


 Cite this: *RSC Adv.*, 2024, 14, 28904

# Synergistic self-driven and heterogeneous effect of a biomass-derived urchin-like $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ Janus micromotor catalyst for efficient degradation of carbamazepine†

 Jie Yang,<sup>a</sup> Wenning Yang,<sup>‡b</sup> Chao Zhang,<sup>ID \*c</sup> Jian Gong,<sup>a</sup> Ming Xu,<sup>d</sup> Jia Li<sup>ID \*e</sup> and Chengzhang Liu<sup>e</sup>

It is well known that obtaining efficient carbamazepine degradation materials or rapid carbamazepine-removal methods is still a challenge in the field of environmental remediation. Hence, the present study aimed to concurrently address these issues by combining a self-driven, heterostructured and low-cost biomass-templated urchin-like Janus micromotor catalyst for highly efficient carbamazepine degradation. The catalyst could autonomously move in a circle-like motion pattern *via*  $\text{O}_2$  bubbles generated from the  $\text{Mn}_3\text{O}_4$ -catalyzed decomposition of  $\text{H}_2\text{O}_2$  with a velocity of  $223.5 \pm 7.0 \mu\text{m s}^{-1}$  in 1%  $\text{H}_2\text{O}_2$ . Benefiting from the well-structured heterojunction at the interface of  $\text{C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$ , carbamazepine (CBZ) was degraded by 61% in 100 min under sunlight irradiation. In addition, density functional theory calculation results proved that the formation of the heterojunction structure promoted the generation of photo-generated carriers. Thus, the presented method provides a promising pathway for the rational construction and preparation of movable catalysts for the efficient removal of organic pollutants from wastewater.

 Received 10th July 2024  
 Accepted 9th August 2024

DOI: 10.1039/d4ra04980b

[rsc.li/rsc-advances](https://rsc.li/rsc-advances)

## Introduction

Antibiotic degradation remains a great challenge to green, low-cost, and feasible approaches for the degradation of medicines and environmental protection. As one of the broad-spectrum antibiotics, carbamazepine (CBZ) has attracted much attention in the treatment of various diseases such as trigeminal neuralgia, epilepsy, and mental illness.<sup>1</sup> According to statistics, about 28% of CBZ is discharged into the water environment due to incomplete human metabolism and medical sewage discharge.<sup>2</sup> To worsen the situation, CBZ is a kind of extremely durable and stable organic environment pollutant and cannot be effectively removed *via* traditional wastewater treatment

technologies.<sup>3,4</sup> Therefore, the development of efficient treatment technology for the degradation of CBZ in sewage is urgent. Compared with the removal of CBZ using conventional technologies, such as biological treatment, adsorption, and other methods,<sup>5–7</sup> the advanced oxidation process (AOP) is regarded as a promising method for the effective removal and mineralization of CBZ.<sup>8–10</sup> Specifically, AOPs based on heterojunction photocatalysis have received more attention owing to their economical preparation, convenient and simple synthesis, and efficient degradation.<sup>11–13</sup> Heterojunction structures have excellent electronic structural characteristics and can exhibit higher catalytic efficiency than traditional catalysts in catalytic reactions. Moreover, the large specific surface area and nanoparticle structure of heterojunction catalysts make their catalytic performance more stable than traditional catalysts. Further, their interactions and interface effects during the reaction process help prevent catalyst failure and deactivation.<sup>14,15</sup> Among the many heterojunction photocatalysts,  $\text{Mn}_x\text{O}_y/\text{C}_3\text{N}_4$ , especially two-dimensional (2D) nanostructured  $\text{Mn}_3\text{O}_4$  heterojunctions, has attracted great interest because of its delocalized conjugative structures, efficient charge separation, outstanding chemical stability, and low cost.<sup>16–18</sup> It has also been reported that  $\text{Mn}_x\text{O}_y/\text{C}_3\text{N}_4$  heterojunctions have good optical absorption properties and can be used for direct charge-transfer collection and are thus applied in a wide range of fields.<sup>19–21</sup> Shi successfully prepared a  $g\text{-C}_3\text{N}_4/\alpha\text{-MnO}_2$  Z-scheme

<sup>a</sup>Department of Pharmaceutical and Bioengineering, Zibo Vocational Institute, Zibo, 255000, P. R. China

<sup>b</sup>Shandong Provincial Key Laboratory of Chemical Energy Storage and Novel Cell Technology, School of Chemistry and Chemical Engineering, Liaocheng University, Liaocheng, 252000, P. R. China

<sup>c</sup>School of Artificial Intelligence and Big Data, Zibo Vocational Institute, Zibo, 255000, P. R. China. E-mail: 652143914@qq.com

<sup>d</sup>State Key Laboratory of Chemical Resource Engineering, Beijing University of Chemical Technology, Beijing, 100029, P. R. China

<sup>e</sup>School of Material Science and Engineering, University of Jinan, Jinan, 250022, China. E-mail: mse\_lij@ujn.edu.cn

 † Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ra04980b>

‡ The author contribute equally: Wenning Yang.



heterojunction, which showed excellent visible-light photocatalytic performance, superior to its pure constituent parts.<sup>22</sup> Moreover, Zhang reported a porous MnO<sub>2</sub>/Mn-modified alkalized g-C<sub>3</sub>N<sub>4</sub> catalyst, which exhibited high catalytic activity and 96.7% tetracycline removal.<sup>23</sup> Besides, Chen synthesized nanodot–nanosheet (Mn<sub>3</sub>O<sub>4</sub>/g-C<sub>3</sub>N<sub>4</sub>) composites, which showed the best performance in persulfuric salt (PMS) activation for the removal of 4-chlorophenol (4-CP).<sup>24</sup> However, it is still a challenge to prepare efficient Mn<sub>x</sub>O<sub>y</sub>/C<sub>3</sub>N<sub>4</sub> heterojunction composites to achieve efficient photocatalytic effects and describe the detailed mechanism of action of heterojunction structures.

At the same time, the satisfactory degradation performance of a catalyst depends not only on its inherent properties but also on the chance of it contacting with the target pollutants. In order to achieve more effective contact, making the catalyst disarray move is a very useful method to ensure achieving effective contact.<sup>25–27</sup> In this regard, micro-/nanomotors, a kind of self-propelled device at the micrometre or nanometre level, that can convert different forms of energy into kinetic energy to perform special tasks under liquid conditions, have considerable application prospects in the fields of biomedicine, sensing, and environmental remediation. Compared with the traditional static catalytic degradation, micro-/nanomotors can shorten the reaction time and reaction site by increasing the contact frequency and changing the displacement of molecules.<sup>28–31</sup> For instance, Zhu introduced a bioinspired flower-shaped hierarchical Pt-free micromotor with a maximal adsorption capacity of 129.51 mg g<sup>-1</sup> through the assistance of movement.<sup>32</sup> Song summarized methods utilizing micro-/nanomotors for improving biosensing, such as the sensitivity, selectivity, detection time, biocompatibility, simplified system operation, and environmental availability.<sup>33</sup> Our group recently constructed a novel glucose-driven catalytic nanomotor with robust dual enzyme-like activities for the sensitive colorimetric sensing of glutathione (GSH) in wastewater.<sup>34</sup> Therefore, it has been confirmed that self-propelled motion and bubble formation together lead to more effective fluid mixing, thereby increasing the degradation efficiency of low-concentration pollutants and

compensating for the low diffusion rate of heterogeneous sensors and catalysts.

Inspired by the previous biotemplate self-driven catalytic micro-/nanomotors and Mn<sub>x</sub>O<sub>y</sub>/C<sub>3</sub>N<sub>4</sub> heterojunction composites, herein we synthesized a novel Mn<sub>3</sub>O<sub>4</sub>/C<sub>3</sub>N<sub>4</sub> Janus micromotor catalyst (Mn<sub>3</sub>O<sub>4</sub>/C<sub>3</sub>N<sub>4</sub>-JMC) with a high specific surface area and 3D hierarchical structure, and applied this to the dynamic photocatalytic degradation of carbamazepine in the presence of hydrogen peroxide (Scheme 1). Sea urchin-shaped sunflower pollen with a grain size of 30 μm was soaked in hot phosphoric acid for several hours, and then washed several times with DI water, acetone, hydrochloric acid, and ethanol in sequence to remove sticky organic matter on the surface of the pollen. Next, a certain amount of melamine was thoroughly mixed and ground with the cleaned pollen. Then the mixture was calcined at 650 °C for 2 h under the protection of Ar gas in a tube furnace to obtain C<sub>3</sub>N<sub>4</sub>/C samples. Next, a certain concentration solution of PMMA was coated on one side of a glass slide. Then a small pinch of the C<sub>3</sub>N<sub>4</sub>/C samples was slowly poured onto the surface of the glass slide until PMMA evaporated to form a uniform thin film layer. Then, alkaline KMnO<sub>4</sub> was dropped onto the slide glass. After that, the resulting hybrid composites were exposed to 100 W incandescent light for 12 h. Finally, Mn<sub>3</sub>O<sub>4</sub>/C<sub>3</sub>N<sub>4</sub>-JMC was obtained after a simple cleaning and drying process. In the presence of H<sub>2</sub>O<sub>2</sub>, on the one hand, the semi-coated Mn<sub>3</sub>O<sub>4</sub> on the surface of the catalyst could decompose H<sub>2</sub>O<sub>2</sub> to produce oxygen gas bubbles for propulsion. Meanwhile, heterojunction g-C<sub>3</sub>N<sub>4</sub>/Mn<sub>3</sub>O<sub>4</sub> was able to degrade harmful carbamazepine in sewage under light irradiation. Benefiting from the synergistic effect of autonomous motion with the high efficiency catalysis of the Mn<sub>3</sub>O<sub>4</sub>/C<sub>3</sub>N<sub>4</sub> heterojunction, the catalyst could highly efficiently degrade CBZ in water. Moreover, the catalyst system also demonstrated high reusability and stability. Therefore a new vision has been developed that has application potential in the field of environmental treatment and remediation.

