the binding energies of 458.88 eV (Ti 2p\textsubscript{3/2}) and 464.58 eV (Ti 2p\textsubscript{1/2}) suggest the existence of Ti\textsuperscript{4+} in the composite.\textsuperscript{39} The O 1s high-resolution spectrum in Fig. 5E is observed at the binding energies of 532.08 eV and 530.48 eV, which are ascribed to the O–H bond and Ti–O bond, respectively.\textsuperscript{40,41} The former can be connected to the presence of a hydroxyl group on the surface of the CDs/TiO\textsubscript{2}/g-C\textsubscript{3}N\textsubscript{4} composite, which is in line with the FT-IR analysis in Fig. 6.

The FT-IR spectra of the TiO\textsubscript{2} microspheres, g-C\textsubscript{3}N\textsubscript{4}, TCN-90 composite, and CTCN-1 composite are shown in Fig. 6. For TiO\textsubscript{2}, a wide absorption band can be observed at 500–700 cm\textsuperscript{-1}, which is attributed to Ti–O stretching.\textsuperscript{42} The peaks at 1640 cm\textsuperscript{-1} and 3420 cm\textsuperscript{-1} are related to the bending vibration of O–H and N–H.\textsuperscript{43} For g-C\textsubscript{3}N\textsubscript{4}, the peak at 1630 cm\textsuperscript{-1} corresponds to the C–N heterocycle stretching vibration modes, whereas the peaks at 1450 cm\textsuperscript{-1} and 1250 cm\textsuperscript{-1} can be assigned to aromatic C–N stretching vibration modes.\textsuperscript{9} The peaks at 808 cm\textsuperscript{-1} and 3170 cm\textsuperscript{-1} are in accordance with the characteristic breathing mode of triazine units and N–H.\textsuperscript{15} A peak at 1409 cm\textsuperscript{-1} is observed in the FTIR spectra of both TCN-90 and CTCN-1, which can be interpreted as the introduction of g-C\textsubscript{3}N\textsubscript{4} and CDs.

The optical properties of the as-prepared samples were characterized by the DRS technique, and the results are shown in Fig. 7. TiO\textsubscript{2} and g-C\textsubscript{3}N\textsubscript{4} each revealed a basal absorption ~5 nm with a lattice spacing of 0.22 nm corresponding to the (100) facet of graphite.\textsuperscript{36}
band at ~390 nm and ~475 nm, respectively, which was consistent with previous reports.\textsuperscript{34,44,45} g-C\textsubscript{3}N\textsubscript{4}/TiO\textsubscript{2} possesses the absorption features of single TiO\textsubscript{2} and g-C\textsubscript{3}N\textsubscript{4}, which indicates the existence of g-C\textsubscript{3}N\textsubscript{4}. As expected, CD/TiO\textsubscript{2}/g-C\textsubscript{3}N\textsubscript{4} shows obvious additional absorption in the 370–450 nm region when doped with different amounts of CDs. The absorption edge of CTCN-1 ternary composite shows a remarkable red shift towards a higher wavelength at ~450 nm; this suggests that the ternary composite is able to utilize the visible-light region better. Herein, Kubelka–Munk transformation was used to estimate the band gap energies of the semiconductors according to eqn (1).

\[
(\alpha h\nu)^{1/2} = A(h\nu - E_g)
\]  

(1)

where \(\alpha\) is the absorption coefficient, \(h\nu\) is the light frequency, \(E_g\) is the band gap energy, and \(A\) is a constant.\textsuperscript{46} The potentials of VB and CB of the catalyst can be estimated according to eqn (2) and (3), respectively.

\[
E_{VB} = X - E_c + 0.5E_g
\]  

(2)

\[
E_{CB} = E_{VB} - E_g
\]  

(3)

where \(X\) is the electronegativity of the semiconductor (\(X\) values are 5.81 eV for TiO\textsubscript{2} and 4.72 eV for g-C\textsubscript{3}N\textsubscript{4}) and \(E_c\) is the energy of free electrons on the hydrogen scale (about 4.5 eV). The energy band structure parameters are listed in Table S1.\textsuperscript{†}

