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# Cobalt phthalocyanine (CoPc) anchored on Ti<sub>3</sub>C<sub>2</sub> MXene nanosheets for highly efficient selective catalytic oxidation†

Simeng Zhu,<sup>a</sup> Peng Liu<sup>b</sup> and Xinlin Hong \*<sup>a</sup>

Quinclorac is an important precursor for pharmaceutical, agricultural, and synthetic chemistry. The state-of-the-art synthesis of quinclorac *via* condensation, chlorination and oxidative hydrolysis often uses homogeneous catalysts and strong acid oxidant agents to promote the catalytic oxidation, which requires huge manpower input for the late-stage purification process and is usually environmentally unfriendly. In this work, we successfully fabricated a stable cobalt phthalocyanine (CoPc) Co-based composite (CoPc/Ti<sub>3</sub>C<sub>2</sub>) by anchoring CoPc on the surface of Ti<sub>3</sub>C<sub>2</sub> nanosheets for the selective oxidation of 3,7-dichloro-8-dichloro methyl quinoline (3,7-D-8-DMQ) into quinclorac. More impressively, CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br exhibits a high selectivity of 91.8% for the catalytic oxidation of 3,7-D-8-DMQ to quinclorac in acetic acid, with a quinclorac yield of 87.5%, which is approximately 2.46 times higher than that of pristine CoPc-Mn-Br. The obtained heterogeneous catalytic system shows good reusability. Detailed mechanistic investigations reveal that the system works through the free radical mechanism *via* the formation of Co<sup>2+</sup>/Co<sup>3+</sup> redox cycles. This work provides a new understanding for the stabilization of reaction intermediates and facilitates the design of catalysts for selective catalytic oxidation.

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## 1. Introduction

Selective oxidation is one of the most important transformations in the chemical industry and widely used for producing aldehydes, acid, and several other chemicals.<sup>1,2</sup> Compared to strong acid oxidants, oxygen is an inexpensive and green oxidant.<sup>3,4</sup> However, directly utilizing clean molecular oxygen as an oxidant is challenging because its ground state is a triplet. According to the Wigner spin selection rules, the direct reaction between the substrate and molecular oxygen is usually spin-forbidden.<sup>5</sup> On this basis, activation of ground-state molecular oxygen is a fundamental step in catalytic oxidation processes.<sup>6-8</sup> In this regard, it is imperative to develop a novel catalytic strategy to coordinate molecular oxygen and substrates in selective oxidation.

The selective oxidation synthesis of quinclorac is an essential laboratory and commercial procedure.<sup>9</sup> Industrially, quinclorac is synthesized by the condensation of aniline and glycerol, chlorination and oxidative hydrolysis, processes that

require large amounts of strong corrosive or toxic reagents, often in the homogeneous phase, which later require complex separation methods.<sup>10-12</sup> To tackle these problems, further exploring the combination of molecular complex catalysts with O<sub>2</sub> under green and eco-friendly conditions is one of the key development directions not only for selective oxidation reactions in organic synthesis, but also for the catalytic oxidation synthesis of quinclorac.<sup>13,14</sup>

Along with the intensive research efforts in developing catalysts with high performance, low cost and environmental friendliness, cobalt-based complex catalysts represented by cobalt phthalocyanine (CoPc) have emerged as a new alternative class of materials.<sup>15,16</sup> Transition-metal complexes, for example, CoPc, with well-defined active sites and adjustable structures have been widely used for various catalytic reactions.<sup>17-19</sup> However, the major challenge of utilizing homogeneous CoPc for catalysis application lies in the unstable properties, such as uncontrollable aggregation, oxidative self-destruction in the oxidizing solvent, and undesirable reduction of the Pc ligand in the reductive media, which resulted in poor long-term stability.<sup>20-22</sup> In addition, homogeneous catalysts suffer from separation and recycling problems. Thus, the heterogenization of CoPc provides a solution to maximize the stability without sacrificing the intrinsic activity and selectivity of homogeneous molecular catalysts.