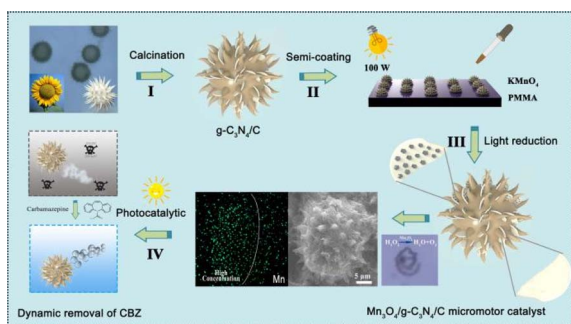
## Experimental

### Chemicals and materials

Sunflower pollen was obtained from Taian Jinzhong Sanitary Material Co., Ltd. Melamine was obtained from Sinopharm Chemical Reagent Co., Ltd. Sodium dodecyl sulfonate (C<sub>12</sub>H<sub>25</sub>SO<sub>3</sub>Na, SDS), isopropanol (IPA), benzoquinone (BQ), sodium oxalate (OA), and phosphoric acid (H<sub>3</sub>PO<sub>4</sub>) were obtained from Tianjin Chemical Reagent Co., Ltd. Potassium permanganate (KMnO<sub>4</sub>), ethanol, acetone, hydrochloric acid (HCl), sodium hydroxide (NaOH), and hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) were purchased from Aladdin. Carbamazepine (CBZ), polymethyl methacrylate (PMMA), and ethyl acetate were purchased from Sigma-Aldrich. All the chemicals were of analytical-grade purity and were used as received without further purification.

### Characterization

X-Ray diffraction patterns were obtained using an X-ray diffractometer (Bruker D8-Advance, Germany, Cu X-ray



**Scheme 1** Schematic of the synthesis and application of a Mn<sub>3</sub>O<sub>4</sub>/C<sub>3</sub>N<sub>4</sub> micromotor catalyst (Mn<sub>3</sub>O<sub>4</sub>/C<sub>3</sub>N<sub>4</sub>-JMC). (I) Growth of C<sub>3</sub>N<sub>4</sub> nanosheets on sunflower pollens *via* calcination in an inert atmosphere. (II) Application of the organic solvent impregnation semi-coating method for catalyst preparation; (III) formation of Mn<sub>3</sub>O<sub>4</sub>/C<sub>3</sub>N<sub>4</sub>-JMC *via* a light-driven reduction process. (IV) Degradation of carbamazepine (CBZ) *via* the synergy of photocatalysis with autonomous motion.



sources, 40 kV), in the  $2\theta$  range from 5–80° and scanning speed of  $2^\circ \text{ min}^{-1}$ . The morphologies were analyzed by transmission electron microscopy (TEM) on a Tecnai F20 instrument at an accelerating voltage of 4 kV for the electron beam and by field emission scanning electron microscopy (FESEM) on a Quanta 400F system fitted with an energy-dispersive X-ray spectroscopy (EDX) unit. The concentration of the suspended sample was  $0.2 \text{ mmol L}^{-1}$ . Fourier-transform infrared (FT-IR) spectra of the obtained samples were obtained using a Thomas Nicolet FT-IR spectrometer. The analyses were performed through KBr pellets and the wavenumber scanning range was  $4000 \text{ to } 400 \text{ cm}^{-1}$ , while the amount of samples was 20 mg, the spectral resolution was better than  $0.4 \text{ cm}^{-1}$ , and the signal to noise ratio was 60 000 : 1. Simultaneous thermogravimetry and derivative thermogravimetry analyses (TGA/DTG) were carried out between 30 °C and 850 °C at a rate of  $10 \text{ }^\circ\text{C min}^{-1}$  on a TA instrument (Netzsch Sta 449) under a flow of air. The surface elemental composition of the samples was determined by X-ray photoelectron spectroscopy (XPS). Videos of the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$  were captured by an optical microscope (Microscope N-300 M), coupled with a digital camera (Tucsen EC300) using the TSview software to analyze the speeds of the Janus micromotor catalyst. The photoelectric current was analyzed using an electrochemical workstation (CHI-600e) coupled with a standard three-electrode, where the sample-coated ITO glass served as the working electrode, Pt wire as the counter electrode, Ag/AgCl as the reference electrode, 0.1 M  $\text{Na}_2\text{SO}_4$  solution as the electrolyte, and a 300 W Xe lamp as the radiation source.

### Synthesis of urchin-like $\text{C}_3\text{N}_4/\text{C}$

In the synthesis, the sunflower pollen grains were suspended in phosphoric acid and mixed to form a homogeneous suspension, which was further heated to 70 °C and stirred gently for 5 h. Then, the pollen grains were collected and extensively washed using DI water, acetone, hydrochloric acid, and ethanol in sequence. Finally, the wet pollen grains were dried at 60 °C for future characterization and experiments. Next, 0.5 g of the extracted samples and 3.0 g melamine were placed in a mortar and thoroughly ground for 5 min. Then the mixed samples were amplified in a tube furnace, under Ar protection at a heating rate of  $2 \text{ }^\circ\text{C min}^{-1}$  for 4 h reaction at 650 °C. The obtained sample was designated as  $\text{C}_3\text{N}_4/\text{C}$ .

### Synthesis of the urchin-like $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ Janus micromotor catalyst ( $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$ )

The  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$  was prepared by a facile semi-coating method, as reported by Tan's research group.<sup>35</sup> First, 50 mL of 10% w/w solution of PMMA was coated on one side of a  $2.5 \text{ cm} \times 7.6 \text{ cm} \times 0.1 \text{ cm}$  glass slide. Next, 0.05 g of the  $\text{C}_3\text{N}_4/\text{C}$  samples was slowly poured onto the surface of the PMMA-coated glass slide until the particles layer evenly covered the PMMA surface. Then, 0.1 mL of 0.45 M alkaline  $\text{KMnO}_4$  ( $\text{KMnO}_4 : \text{NaOH} \frac{1}{4} : 1 : 1$  in molar ratio) was dropped onto the glass slide containing the  $\text{C}_3\text{N}_4/\text{C}$ . After that, the resulting hybrid composites were exposed to 100 W incandescent light for 12 h. Then the samples were cleaned with ethyl acetate and DI water

followed by drying at 60 °C for 8 h, and the obtained products were denoted as  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$ . Also, a comparative sample,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  without carbonized pollen, was prepared using an identical process.

### Removal of CBZ by $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$

The catalytic property of the as-synthesized products was reflected by CBZ removal under 300 W Xe lamp irradiation without sodium dodecyl sulfonate (SDS). In each experiment, 0.05 g of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$  samples was dispersed in a breaker containing 50 mL CBZ solution ( $50 \text{ mg L}^{-1}$ ). This was then stirred in the dark for 30 min to achieve adsorption–desorption equilibrium and exposed to a xenon lamp at  $30 \pm 2 \text{ }^\circ\text{C}$ . Then a quantity of  $\text{H}_2\text{O}_2$  (the total mass fraction, 0.5%) was added to the mixed solution. About 3 mL of the aliquot solution was withdrawn at different time intervals (0, 1, 3, 5, 10, 15, 20, 40, 50, 60, 100, 150, 200 min, *etc.*) from the reaction mixture, and was subjected to UV-vis spectrophotometry analysis. The absorbance at 284 nm was tested and recorded. The degradation efficiency of CBZ was calculated using the equation.

$$D(\%) = (C_0 - C_t)/C_0 \times 100\% \quad (1)$$

where  $D$  is the degradation efficiency of CBZ,  $C_0$  is the initial concentration, and  $C_t$  is the concentration after  $t$  min reaction of the solution.

### Stability and reusability of $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$ micromotor catalyst

The operational reusability of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$  was evaluated in a series of repeated batch experiments, and the activity retention of the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$  was tested, as described in relation to the activity assays. After each batch,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4\text{-JMC}$  was collected and washed with deionized water three times to remove any residual substrate and then reintroduced into the fresh reaction medium.