PL spectra were obtained (Fig. 7B) to detect the recombination of electron–hole pairs. Generally, lower PL intensity corresponds to lower recombination of electron–hole pairs, contributing to higher photocatalytic activity.\textsuperscript{47} The g-C\textsubscript{3}N\textsubscript{4} has an intense emission peak at around 460 nm, which can be attributed to the fast recombination of the photoinduced electron–hole pairs of g-C\textsubscript{3}N\textsubscript{4}. The intensities of the peaks for CTCN-0.5, CTCN-1, CTCN-2, and CTCN-3 are much lower than those for TCN-90, revealing that CDs conduce to slow the recombination rate of electron–hole pairs. Further, the emission peak of CDs/TiO\textsubscript{2}/g-C\textsubscript{3}N\textsubscript{4} shows an obvious red-shift from 455 nm to 470 nm after doping of CDs, which demonstrates the existence of CDs from another perspective. The PL results confirm the importance of CDs/TiO\textsubscript{2}/g-C\textsubscript{3}N\textsubscript{4} in blocking the recombination of electrons and holes.

PL spectra of CDs are shown in Fig. 8, which demonstrate superior up-converted photoluminescence property of CDs. When CDs were excited by long-wavelength light (from 600 nm to 900 nm), the up-converted emissions were significantly located in the range from 350 nm to 600 nm. These results imply that CDs might enhance the photocatalytic efficiency \textit{via} the conversion of near-infrared wavelength to visible light, which markedly promotes the light utilization of the photocatalysts.

**Photocatalytic performance of composites**

Fig. 9 displays the photocatalytic degradation of ENX over various photocatalysts. As can be seen, the self-photolysis of ENX could be neglected in the blank experiment. For TCN-90 (see Fig. 9A), the concentration of ENX is reduced to 45.95% in 1 h. This means TCN-90 performs better than other doping ratios of g-C\textsubscript{3}N\textsubscript{4}, which confirms that the small doping ratio of g-C\textsubscript{3}N\textsubscript{4} is in favour of ENX degradation. After doping of CDs (1 wt%), enhanced photocatalytic activity was exhibited by both TiO\textsubscript{2} and g-C\textsubscript{3}N\textsubscript{4}, which could be attributed to the up-converted PL property. In Fig. 9B, we drew a comparison between TiO\textsubscript{2}, TiO\textsubscript{2}/CDs, g-C\textsubscript{3}N\textsubscript{4}, g-C\textsubscript{3}N\textsubscript{4}/CDs, TCN-90, CTCN-0.5, CTCN-1, CTCN-2, and CTCN-3 in the photocatalytic degradation of ENX. CTCN-1 was found to be the most optimal photocatalyst among all the photocatalysts as it reduced ENX to 8.34% in 1 h.
which corresponds to the DRS results in Fig. 7A. Compared with the single-component (TiO$_2$ and g-C$_3$N$_4$) samples and two-component samples (TiO$_2$/CDs, g-C$_3$N$_4$/CDs, TCN-90), CD/TiO$_2$/g-C$_3$N$_4$ exhibits remarkably enhanced photocatalytic activity, owing to the introduction of CDs as an electron reservoir and transporter. However, further CD loading (>1.0 wt%) leads to decreased degradation rates, which can be attributed to the inner filter effect of CDs via scrambling for the absorption of photons and reduction of the formation of reactive species.

Reusability is a critical factor for practical utilization. To investigate the stability of CD/TiO$_2$/g-C$_3$N$_4$, five photocatalytic experiment cycle runs were completed, and the results are shown in Fig. 10. As a result, 88.7% of the ENX was decomposed in 60 min in the fifth cycle, which was slightly reduced as compared to that in the first round (91.6%). Thus, the CTCN-1 can be identified as a stable photocatalyst.