Two-dimensional (2D) titanium carbide (Ti<sub>3</sub>C<sub>2</sub>) nanosheets, as a typical 2D MXene nanomaterial, contain abundant active

<sup>a</sup>College of Chemistry and Molecular Sciences, Wuhan University, Wuhan, China. E-mail: hongxl@whu.edu.cn

<sup>b</sup>Hubei Province Engineering and Technology Research Center for Fluorinated Pharmaceuticals, School of Pharmaceutical Sciences, Wuhan University, Wuhan, China

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sites, large surface area and excellent properties. Hence,  $\text{Ti}_3\text{C}_2$  MXene could be applied as the solid support for CoPc, increasing its uniform distribution and anchoring impact in  $\text{Ti}_3\text{C}_2$ , as well as its catalytic capability.<sup>23–25</sup> The attempt to produce quinclorac with molecularly dispersed metal phthalocyanine catalysts in the presence of molecular oxygen has never been reported, to the best of our knowledge.<sup>26–29</sup>

In this work, we report the realization of quinclorac synthesis enabled by anchoring CoPc onto  $\text{Ti}_3\text{C}_2$  MXene nanosheets for the design of a new kind of CoPc/ $\text{Ti}_3\text{C}_2$ -Mn(OAc)<sub>2</sub>-HBr (CoPc/ $\text{Ti}_3\text{C}_2$ -4.5%-Mn-Br) system with oxygen as the oxidant. This enables CoPc/ $\text{Ti}_3\text{C}_2$ -4.5%-Mn-Br to exhibit a high catalytic activity for the production of high value-added quinclorac with a yield of 87.5%, which is even 2.46-fold higher than that of the CoPc-Mn-Br system. By investigating the chemical state of the reaction active species, the favorable effect of  $\text{Co}^{2+}$  and  $\text{Co}^{3+}$  species leading to the excellent performance is revealed. A catalytic mechanism involving surmised radical generation and the oxidation intermediate species can be proposed based on experimental and theoretic analysis. This strategy provides the exact answer for selective catalytic oxidation and opens a new horizon for efficient organic synthesis conversion owing to its extremely high activity and cost-effective nature.

## 2. Experimental section

### 2.1 Synthesis of $\text{Ti}_3\text{C}_2$ MXene nanosheets

$\text{Ti}_3\text{AlC}_2$  powder (1.0 g) was added slowly into 20 mL of 40% HF. The mixture was stirred and heated at 80 °C overnight for selectively exfoliating the Al layers from  $\text{Ti}_3\text{AlC}_2$ . The as-synthesized product was washed with distilled water by centrifugation until the supernatant reached a pH value of approximately 7. To delaminate the etched MAX powder, the precipitate was stirred vigorously in 20 mL of tetramethylammonium hydroxide (TMAOH) for 24 hours. And the suspension was

washed with distilled water several times to completely eliminate the excess TMAOH. The remaining precipitate was further dispersed in fresh distilled water. Then, the solution was ultrasonically treated for 6 h in an ice bath. The resulting material was then treated for the defluorinative functionalization at 300 °C for 4 h under  $\text{H}_2$  flow. Finally, the resultant products were washed with distilled water followed by vacuum-drying at 60 °C overnight.

### 2.2 Synthesis of CoPc/ $\text{Ti}_3\text{C}_2$

Preparation of composite catalyst CoPc/ $\text{Ti}_3\text{C}_2$ -X using the self-assembly method. CoPc (0.2 g) was dispersed in 10.0 mL of  $\text{H}_2\text{O}$ , and then a certain amount of  $\text{Ti}_3\text{C}_2$  (containing 3 mg, 6 mg, 9 mg, 12 mg and 15 mg of  $\text{Ti}_3\text{C}_2$ , respectively) was added and stirred for 1 h. The final product was centrifuged, washed and vacuum-dried at 60 °C overnight. A series of products were prepared and labeled as CoPc/ $\text{Ti}_3\text{C}_2$ -X, X meant the mass ratio of  $\text{Ti}_3\text{C}_2$  to CoPc, which was 1.5%, 3%, 4.5%, 6% and 7.5%, respectively.