### DFT calculations

Density functional theory (DFT) calculations were performed using the Vienna *Ab initio* Simulation Package (VASP) with the projector augmented wave pseudopotentials.<sup>36</sup> The Perdew–Burke–Ernzerhof (PBE) exchange–correlation functional within the generalized gradient approximation (GGA) was chosen in consideration of a balance between the accuracy and computational cost.<sup>37</sup> The plane wave energy cut-off was 400 eV for the slabs. These periodic slabs were separated by 20 Å vacuum space along the  $z$  direction to isolate interactions between replicas. The electronic and force convergence standards were respectively set to  $10^{-8} \text{ eV}$  and  $0.02 \text{ eV } \text{Å}^{-1}$ . The Brillouin zone was sampled on a  $5 \times 5 \times 5$  Monkhorst–Pack  $k$ -point grid for  $\text{Mn}_3\text{O}_4$  bulk and  $4 \times 4 \times 1$  for the heterostructure and  $g\text{-C}_3\text{N}_4$ .<sup>38</sup> The charge density difference was obtained by subtraction of that for the total heterostructure of  $g\text{-C}_3\text{N}_4/\text{Mn}_3\text{O}_4$  minus that for  $g\text{-C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$ . The latter two models' schemes did not undergo optimization.





## Results and discussion

Characterization of the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC micromotor catalyst

Fig. 1a depicts the XRD patterns of the carbonized sunflower pollen and the final product  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC. The XRD pattern of a control sample  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  is also provided for comparison. As shown, the carbonized sunflower pollen exhibited a high diffraction baseline and two broad diffraction peaks around  $26^\circ$  and  $43^\circ$ , ascribed to the (002) and (100) planes, indicating the presence of amorphous organics. Nevertheless, the presence of a broad diffraction peak at  $13^\circ$  marked by a symbol of spades could be ascribed to the graphite phase. In the XRD patterns of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC, the corresponding (100) facet at  $12.81^\circ$  indicated the periodic arrangement in triazine of  $\text{C}_3\text{N}_4$ ; while the corresponding (002) crystal plane at  $27.91^\circ$  indicated the accumulation by the conjugated direction system of carbon nitride. Besides, the diffraction peaks at  $17.84^\circ$ ,  $25.87^\circ$ ,  $30.61^\circ$ , and  $36.05^\circ$  could be ascribed to the (020), (022), (110), and (112) planes of  $\text{Mn}_3\text{O}_4$  (JCPDS 75-0765). At the same time, a low diffraction intensity was observed for the catalyst due to the large amount of amorphous carbon in the samples. Fig. 1b shows the FT-IR spectra of carbonized sunflower pollen,  $\text{C}_3\text{N}_4$ ,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ ,

and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC. The peaks at approximately  $811\text{ cm}^{-1}$  were mainly caused by the stretching vibration of the triazine ring in the graphite carbon nitride,<sup>39</sup> while many peaks in the range of  $1200\text{--}1700\text{ cm}^{-1}$  were attributed to the typical breathing and stretching vibration modes of the heptazine heterocyclic ring,<sup>40</sup> and the peak at  $3140\text{ cm}^{-1}$  corresponded to the  $\text{--OH}$  bond of water. Besides, the spectrum of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC still appeared to be the same as the spectrum of  $\text{C}_3\text{N}_4$ , indicating the phase stability of  $\text{C}_3\text{N}_4$  in  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC.

The nitrogen adsorption–desorption isotherms and pore-size distributions of  $\text{C}_3\text{N}_4$ ,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ , and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC are shown in Fig. 1c. The isotherms of all these samples belonged to type IV isotherms with an H3 hysteresis loop, indicating the presence of mesopores in the obtained samples.<sup>41</sup> Through calculation and analysis, the specific surface areas of  $\text{C}_3\text{N}_4$ ,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ , and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC were  $88.39$ ,  $89.66$ , and  $81.35\text{ m}^2\text{ g}^{-1}$ , respectively (Table S1†). In addition, the average pore sizes of  $\text{C}_3\text{N}_4$ ,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ , and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC were  $7.1$ ,  $6.3$ , and  $9.2\text{ nm}$ , respectively. As shown in the inset of Fig. 1c, the distribution of pores was mainly concentrated in the range of  $2\text{--}50\text{ nm}$  and multi-peaks were observed for all three samples.

The elements C, N, Mn, and O could be clearly detected in the XPS spectra of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC, as shown in Fig. 1d. The carbon peak that appeared at  $284.8\text{ eV}$  was due to the hydrocarbon originating from the XPS instrument itself used as the standard. The C 1s spectrum (Fig. 1e) could be deconvoluted into three peaks centred at  $284.7$ ,  $288.2$ , and  $293.4\text{ eV}$ , corresponding to  $\text{C}=\text{C}/\text{C}-\text{C}$ ,  $\text{C}-\text{N}$  and  $\text{C}-\text{F}$ , respectively. The  $\text{C}=\text{C}$  and  $\text{C}-\text{N}$  were derived from the  $\text{C}_3\text{N}_4$  in  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC, while  $\text{C}-\text{F}$  was derived from the carbonized pollen. The three peaks of N 1s at  $398.8$ ,  $400.5$ , and  $404.6\text{ eV}$  (Fig. 1f) could be attributed to  $\text{C}=\text{N}-\text{C}$ ,  $\text{C}-\text{N}-\text{H}$ , and  $\text{O}-\text{N}$ . The Mn 2p peaks (Fig. 1g) located at  $642.2$  and  $653.3\text{ eV}$  were fitted to Mn  $2p_{3/2}$  Mn  $2p_{1/2}$ , respectively. Moreover, the O 1s spectra showed three components (Fig. 1h), with the  $\text{C}-\text{O}-\text{Mn}$  bond ( $532.2\text{ eV}$ ) for  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC revealing that some of the O of  $\text{Mn}_3\text{O}_4$  was involved in bonding with the C of  $\text{C}_3\text{N}_4$ .

Fig. S1† shows the DTA curves of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC, pure  $\text{C}_3\text{N}_4$ , and  $\text{Mn}_3\text{O}_4$ . The small and linear loss of weight below  $100^\circ\text{C}$  was due to the evaporation of water. Compared to the pure  $\text{C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$  samples, the weight of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC showed a slight decrease from  $120^\circ\text{C}$  to  $500^\circ\text{C}$ , while the DTA curve displayed a broad exothermic peak, due to the degradation of carbonized sunflower pollen (C) and the partial decomposition of  $\text{C}_3\text{N}_4$  in the composites. Moreover, the DTA curve of the pure  $\text{C}_3\text{N}_4$  samples also indicated that  $\text{C}_3\text{N}_4$  was partially pyrolyzed in this temperature range. The next mass loss stage of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC between  $500^\circ\text{C}$  and  $700^\circ\text{C}$  could be ascribed to the decomposition of  $\text{C}_3\text{N}_4$ , along with an exothermic reaction, as shown in the DTA curve. The same result was also reflected in the DTA curve of pure  $\text{C}_3\text{N}_4$ . Compared with the relevant literature,<sup>42</sup> the exothermic peak of  $\text{C}_3\text{N}_4$  was shifted towards a lower temperature. It can be inferred from this that the formation of the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  heterojunction affected the temperature resistance of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC.<sup>43</sup>

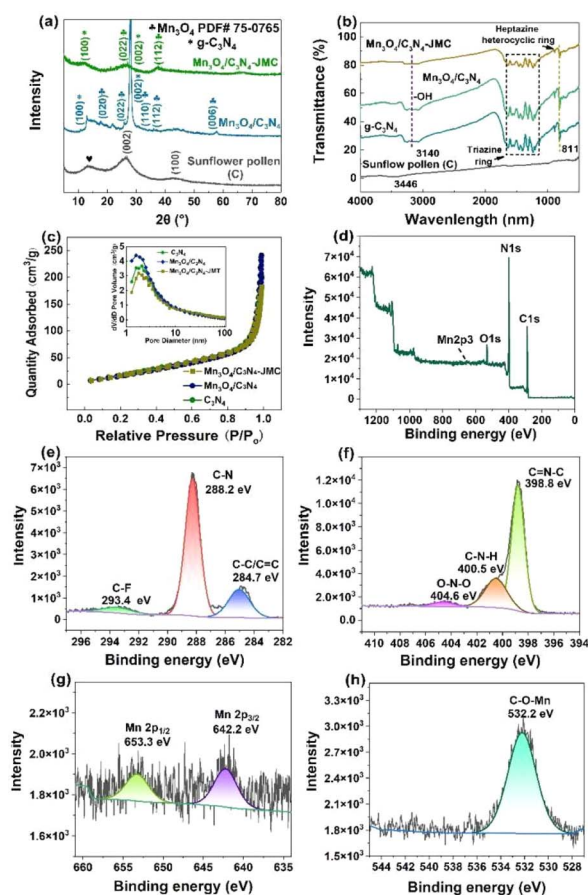


Fig. 1 XRD patterns (a) and IR spectra (b) of the obtained samples;  $\text{N}_2$  adsorption–desorption isotherms and pore-size distributions of the obtained samples (c). Full-XPS spectra (d) and spectra of C 1s (e), N 1s (f), Mn 2p (g), and O 1s (h) for  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC.