To study the generation of radicals in the photocatalytic system under visible-light irradiation ($\lambda > 420$ nm), the ESR spin-trap with DMPO technique was used. As shown in Fig. 11, the signals of O$_2$$^-$ and ·OH increase with time elapsing in 10 minutes, whereas no signals appear in the dark. The weak signals of O$_2$$^-$ and ·OH radicals in TiO$_2$ can be attributed to fast recombination of photogenerated electron–hole pairs. Because the VB position of g-C$_3$N$_4$ (+1.36 eV) is more negative than the standard redox potential of ·OH/OH$^-$ (+1.99 eV) and ·OH/H$_2$O (+2.73 eV), h$^+$ of g-C$_3$N$_4$ fails to provide ·OH by oxidizing OH$^-$ or H$_2$O. The signals of ·OH radicals appear in g-C$_3$N$_4$ due to the reaction of O$_2$$^-$ and h$^+$. Compared with TCN-90, the signals of O$_2$$^-$ radicals are greatly enhanced for CTCN-1, indicating that CDs play a significant role as electron mediator to improve the photocatalytic activity of CD/TiO$_2$/g-C$_3$N$_4$. Simultaneously, the signals of O$_2$$^-$ radicals are also improved, demonstrating the increased photocatalytic activity of CD-modified TiO$_2$/g-C$_3$N$_4$.
Fig. 12 shows the kinetic constants of TiO₂, g-C₃N₄, TCN-90, and CTCN-1 in the presence of different quenchers. For the addition of isopropanol (IPA, -OH scavenger) and KI (-OH and h⁺ scavenger), a slight decrease can be observed in the degradation of ENX by CTCN-1, indicating the minor role of h⁺. For the TCN-90 composite, similar results were obtained, whereas a slight decrease of $k$ was observed after the addition of NaN₃ (¹O₂ scavenger) for the TCN-90 and CTCN-1 composites. Although benzoquinone (BQ, O₂⁻⁻ scavenger) has a slight effect in the presence of TCN-90, it plays a quite important role in CTCN-1 probably because CDs can act as electron mediators and generate more O₂⁻⁻.

Based on the abovementioned results and the previous study, the potential photocatalytic mechanisms of the as-prepared photocatalysts with different quenchers are proposed in Fig. 13. For the conventional TiO₂/g-C₃N₄ heterojunctions, electron–hole pairs are formed at the $E_{CB}$ and $E_{VB}$ of TiO₂ and g-C₃N₄ under visible light. Subsequently, h⁺ generates at the $E_{VB}$ of TiO₂ and transfers to the $E_{CB}$ of g-C₃N₄, and the e⁻ from $E_{CB}$ of g-C₃N₄ can transfer to the $E_{VB}$ of TiO₂. However, the h⁺ from $E_{VB}$ of g-C₃N₄ cannot oxidize OH⁻ or H₂O to generate ·OH because of the negative valence band position of g-C₃N₄ (+1.51 eV), which is less than the standard redox potential of ·OH/OH⁻ (+1.99 eV) and ·OH/H₂O (+2.73 eV). The e⁻ from TiO₂ may be captured by molecular oxygen to generate O₂⁻⁻ radicals, which can directly attack pollutants. If TiO₂/g-C₃N₄ follows this mechanism, h⁺ should play a major role in the photocatalytic system, whereas IPA has minor effects. However, the quenching experiments and ESR spectra results have contradicted this hypothesis. Moreover, the results of the ¹O₂ quenching experiment are inconsistent with this mechanism. Thus, the TiO₂/g-C₃N₄ photocatalyst follows the direct Z-scheme type mechanism.

For the TiO₂/g-C₃N₄ Z-scheme with the introduction of CDs, the near-infrared light wavelength could be converted to visible light due to the up-converted PL properties of CDs⁵³. Herein, the presence of CDs can facilitate a Z-scheme transfer mechanism, which differs from the general heterostructured mechanism. The general heterostructured mechanism can hardly interpret the formation of ·OH, and the transfer speed of electrons and holes might be very slow.⁵⁴ The active species, h⁺, O₂⁻⁻, ¹O₂, and ·OH can effectively degrade ENX into small molecule compounds or directly into final products such as CO₂ and H₂O.

Conclusions

In this study, CDs/TiO₂@g-C₃N₄ was synthesized via a facile hydrothermal method, which exhibited high photocatalytic performance towards the degradation of ENX under visible-light irradiation. The results of the characterization verified the formation of CD/TiO₂@g-C₃N₄, in which the CDs were embedded in TiO₂ nanoparticles and deposited onto the surface of the g-C₃N₄ nanosheets. The enhanced photocatalytic performance of CDs/TiO₂@g-C₃N₄ might be attributed to the unique up-converted PL property, efficient charge separation, and efficient electron transportation of CDs. The synergistic effects could be concluded as follows: (1) the extension of visible-light response scope; (2) efficient photogenerated electron–hole pair separation; and (3) retardation of charge recombination. The active species h⁺, O₂⁻⁻, ¹O₂, and ·OH were produced in the Z-scheme system. The present study confirms that the CD/TiO₂@g-C₃N₄ catalyst is efficient, stable, and reusable for environmental applications.

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