### 2.3 Catalytic activity test and characterization

The catalytic experiment and characterization are shown in the ESI.†

## 3. Results and discussion

### 3.1 Synthesis and characterization of catalysts

As shown in Fig. 1, the CoPc anchored  $\text{Ti}_3\text{C}_2$  MXene could be fabricated by a facile self-assembly strategy. Scanning electron microscopy (SEM) measurements of the as-prepared CoPc/ $\text{Ti}_3\text{C}_2$  hybrid show that the composite comprises comparatively uniform 2D nanosheets (Fig. 2a and b). High-resolution transmission electron microscopy (HRTEM) images of the composite confirm the high crystallinity of  $\text{Ti}_3\text{C}_2$  nanosheets and the

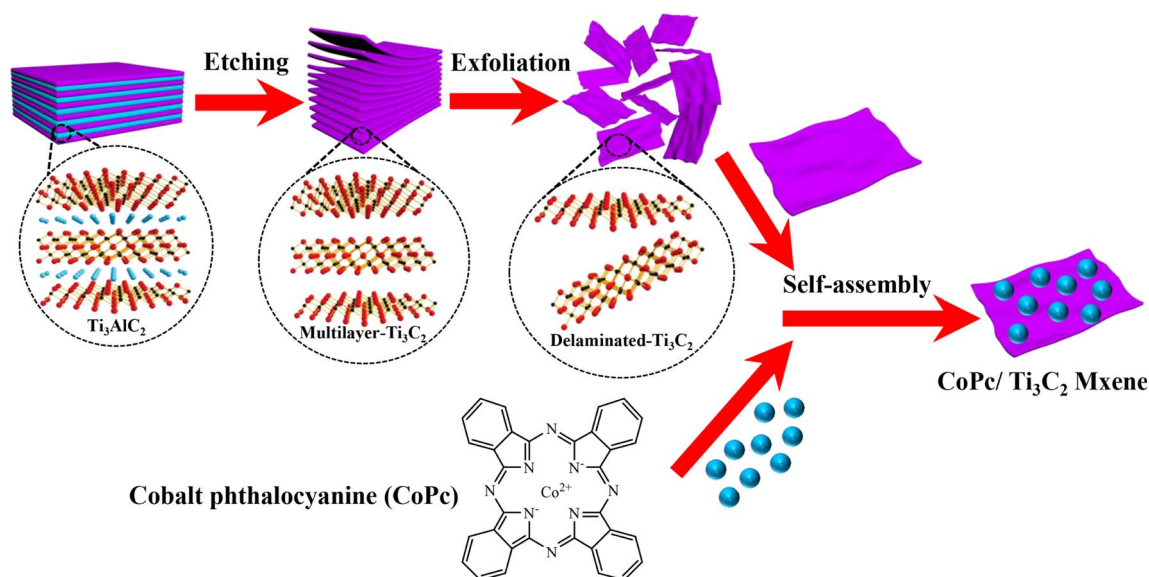


Fig. 1 Schematically synthetic procedure of the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid.