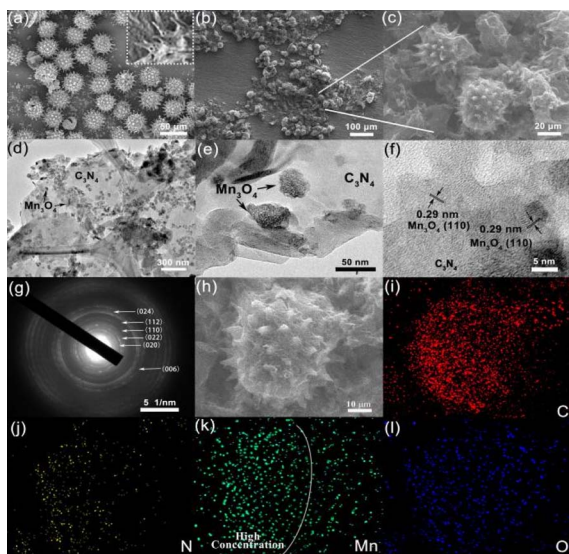


Fig. 2 FESEM images of sunflower pollens (a) and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC (b and c). TEM images (d–f), selected area electron diffraction (g) of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC. Energy-dispersive X-ray mappings of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC (h–l).

The FESEM and TEM images further revealed the micro- and nanostructure of the obtained samples. As shown in Fig. 2a, the sunflower pollen had a hollow and spiked structure with a smooth surface and a uniform diameter of around 30  $\mu\text{m}$ . It is worth noting that the needle structure offers a large surface area for the subsequent synthesis of  $\text{C}_3\text{N}_4$ , while the hollow structure reduces the weight of the micromotor catalyst and contributes to fast motion. The FESEM image of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC showed that free-standing  $\text{C}_3\text{N}_4$  nanosheets were grown uniformly on the surface of the sunflower pollen in an edge-to-face stacking mode to form a 3D hierarchical structure, avoiding the agglomeration of  $\text{C}_3\text{N}_4$  sheets (Fig. 2b and c). Furthermore, the TEM images further revealed the detailed structural characteristics of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC. Irregular spherical  $\text{Mn}_3\text{O}_4$  particles with an average grain size of 30 nm were uniformly grown on the surface of the interlaced  $\text{C}_3\text{N}_4$  nanosheets (Fig. 2d and e). As shown in Fig. 2f, there was a lattice fringe of 0.29 nm ascribed to the (110) plane of  $\text{Mn}_3\text{O}_4$ . Besides, Fig. 2g presents the SAED photograph of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC, revealing clear diffraction rings corresponding to the (020), (022), (110), and (112) crystal planes of  $\text{Mn}_3\text{O}_4$ , consistent with the XRD characterization results. Fig. 2h–l show the energy-dispersive X-ray (EDX) mappings of the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC. The elements C, N, C, Mn, and O were well distributed on the surface of the catalyst, but the distribution of Mn was more focused on one side of the microsphere (Fig. 2l), confirming the successful semi-coating of  $\text{Mn}_3\text{O}_4$  on the surface of the catalyst.

### Motion behaviours of the $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC micromotor catalyst

The self-propelled movement of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC was powered by the  $\text{O}_2$  bubbles generated from the  $\text{H}_2\text{O}_2$  decomposition by  $\text{Mn}_3\text{O}_4$ . Video S1† displays the movement of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC in different concentrations of  $\text{H}_2\text{O}_2$  solution containing 0.5%

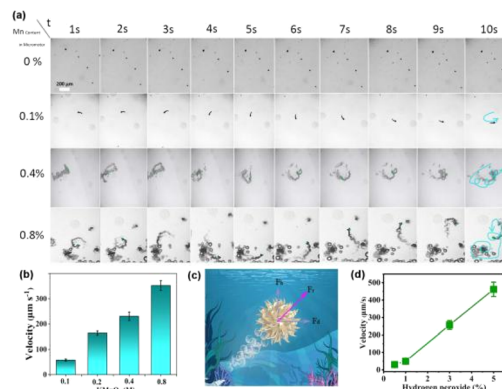


Fig. 3 Time-lapse images of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC in 1 wt%  $\text{H}_2\text{O}_2$  solution containing 0.5 wt% sodium dodecyl sulfonate under different Mn contents (a). Bar chart of micromotor catalyst velocity under different Mn contents (b). The force condition of the micromotor catalyst during movement (c). Effect of fuel  $\text{H}_2\text{O}_2$  concentration on the movement rate (d).

sodium dodecyl sulfonate. No regular curve or circular motion were observed in Video S1.† Moreover, the amount of manganese precursor had an important effect on the motor power. The effect of  $\text{KMnO}_4$  concentration on the velocity of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC is shown in Fig. 3a. The average velocity increased from  $57.83 \pm 2.24 \mu\text{m s}^{-1}$  to  $352.45 \pm 19.23 \mu\text{m s}^{-1}$  as the  $\text{KMnO}_4$  concentration was increased from 0.1 to 0.8 M (Fig. 3b). In addition, the hollow structure of the catalyst generated buoyancy in the fluid, which also had a positive effect on the rapid movement of the catalyst. As shown in Fig. 3c, the resultant force ( $F_r$ ) for motion was a combination of the driving force ( $F_d$ ) produced by the bubbles and the buoyancy ( $F_b$ ) of the motor itself. The  $\text{O}_2$  bubbles generated a strong momentum that propelled the catalyst forward with a velocity of  $30.4 \pm 3.4$ ,  $49.6 \pm 7.2$ ,  $256.8 \pm 25.2$ ,  $462.6 \pm 41.2 \mu\text{m s}^{-1}$  in 0.5, 1, 3, and 5 wt%  $\text{H}_2\text{O}_2$  (Fig. 3d), respectively. Besides, the drag force, mechanical work, chemical input power, and working efficiency are summarized in Table 1, where it can be observed that as the concentration of hydrogen peroxide increased, all four parameters gradually increased. The work efficiency of the micromotor catalyst was in the order of  $10^{-7}$ . These excellent kinetic and mechanical parameters lay a foundation for the application of such catalysts in micro-space or trace concentration fields.

### Micromotor-assisted degradation of CBZ

Fig. 4a shows the CBZ degradation by different samples. About 79.3% and 55.5% CBZ were degraded by  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  in 300 min, which were much higher rates than those achieved by g- $\text{C}_3\text{N}_4$ ,  $\text{C}_3\text{N}_4/\text{C}$ , and  $\text{H}_2\text{O}_2$ . The degradation rate of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC was always higher than that of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ , especially in the degradation stage at low CBZ concentration, which was due to the active contact of the catalyst with the pollutant molecules. Besides, several control experiments were carried out to verify the roles of movement in the degradation of CBZ. Fig. 4b shows CBZ degradation by under different conditions:  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC,  $\text{C}_3\text{N}_4/\text{C}$ -100-stirring



Table 1 Velocity and work efficiency of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC in different concentrations of  $\text{H}_2\text{O}_2$ 

$\text{H}_2\text{O}_2/\%$	$v/10^{-6} \text{ m s}^{-1}$	$F_{\text{drag}}/10^{-9} \text{ N}$	$P_{\text{mecha}}/10^{-15} \text{ W per motor}$	$P_{\text{chem}}/10^{-8} \text{ W per motor}$	$\eta/10^{-7}$
0.5	30.4	1.79	489.7	45.6	5.67
1	49.6	2.79	623.01	62.8	9.92
3	256.8	6.95	2445.01	83.1	29.42
5	462.6	20.06	11 991.87	314.7	38.11

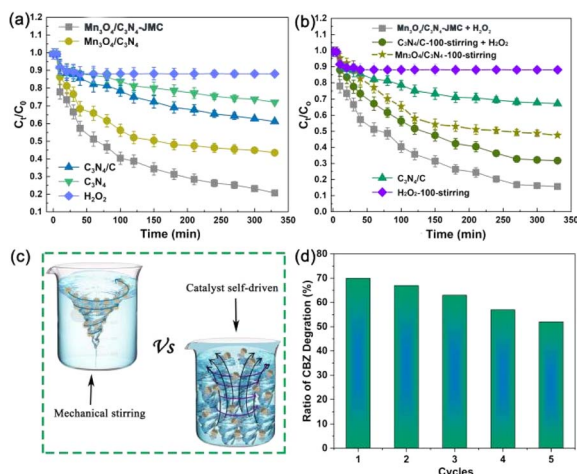


Fig. 4 Degradation of CBZ using different samples in 3% mass fraction of  $\text{H}_2\text{O}_2$  (a). CBZ degradation by  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC,  $\text{C}_3\text{N}_4/\text{C}$ -100-stirring (mechanical stirring at a speed of 100 rpm with  $\text{H}_2\text{O}_2$ ),  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -100-stirring (mechanical stirring at a speed of 100 rpm without  $\text{H}_2\text{O}_2$ ),  $\text{C}_3\text{N}_4/\text{C}$  without  $\text{H}_2\text{O}_2$  and with  $\text{H}_2\text{O}_2$  with mechanical stirring at 100 rpm (b) schematic of the CBZ degradation principle via mechanical agitation and micromotor catalyst motion (c); reusability of the catalyst for CBZ degradation (d).

(mechanical stirring with speed of 100 rpm with  $\text{H}_2\text{O}_2$ ),  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -100-stirring (mechanical stirring with speed of 100 rpm without  $\text{H}_2\text{O}_2$ ),  $\text{C}_3\text{N}_4/\text{C}$  without  $\text{H}_2\text{O}_2$  and  $\text{H}_2\text{O}_2$  with mechanical stirring 100 rpm. The comparison results showed that the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC exhibited higher catalytic activity towards CBZ than static  $\text{C}_3\text{N}_4/\text{C}$  and  $\text{C}_3\text{N}_4/\text{C}$ -100 under mechanical stirring with a speed of 100 rpm, indicating the positive contribution of autonomous movement. It is an important point that the random autonomous movement of the catalyst results in turbulence of the reactive liquid, facilitating more efficient contact between the active sites and the contaminants. Conventional mechanical agitation results in liquid flow in a certain direction, and the active site is in contact with contaminants only to a certain extent (Fig. 4c). Therefore, the degradation of pollutants by our catalyst system is superior to mechanical agitation in a certain range. Meanwhile, the reusability of the catalyst for the catalytic degradation of CBZ was studied. As shown in Fig. 4d, the CBZ removal rate was 54.7% at the end of the 5th cycle in 100 min, slightly lower than that for the first cycle (69.3%), indicating its high reusability for CBZ degradation. Besides, to clarify the active species responsible for the catalysis activity, and as shown in Fig. S2,† IPA, EDTA, and *P*-benzoquinone were used as scavengers for  $\cdot\text{OH}$ , oxygen vacancies, and  $\cdot\text{O}^{2-}$ .<sup>44</sup> It was found that hydroxyl free radicals

( $\cdot\text{OH}$ ) and superoxide free radicals ( $\cdot\text{O}^{2-}$ ) were produced in large quantities in the process of degradation. Comparatively,  $\cdot\text{OH}$  radicals were the main active species.

The kinetic study for the photocatalytic degradation of CBZ was investigated using pseudo-first-order and pseudo-second-order kinetic models.<sup>45,46</sup> Fig. 4a presents the photocatalytic degradation plot of CBZ using  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC, in which the operating condition of 50 mg per L photocatalyst loading was applied. The pseudo-first-order kinetics based on Langmuir-Hinshelwood kinetics when a small initial concentration of the reactant is used can be described by eqn (2):

$$-\ln C_t/C_0 = k_1 t \quad (2)$$

where  $C_0$  ( $\text{mg L}^{-1}$ ) and  $C_t$  ( $\text{mg L}^{-1}$ ) are CBZ concentrations at initial and reaction times, respectively,  $t$  is the irradiation time, and  $k_1$  is the apparent first-order rate constant of CBZ degradation. Fig. S3(a)† depicts the plot of  $-\ln C_t/C_0$  vs. time for  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC within 100 min irradiation. The slope of the linear fitted plot depicts  $k_1$ , which was calculated as  $0.0088 \text{ min}^{-1}$  and the coefficient of determination ( $R^2$ ) was obtained as 0.9253. Therefore, the pseudo-first-order kinetics model showed an unsatisfactory quality of linear fitting.

The pseudo-second-order kinetic model can be described by eqn (3):

$$1/C_t - 1/C_0 = k_2 t \quad (3)$$

where  $k_2$  ( $\text{L mg}^{-1} \text{ min}^{-1}$ ) is the second-order kinetics rate constant and was determined from a linear fitting of the data.

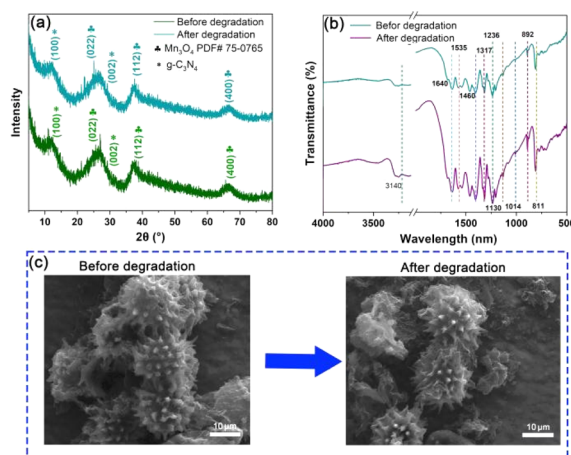


Fig. 5 XRD patterns (a) and FT-IR spectra (b) before and after CBZ degradation by  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC. FESEM images of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC before and after CBZ degradation (c).





Fig. S3(b)† shows the plot of  $(1/C_t - 1/C_0)$  vs.  $t$  within 300 min irradiation. From the slope of the linear fitted plot,  $k_2$  could be calculated as  $0.0144 \text{ L mg}^{-1} \text{ min}^{-1}$  and  $R^2$  was obtained as 0.9793. A much higher fitting quality was obtained by the pseudo-second-order kinetics model equation. Based on these results, it could be concluded that the photodegradation of CBZ using  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC followed a pseudo-second-order reaction, indicating that the rate of reaction with the autonomous movement catalyst was not only related to the concentration of reactants, but also to the concentration of the intermediate or transformation products generated by the reaction.

### Stability of the $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC micromotor catalyst

Based on the degradation efficiency of CBZ, the stability of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC was tested under the same conditions, as shown in Fig. 5a and b. Comparing the XRD and FT-IR patterns before and after the degradation of CBZ, there was no significant change in the phase composition and bonding state in the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC samples, indicating the high chemical stability and reusability of the catalysts in the process of CBZ degradation. Furthermore, the FESEM image of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC before and after five cycles (Fig. 5c) shows that its morphology remained almost unchanged after the cyclic testing, demonstrating its high stability and reusability.

### Energy band structure analysis of $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC by electrochemical studies and DFT theory

To reveal the degradation mechanism of carbamazepine, light-response, photocurrent, and electrochemical impedance tests were carried out to evaluate the photocatalytic activity of the catalyst. As shown in Fig. 6a,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  exhibited a shift towards a long wavelength region compared to  $\text{C}_3\text{N}_4$ , indicating that  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  had a higher adsorption ability for visible light

than  $\text{C}_3\text{N}_4$ ; while  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC showed a higher adsorption ability of  $\text{C}_3\text{N}_4$ , due to the carbonization of pollen (C), which increased the separation of photoelectrons. Meanwhile,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC had a higher photocurrent intensity than  $\text{g-C}_3\text{N}_4$  (Fig. 6b), suggesting the stronger charge-transfer ability of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC than  $\text{g-C}_3\text{N}_4$ . Besides, the electrochemical analysis showed that  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC displayed a smaller hemicycle radius than  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  and  $\text{g-C}_3\text{N}_4$  in the electrochemical impedance spectroscopy analysis (Fig. 6c), indicating a more efficient production of photoexcited electrons and holes.

To further understand the charge-transport effect between  $\text{g-C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$ , DFT calculations were performed to illustrate the charge-transfer property. Fig. 6(d)-1 and (d)-2 show the charge-transfer situation in the interface of  $\text{g-C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$ . Moreover, Fig. 6(d)-3 exhibits the charge-transfer situation in the interface of the heterojunction  $\text{g-C}_3\text{N}_4/\text{Mn}_3\text{O}_4$ . It can be clearly observed from Fig. 6(d)-3 that there was obvious charge transfer between the two phases, verifying the hetero-effect that the loading of  $\text{Mn}_3\text{O}_4$  could affect the electronic structure on the surface of the 2D  $\text{g-C}_3\text{N}_4$ .<sup>47–49</sup>

Fig. 7(a)–(d) exhibit the charge difference distribution and the corresponding electronic location function images to reveal the charge separation condition between  $\text{Mn}_3\text{O}_4$  and  $\text{g-C}_3\text{N}_4$ . It is obvious that when the heterostructure between the  $\text{g-C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$  was formed, the cyan and yellow region show that there was an accumulation of electrons migrating from  $\text{g-C}_3\text{N}_4$  to the  $\text{Mn}_3\text{O}_4$ . Meanwhile, the electronic location function image demonstrated that there existed a distinct covalent interaction between the C atom of  $\text{g-C}_3\text{N}_4$  triazine and the Mn layer, forming a charge-transfer channel for transfer from  $\text{g-C}_3\text{N}_4$  to the  $\text{Mn}_3\text{O}_4$  nanosheets.<sup>50–52</sup> So the above experimental test and theoretical calculation together confirmed that when  $\text{g-C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$  formed a composite, the charge separation and transfer ability between them could be dramatically improved compared to that of their counterparts alone.

The results and analysis in Fig. 5(a)–(c) prove that the heterojunction structure can significantly improve the electron conduction efficiency. Likewise, the band structure and density of states calculation were conducted to confirm the results. Fig. 8 shows the band structure of the heterostructure and density of states results of  $\text{g-C}_3\text{N}_4/\text{Mn}_3\text{O}_4$  and  $\text{g-C}_3\text{N}_4$ . It is worth noting that the band gap was underestimated by the functional

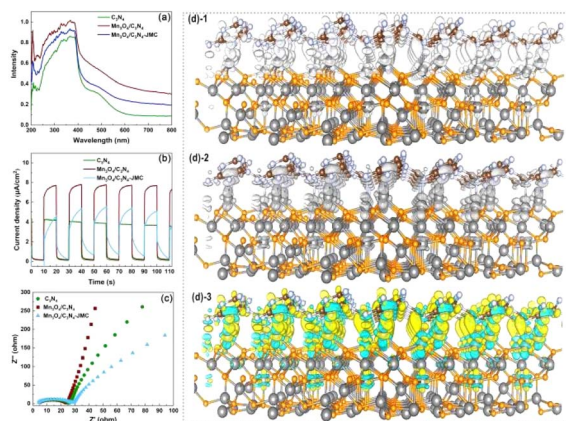


Fig. 6 UV-visible diffuse reflectance spectra of the as-prepared  $\text{C}_3\text{N}_4$ ,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ , and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC samples (a). Transient photocurrent density of the samples (b). Electrochemical impedance spectra of  $\text{C}_3\text{N}_4$ ,  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ , and  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC (c). Charge density difference between  $\text{Mn}_3\text{O}_4$  and  $\text{g-C}_3\text{N}_4$  interface. Positive part (d)-1, negative part (d)-2, and mixed distribution (d)-3. Yellow stands for the positive part, which indicates charge accumulation, whereas light green indicates charge depletion after hetero-contact formation. Up:  $\text{g-C}_3\text{N}_4$ , down:  $\text{Mn}_3\text{O}_4$ .