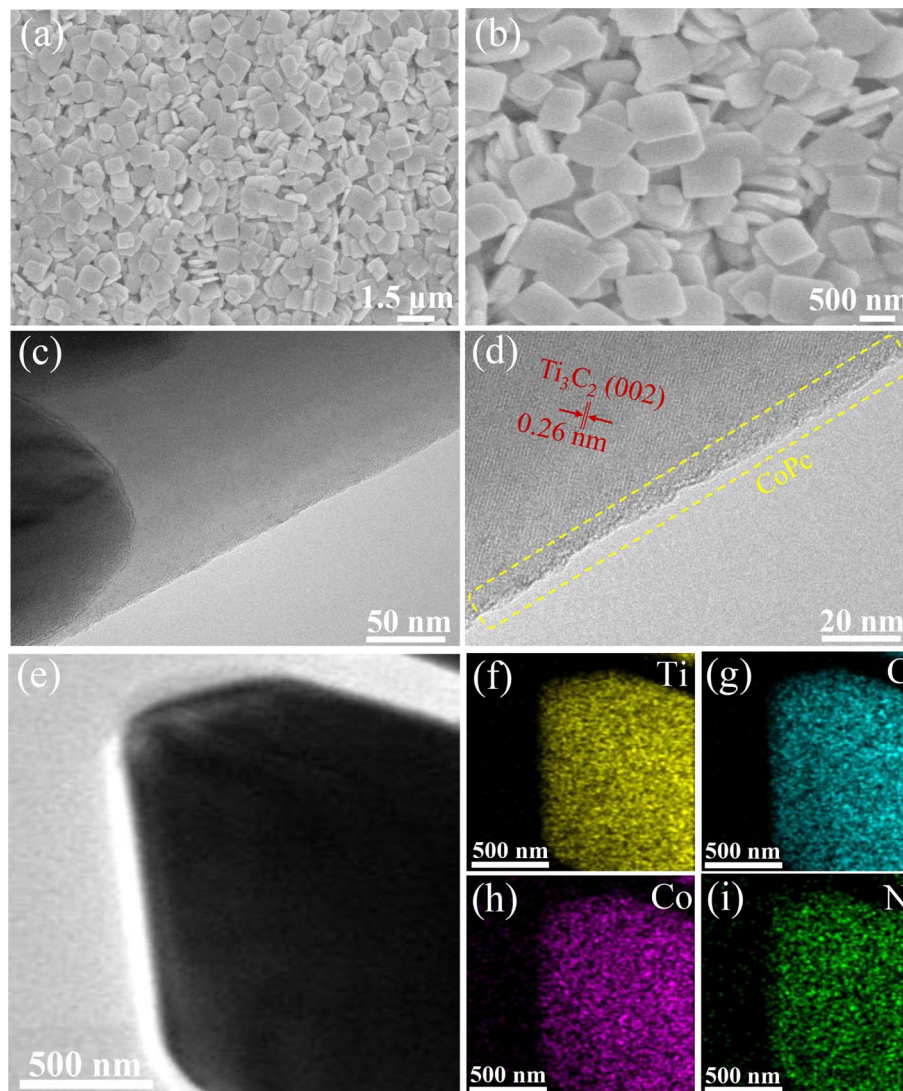


Fig. 2 SEM images (a and b) and high-resolution TEM (HRTEM) images (c and d) of CoPc/Ti<sub>3</sub>C<sub>2</sub>. (e) High-angle annular dark-field (HAADF) image of CoPc/Ti<sub>3</sub>C<sub>2</sub>. (f–i) The corresponding EDS elemental mapping images of Ti, C, Co and N elements.

amorphous morphology of CoPc molecules (Fig. 2c). As seen in Fig. 2d, the estimated lattice spacing of 0.260 nm can be attributed to the (002) facet of Ti<sub>3</sub>C<sub>2</sub>. The presence of rough spots in the HRTEM image of hybrid indicates the anchored CoPc units on Ti<sub>3</sub>C<sub>2</sub>, and no remarkable lattice spaces are observed for CoPc, which is attributed to the distribution of amorphous CoPc on the surface of Ti<sub>3</sub>C<sub>2</sub> nanosheets. High angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) and corresponding energy dispersive spectroscopy (EDS) elemental analysis have been utilized to observe the distribution of various species in the catalysts. Interestingly, a nanosheet-like composite material is formed according to EDS mapping, which exhibits a relatively homogeneous distribution of Ti, C, Co and N (Fig. 2e–i).

Based on the XRD pattern in Fig. 3a, pristine Ti<sub>3</sub>C<sub>2</sub> nanosheets exhibit three peaks at 8.8°, 27.9°, and 36.5°, corresponding to the (002), (004) and (101) facets of Ti<sub>3</sub>C<sub>2</sub>. Moreover, the presence of Ti<sub>3</sub>C<sub>2</sub> and CoPc diffraction signal peaks in the

CoPc/Ti<sub>3</sub>C<sub>2</sub> hybrid can be observed, inferring that the combination of CoPc molecules and Ti<sub>3</sub>C<sub>2</sub> does not alter their respective structures. Furthermore, X-ray photoelectron spectroscopy (XPS) was conducted to explore the elementary composition and electronic structure of these catalysts. According to the XPS survey spectra in Fig. 3b, the four elements of Ti, C, Co and N can be evidently observed with the binding energies of Ti 2p, C 1s, Co 2p and N 1s electrons, revealing that the CoPc/Ti<sub>3</sub>C<sub>2</sub> hybrid consists of CoPc and Ti<sub>3</sub>C<sub>2</sub>, which is in accordance with the XRD and EDS results. Besides, as for Ti, the peaks centered at 461.6 and 462.5 eV can be assigned to the 2p<sub>3/2</sub> and 2p<sub>1/2</sub> of Ti, respectively (Fig. 3c). The C 1s spectrum (Fig. 3d) contains three components located at 284.5 eV (C=C), 285.8 eV (C–C) and 288.3 eV (C–Ti), while for Co, two characteristic peaks centered at 782.1 and 797.8 eV of Co 2p<sub>3/2</sub> and Co 2p<sub>1/2</sub> are observed in Fig. 3e, respectively, suggesting that the valence state of Co is +2. In Fig. 3f, the N 1s peaks centered at 400.5 and 402.1 eV are assigned to the Co–N and aza N in the Pc