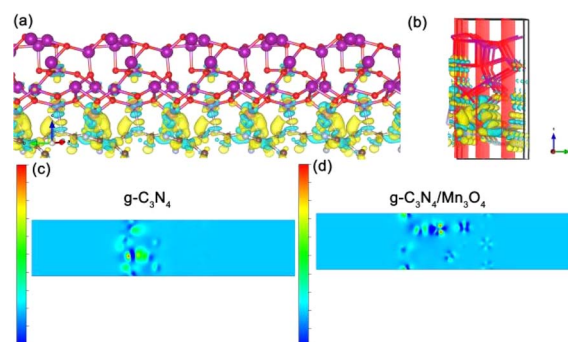


Fig. 7 Charge difference distribution (a) (b) and electronic location function analysis of the  $\text{g-C}_3\text{N}_4$  (c) and  $\text{g-C}_3\text{N}_4/\text{Mn}_3\text{O}_4$  composite (d).



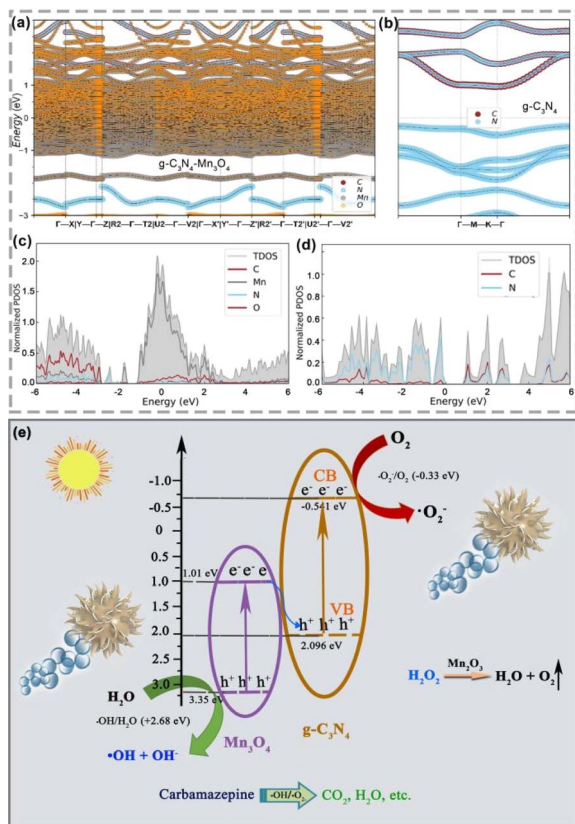


Fig. 8 Projected-to-element band structures of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  (a) and  $\text{g-C}_3\text{N}_4$  (b). Projected-to-element density of states (DOS) of  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  (c) and  $\text{g-C}_3\text{N}_4$  (d). Schematic of the photogenerated electron and hole migration and separation on the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$ -JMC heterojunction under visible-light irradiation (e).

PBE, but it made the comparison in the identity accuracy level. As shown in Fig. 8a, the O and Mn element bands occupied the dominant Fermi level and overlapped the C and N elements, indicating the hetero-effect charge transfer and sharing between the two different phases. In terms of the band structure (Fig. 8b), it could be observed that the pristine  $\text{g-C}_3\text{N}_4$  had a clear band gap, while the band gap disappeared in the heterogeneous structure. Furthermore, intensive crossing to the Fermi level (0 eV calibrated) indicated the better electronic conductivity. Moreover, the projected-to-element density of states was investigated and the results analyzed. From Fig. 8c, it could be observed that the most prominent peak occupied the Fermi level while isolated  $\text{g-C}_3\text{N}_4$  was not found, in line with the experimental observation and previous band calculations, which confirmed the better electronic conductivity of the hetero-interface of  $\text{g-C}_3\text{N}_4/\text{Mn}_3\text{O}_4$  (Fig. 8d). Therefore, constructing a multi-interface is a critical step for  $\text{g-C}_3\text{N}_4$ -based materials. To explain the mechanism of photocatalysis, the valence band (VB) and conduction band (CB) edges of  $\text{g-C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$  nanosheets were estimated. The CB edge values were calculated to be 1.01 and  $-0.541$  eV, while the VB edge values were 3.35 and 2.096 eV for  $\text{Mn}_3\text{O}_4$  and  $\text{g-C}_3\text{N}_4$ , respectively.<sup>53</sup> According to the above results, both  $\text{g-C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$  nanosheets could generate electrons and holes in the

heterojunction under visible-light irradiation (Fig. 8e). The result shows that the CB edge of  $\text{Mn}_3\text{O}_4$  was lower than the redox potential of  $\text{O}_2/\cdot\text{O}_2^-$  ( $-0.28$  eV), which means the electrons in the CB of  $\text{Mn}_3\text{O}_4$  could not form  $\cdot\text{O}_2^-$  radicals.<sup>54</sup> Similarly, since the VB edge of  $\text{g-C}_3\text{N}_4$  was higher than the redox potential of  $\cdot\text{OH}/\text{OH}^-$  (2.68 eV), the holes left in the VB of  $\text{g-C}_3\text{N}_4$  could not form  $\cdot\text{OH}$  radicals.<sup>55,56</sup> However, the photocatalytic degradation performance also decreased after adding the hole-trapping agent (Fig. S2†), indicating that both electrons and holes were involved in photocatalytic degradation. Therefore, the  $\text{g-C}_3\text{N}_4/\text{Mn}_3\text{O}_4$  nanosheets should form a Z-scheme heterojunction.<sup>57–59</sup> When exposed to visible light, the electrons transit from the VB of  $\text{g-C}_3\text{N}_4$  and  $\text{Mn}_3\text{O}_4$  to CB, leaving holes in the VB. The photogenerated electrons in the CB of  $\text{Mn}_3\text{O}_4$  transfer to the VB of  $\text{g-C}_3\text{N}_4$ . Therefore, holes are left in the VB of  $\text{Mn}_3\text{O}_4$ , while electrons are left in the CB of  $\text{g-C}_3\text{N}_4$ . The holes and electrons will react with  $\text{OH}^-$  and  $\text{O}_2$  to form free radicals with strong oxidation and deoxidization, as confirmed by the experimental capture of active species, which further oxidize CBZ molecules.

In addition, it is worth mentioning that, as an artificial active object, the micromotor catalyst can enhance mass transfer in the solution and improve the interaction between the active surface and the target pollutants. Hence, combined with the degradation path and assisted by the self-propelled motion of the catalyst, a very high CBZ degradation rate was obtained.

## Conclusions

In summary, an urchin-like  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  Janus micromotor catalyst was precisely designed and synthesized for the dynamic photocatalytic degradation of carbamazepine from sewage. The asymmetric distribution of  $\text{Mn}_3\text{O}_4$  on the surface of the catalyst was achieved by a facile semi-coating method. The self-propulsion of the Janus micromotor catalyst was achieved with a speed of  $223.5 \pm 7.0 \mu\text{m s}^{-1}$  through the  $\text{O}_2$  bubbles generated from the decomposition of  $\text{H}_2\text{O}_2$  by  $\text{Mn}_3\text{O}_4$ . Furthermore, the photoelectrons generated from the  $\text{Mn}_3\text{O}_4/\text{C}_3\text{N}_4$  heterojunction could create active species and promote the photocatalytic degradation activity of the catalyst under simulated sunlight irradiation. Consequently, nearly 70% of CBZ could be degraded within 5 h with the help of the movement. Besides, density functional theory calculations proved that the formation of the heterojunction structure promoted the generation of photogenerated carriers. Therefore, these features endow the micromotor catalyst with exciting potential in environmental remediation fields.

## Data availability

The authors confirm that the data supporting the findings of this study are available within the article and its ESI.†

## Author contributions

Jie Yang: conceptualization; data curation; formal analysis; investigation; visualization; writing – original draft. Wenning Yang: data curation; formal analysis. Chao Zhang:





conceptualization; formal analysis. Jian Gong: investigation; supervision. Jia Li: conceptualization; funding acquisition; supervision; writing – review & editing; Ming Xu: conceptualization; resources; funding acquisition. Chengzhang Liu: data curation, resources.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by the Natural Science Foundation of Shandong [Grant No. ZR2020KE057], Fundamental Research Funds for the Central Universities (buctrc 202112) and the National Natural Science Foundation of China (NSFC: 22102007). Computational resources were provided by Extreme Science and Engineering Discovery Environment (XSEDE Allocation TG-CHE190010) and the Texas Advanced Computing Center (TACC).

## Notes and references

- Z. Zhou, Y. Wu, Y. Xu, Z. Wang, H. Fu and Y. Zheng, Carbamazepine degradation and genome sequencing of a novel exoelectrogen isolated from microbial fuel cells, *Sci. Total Environ.*, 2022, **838**, 156161.
- P. J. Mafa, M. E. Malefane, A. O. Idris, B. B. Mamba, D. Liu, J. Z. Gui and A. T. Kuvarega, Cobalt oxide/copper bismuth oxide/samarium vanadate ( $\text{Co}_3\text{O}_4/\text{CuBi}_2\text{O}_4/\text{SmVO}_4$ ) dual Z-scheme heterostructured photocatalyst with high charge-transfer efficiency: Enhanced carbamazepine degradation under visible light irradiation, *J. Colloid Interface Sci.*, 2021, **603**, 666–684.
- L. Huang, T. Zeng, X. Xu, Z. He, J. Chen and S. Song, Immobilized hybrids between nitrogen-doped carbon and stainless steel derived  $\text{Fe}_3\text{O}_4$  used as a heterogeneous activator of persulfate during the treatment of aqueous carbamazepine, *Chem. Eng. J.*, 2019, **372**, 862–872.
- S. Wang and J. Wang, Carbamazepine degradation by gamma irradiation coupled to biological treatment, *J. Hazard. Mater.*, 2017, **321**, 639–646.
- X. He, Y. Luo, Y. Yi, S. Su and W. Qin, Peroxymonosulfate activation by Fe-Mn Co-doped biochar for carbamazepine degradation, *RSC Adv.*, 2024, **14**, 1141–1149.
- J. Wang and S. Wang, Removal of pharmaceuticals and personal care products (PPCPs) from wastewater: a review, *J. Environ. Manage.*, 2016, **182**, 620–640.
- Y. Deng, B. Eitzer, J. C. White and B. Xing, Impact of multiwall carbon nanotubes on the accumulation and distribution of carbamazepine in collard greens (*Brassica oleracea*), *Environ. Sci.: Nano*, 2017, **4**, 149–159.
- J. Zhai, X. Wang, J. Yan, C. Gong, W. Zhu, Y. Luo, D. Yang and X. Gao, Oxidized pyrite as an efficient Fenton reagent to generate active species for the degradation of carbamazepine in a wide pH range, *Appl. Surf. Sci.*, 2023, **622**, 158098.
- B. Zhang, L. Zhang, K. Akiyama, P. Bingham, Y. Zhou and K. Shiro, Self-assembly of nanosheet-supported Fe-MOF heterocrystals as a reusable catalyst for boosting advanced oxidation performance via radical and nonradical pathways, *ACS Appl. Mater. Interfaces*, 2021, **13**, 22694–22707.
- Y. Zhou, Z. Wang, W. Fang, R. Qi, Z. Wang, C. Xia, K. Lei, B. You, X. Yang and Y. Liu, Modulating O-H activation of methanol oxidation on nickel-organic frameworks for overall  $\text{CO}_2$  electrolysis, *ACS Catal.*, 2023, **13**, 2039–2046.
- B. Mapleback, N. Brack, L. Thomson, M. Spencer, D. Osborne, S. Doshi, E. Thostenson and A. Rider, Development of stable boron nitride nanotube and hexagonal boron nitride dispersions for electrophoretic deposition, *Langmuir*, 2020, **36**, 3425–3438.
- A. Kumar, M. Khan, J. He and I. Lo, Visible-light-driven magnetically recyclable terephthalic acid functionalized  $\text{g-C}_3\text{N}_4/\text{TiO}_2$  heterojunction nanophotocatalyst for enhanced degradation of PPCPs, *Appl. Catal., B*, 2020, **270**, 11889.
- L. Yang, L. Liang, L. Wang, J. Zhu, S. Gao and X. Xia, Accelerated photocatalytic oxidation of carbamazepine by a novel 3D hierarchical protonated  $\text{g-C}_3\text{N}_4/\text{BiOBr}$  heterojunction: Performance and mechanism, *Appl. Surf. Sci.*, 2019, **473**, 527–539.
- Y. Zhu, Y. Liu, Q. Ai, G. Gao, L. Yuan, Q. Fang, X. Tian, X. Zhang, E. Egap and P. Ajayan, In situ synthesis of lead-free halide perovskite-COF nanocomposites as photocatalysts for photoinduced polymerization in both organic and aqueous phases, *ACS Mater. Lett.*, 2022, **4**, 464–471.
- Z. Xia, B. Liu, Y. Xiao, W. Hu, M. Deng and C. Lu, Integrating hybrid perovskite nanocrystals into metal-organic framework as efficient S-scheme heterojunction photocatalyst for synergistically boosting controlled radical photopolymerization under 980 nm NIR light, *ACS Appl. Mater. Interfaces*, 2023, **15**, 57119–57133.
- A. Khundi, A. Habibi-Yangjeh, M. Abitorabi and S. Pouran, Review on photocatalytic conversion of carbon dioxide to value-added compounds and renewable fuels by graphitic carbon nitride-based photocatalysts, *Catal. Rev.*, 2019, **61**, 595–628.
- A. Khundi, A. Badieli, G. Ziarani, A. Habibi-Yangjeh, M. Munoz-Batista and R. Luque, Graphitic carbon nitride-based photocatalysts: toward efficient organic transformation for value-added chemicals production, *Mol. Catal.*, 2020, **488**, 110902.
- L. Shi, K. Chang, H. Zhang, X. Hai, L. Yang, T. Wang and J. Ye, Drastic enhancement of photocatalytic activities over phosphoric acid protonated porous  $\text{g-C}_3\text{N}_4$  nanosheets under visible light, *Small*, 2016, **12**, 4431–4439.
- M. Nakhaei, Z. Barzgar, S. Mohammadi and A. Ghazizadeh, Preparation of  $\text{MnO}_2/\text{bentonite}$  nanocomposite with enhanced photocatalytic activity under sunlight irradiation, *Res. Chem. Intermed.*, 2019, **45**, 4995–5005.
- S. Das, A. Sarnanta and S. Jana, Light-assisted synthesis of hierarchical flower-like  $\text{MnO}_2$  nanocomposites with solar light induced enhanced photocatalytic activity, *ACS Sustain. Chem. Eng.*, 2017, **5**, 9086–9094.