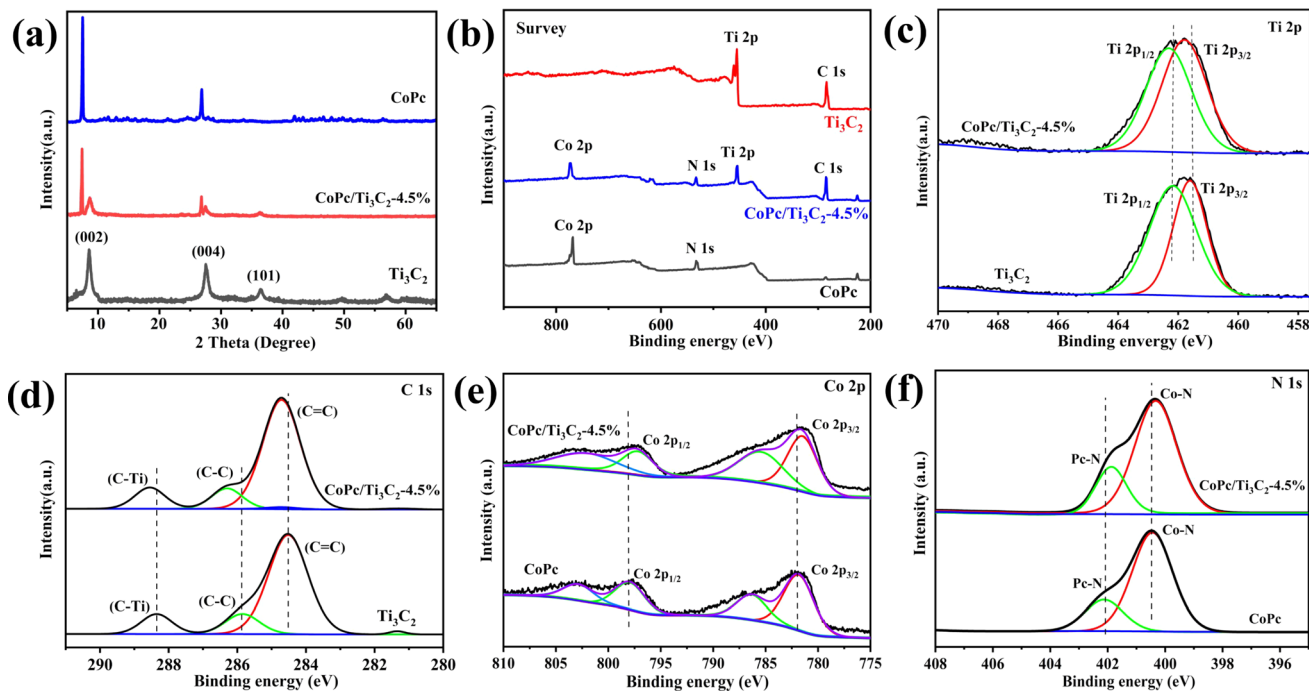


Fig. 3 (a) XRD patterns of as-prepared  $\text{Ti}_3\text{C}_2$ , CoPc and the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid. XPS survey spectrum (b), and high-resolution XPS spectra of Ti 2p (c), C 1s (d), Co 2p (e) and N 1s (f) of  $\text{Ti}_3\text{C}_2$ , CoPc and the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid.

moieties, respectively. It is interesting to note that the peaks of Ti 2p and C 1s for CoPc/ $\text{Ti}_3\text{C}_2$  shift to the higher energy direction compared with pristine  $\text{Ti}_3\text{C}_2$ , while the binding energies of

Co 2p and N 1s peaks of the composite are lower than that of pristine CoPc. Note that the electron binding energy of elements originates from the strong coulombic attraction between the