- 21 A. Baral, D. Das, M. Minakshi, M. Ghosh and D. Padhi, Probing environmental remediation of RhB organic dye using  $\alpha$ -MnO<sub>2</sub> under visible-light irradiation: structural, photocatalytic and mineralization studies, *ChemistrySelect*, 2016, **1**, 4277–4285.
- 22 Y. Shi, M. Zhang, Y. Li, G. Liu, R. Jin, Q. Wang, H. Xu and S. Gao, 2D/1D protonated g-C<sub>3</sub>N<sub>4</sub>/ $\alpha$ -MnO<sub>2</sub> Z-scheme heterojunction with enhanced visible-light photocatalytic efficiency, *Ceram. Int.*, 2020, **46**, 25905–25914.
- 23 Q. Zhang, Y. Peng, F. Deng, M. Wang and D. Chen, Porous Z-scheme MnO<sub>2</sub>/Mn-modified alkalized g-C<sub>3</sub>N<sub>4</sub> heterojunction with excellent Fenton-like photocatalytic activity for efficient degradation of pharmaceutical pollutants, *Sep. Purif. Technol.*, 2020, **246**, 116890.
- 24 C. Chen, M. Xie, L. Kong, W. Lu, Z. Feng and J. Zhan, Mn<sub>3</sub>O<sub>4</sub> nanodots loaded g-C<sub>3</sub>N<sub>4</sub> nanosheets for catalytic membrane degradation of organic contaminants, *J. Hazard. Mater.*, 2020, **390**, 122146.
- 25 K. Vikrant and K. H. Kim, Metal-organic framework micromotors: perspectives for environmental applications, *Catal. Sci. Technol.*, 2021, **11**, 6592–6600.
- 26 X. Zhang, C. Liu, J. Li, R. Chu, Y. Lyu and Z. Lan, Dual source-powered multifunctional Pt/FePc@Mn-MOF spindle-like Janus nanomotors for active CT imaging-guided synergistic photothermal/chemodynamic therapy, *J. Colloid Interface Sci.*, 2024, **657**, 799–810.
- 27 X. Yang, C. Liu, S. Gao, X. Zhang, Z. Lan, M. Zuo and J. Li, A novel bio-template route to synthesize enzyme-immobilized MOF/LDH tubular magnetic micromotors and their application in water treatment, *Environ. Sci.: Nano*, 2024, **11**, 1142–1156.
- 28 H. Zeng, Y. Wang, T. Jiang, H. Xia, X. Gu and H. Chen, Recent progress of biomimetic motions-from microscopic micro/nanomotors to macroscopic actuators and soft robotics, *RSC Adv.*, 2021, **11**, 27406–27419.
- 29 L. Chen, H. Yuan, S. Chen, C. Zheng, X. Wu, Z. Li, C. Liang, P. Dai, Q. Wang, X. Ma and X. Yan, Cost-Effective, high-yield production of biotemplated catalytic tubular micromotors as self-Propelled microcleaners for water treatment, *ACS Appl. Mater. Interfaces*, 2021, **13**, 31226–31235.
- 30 H. Wang, M. Potroz, J. Jackman, B. Khezri, T. Marić, N. Cho and M. Pumera, Bioinspired spiky micromotors based on sporopollenin exine capsules, *Adv. Funct. Mater.*, 2017, **27**, 1702338.
- 31 S. Gao, C. Liu, X. Yang, Z. Lan, M. Zuo, P. Yang and J. Li, New synthetic strategy toward a natural enzyme-nanozyme hybrid dual-function nanomotor and its application in environmental remediation, *Catal. Sci. Technol.*, 2024, **14**, 1239–1254.
- 32 Y. Zhu, Z. Yuan, J. Rong, T. Zhang, D. Yang, J. Pan and F. Qiu, Engineering flower-shaped hierarchical micromotors on a sustainable biotemplate by teamed boronate affinity-based surface imprinting for effective separation of shikimic acid, *Sep. Purif. Technol.*, 2024, **336**, 126345.
- 33 Y. Song, Z. Song, J. Wu, Z. Li, X. Gu, C. Wang, L. Wang and J. Liang, Focus on the performance enhancement of micro/nanomotor-based biosensors, *Biosens. Bioelectron.*, 2023, **241**, 115686.
- 34 Z. Lan, T. Li, Q. Li, C. Liu and J. Li, Multifunctional glucose-powered nanomotors with robust dual enzyme mimic activities, *Sep. Purif. Technol.*, 2024, **333**, 125860.
- 35 W. Wang, T. Chiang, D. Velegol and T. Mallouk, Understanding the efficiency of autonomous nano-and microscale motors, *J. Am. Chem. Soc.*, 2013, **135**, 10557–10565.
- 36 G. Kresse and J. Furthmüller, Efficient iterative schemes for ab initio total-energy calculations using a plane-wave basis set, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **54**, 11169–11186.
- 37 J. Perdew, J. Chevary, S. Vosko, K. Jackson, M. P. D. Singh and C. Fiolhais, Atoms, molecules, solids, and surfaces: Applications of the generalized gradient approximation for exchange and correlation, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1992, **46**, 6671–6687.
- 38 S. Froyen, Brillouin-zone integration by Fourier quadrature: Special points for superlattice and supercell calculations, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1989, **139**, 5–15.
- 39 K. Wang, X. Wang, H. Pan, Y. Liu, S. Xu and S. Cao, In situ fabrication of CDs/g-C<sub>3</sub>N<sub>4</sub> hybrids with enhanced interface connection via calcination of the precursors for photocatalytic H<sub>2</sub> evolution, *Int. J. Hydrogen Energy*, 2018, **43**, 91–99.
- 40 D. Zeng, W. Ong, H. Zheng, M. Wu, Y. Chen, D. Peng and M. Han, Ni<sub>12</sub>P<sub>5</sub> nanoparticles embedded into porous g-C<sub>3</sub>N<sub>4</sub> nanosheets as a noble-metal-free heterostructure photocatalyst for efficient H<sub>2</sub> production under visible light, *J. Mater. Chem. A*, 2017, **5**, 16171–16178.
- 41 K. Sing, H. Everett, R. Haul, L. Moscou, R. Pierotti, J. Rouquerol and T. Siemieniowska, Reporting physisorption data for gas/solid systems with special reference to the determination of surface area and porosity, *Pure Appl. Chem.*, 1985, **57**, 603–619.
- 42 H. Sampatkumar, A. Antony, M. Trivedi, M. Sharma, M. Ghate, M. Baidya, R. Dateer and S. Patil, In situ biosynthesis of palladium nanoparticles on banana leaves extract-coated graphitic carbon nitride: An efficient and reusable heterogeneous catalyst for organic transformations and antimicrobial agent, *Biomass Convers. Biorefin.*, 2024, **14**, 10045–10066.
- 43 J. Cao, C. Qin, Y. Wang II, B. Zhang, Y. Gong, H. Zhang, G. Sun, H. Bala and Z. Zhang, Calcination method synthesis of SnO<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> composites for a high-performance ethanol gas sensing application, *Nanomaterials*, 2017, **7**, 98.
- 44 J. Parmar, D. Vilela, K. Villa, J. Wang and S. Sanchez, Micro-and nanomotors as active environmental microcleaners and sensors, *J. Am. Chem. Soc.*, 2018, **140**, 9317–9331.
- 45 H. Tran, D. Nguyen, P. Do and U. Tran, Kinetics of photocatalytic degradation of organic compounds: a mini-review and new approach, *RSC Adv.*, 2023, **13**, 16915–16925.
- 46 K. Kalantaria and E. Asgari, Photocatalytic degradation of Direct Red 80 using ZnO/ZnTiO<sub>3</sub>/Zn<sub>2</sub>Ti<sub>3</sub>O<sub>8</sub> ternary nanocomposite, *Desalin. Water Treat.*, 2023, **284**, 240–250.



- 47 M. F. R. Samsudin, H. Ullah, A. A. Tahir, X. Li, Y. H. Ng and S. Sufian, Superior photoelectrocatalytic performance of ternary structural BiVO<sub>4</sub>/GQD/g-C<sub>3</sub>N<sub>4</sub> heterojunction, *J. Colloid Interface Sci.*, 2021, **586**, 785–796.
- 48 Y. Huang, K. Wang, T. Gao, J. Li, X. Wu and G. Zhang, Construction of 2D/2D Bi<sub>2</sub>Se<sub>3</sub>/g-C<sub>3</sub>N<sub>4</sub> nanocomposite with High interfacial charge separation and photo-heat conversion efficiency for selective photocatalytic CO<sub>2</sub> reduction, *Appl. Catal., B*, 2020, **277**, 119232.
- 49 Y. Li, M. Zhou, B. Chen and Y. Shao, Recent advances in g-C<sub>3</sub>N<sub>4</sub>-based heterojunction photocatalysts, *J. Mater. Sci. Nanotechnol.*, 2020, **56**, 1–17.
- 50 J. Fan, Q. Wang, W. Yan, J. Chen, X. Zhou and H. Xie, Mn<sub>3</sub>O<sub>4</sub>-g-C<sub>3</sub>N<sub>4</sub> composite to activate peroxydisulfate for organic pollutants degradation: Electron transfer and structure-dependence, *J. Hazard. Mater.*, 2022, **434**, 128818.
- 51 J. Xu, Y. Wang, J. Wan and L. Wang, Facile synthesis of carbon-doped CoMn<sub>2</sub>O<sub>4</sub>/Mn<sub>3</sub>O<sub>4</sub> composite catalyst to activate peroxydisulfate for ciprofloxacin degradation, *Sep. Purif. Technol.*, 2022, **287**, 120576.
- 52 W. Che, W. Cheng, T. Yao, F. Tang, W. Liu, H. Su, Y. Huang, Q. Liu, J. Liu, F. Hu, Z. Pan, Z. Sun and S. Wei, Fast photoelectron transfer in (cring)-C<sub>3</sub>N<sub>4</sub> Plane heterostructural nanosheets for overall water splitting, *J. Am. Chem. Soc.*, 2017, **139**, 3021–3026.
- 53 Y. Yi, J. Wang, Y. Niu, Y. Yu, S. Wu and K. Ding, Exploring the evolution patterns of melam from thermal synthesis of melamine to graphitic carbon nitride, *RSC Adv.*, 2022, **12**, 24311–24318.
- 54 B. Lin, H. Li, H. An, W. Hao, J. Wei, Y. Dai, C. Ma and G. Yang, Preparation of 2D/2D g-C<sub>3</sub>N<sub>4</sub> nanosheet@ZnIn<sub>2</sub>S<sub>4</sub> nanoleaf heterojunctions with well-designed high-speed charge transfer nanochannels towards high efficiency photocatalytic hydrogen evolution, *Appl. Catal., B*, 2018, **220**, 542–552.
- 55 Y. Zhang, H. Li, L. Zhang, R. Gao and W. Dai, Construction of highly efficient 3D/2D MnO<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> nanocomposite in the epoxidation of styrene with TBHP, *ACS Sustain. Chem. Eng.*, 2019, **7**, 17008–17019.
- 56 M. Xiong, L. Chen, Q. Yuan, J. He, S. Luo, C. Au and S. Yin, Controlled synthesis of graphitic carbon nitride/beta bismuth oxide composite and its high visible-light photocatalytic activity, *Carbon*, 2015, **86**, 217–224.
- 57 Y. Nosaka and A. Nosaka, Generation and detection of reactive oxygen species in photocatalysis, *Chem. Rev.*, 2017, **117**, 11302–11336.
- 58 P. Xia, B. Zhu, B. Cheng, J. Yu and J. Xu, 2D/2D g-C<sub>3</sub>N<sub>4</sub>/MnO<sub>2</sub> nanocomposite as a direct Z-scheme photocatalyst for enhanced photocatalytic activity, *ACS Sustain. Chem. Eng.*, 2018, **6**, 965–973.
- 59 P. Xia, S. Cao, B. Zhu, M. Liu, M. Shi, J. Yu and Y. Zhang, Designing a 0D/2D s-scheme heterojunction over polymeric carbon nitride for visible-light photocatalytic inactivation of bacteria, *Angew. Chem., Int. Ed.*, 2020, **59**, 5218–5225.