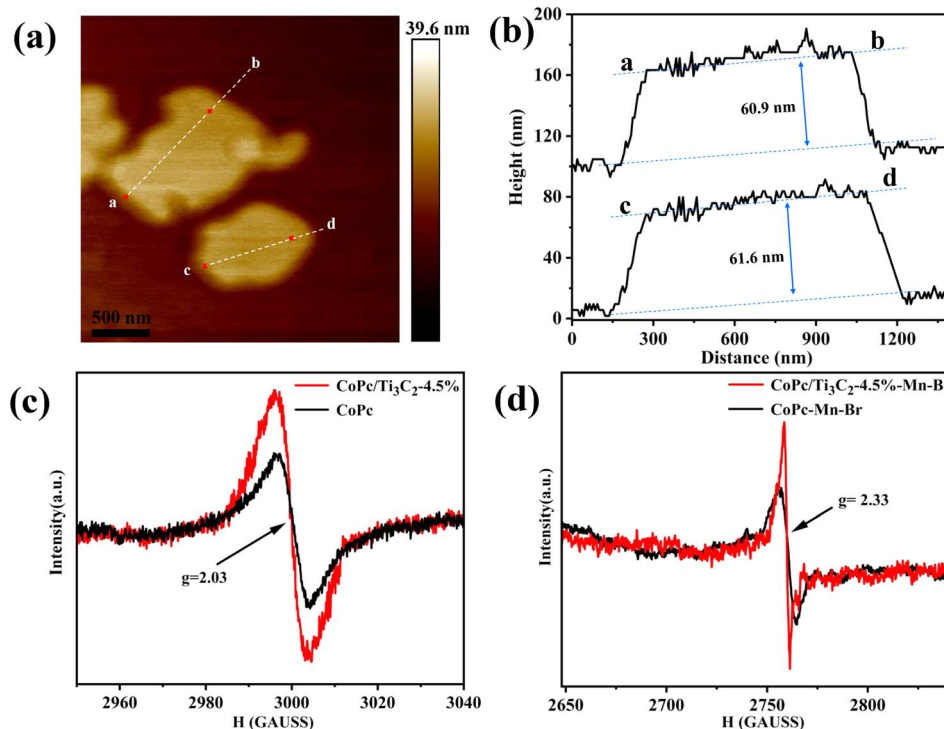


Fig. 4 (a) AFM images, and (b) height profiles from AFM analysis of the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid. EPR spectra of CoPc and the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid (c) without acetic acid as the solvent, (d) with acetic acid as the solvent during the reaction.



outer electrons of the atom and atomic nucleus. Meanwhile, the changes of element binding energy can directly reflect the changes of electron density. Therefore, these XPS results provide important evidence that CoPc molecules can be anchored on  $\text{Ti}_3\text{C}_2$  nanosheets *via* strong  $\pi$ - $\pi$  interactions, matching well with the fact that CoPc molecules are uniformly dispersed on  $\text{Ti}_3\text{C}_2$  nanosheets.

Moreover, based on the atomic force microscopy (AFM) images (Fig. 4a and b), the average statistical lateral size of the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid is about 1.2  $\mu\text{m}$ , and the thickness of the hybrid is about 61.3 nm, confirming its nanosheet structure. To reveal the crucial role of Co valence, we apply electron paramagnetic resonance (EPR) to acquire related information. Fig. 4c displays a relatively strong signal centered at  $g = 2.03$ , which can be attributed to a spin ( $S = 1/2$ )  $\text{Co}^{2+}$  component in both CoPc and CoPc/ $\text{Ti}_3\text{C}_2$ , which coincides well with the XPS results. As shown in Fig. 4d, prominent paramagnetic signals located at  $g = 2.33$  are observed, which can be assigned to the  $\text{Co}^{3+}$  species. Furthermore, significantly stronger signals shown at the same  $g$  value are observed for the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid than CoPc, suggesting the most favorable conversion of  $\text{Co}^{2+}$  to  $\text{Co}^{3+}$  in the CoPc/ $\text{Ti}_3\text{C}_2$ -Mn-Br catalyst system. This again indicates that the presence of the CoPc/ $\text{Ti}_3\text{C}_2$  structure benefits the formation of more  $\text{Co}^{2+}$ / $\text{Co}^{3+}$  during the reaction.

Nitrogen ( $\text{N}_2$ ) adsorption and desorption isotherms were collected to determine the BET surface area and porous structures of  $\text{Ti}_3\text{C}_2$  and the CoPc/ $\text{Ti}_3\text{C}_2$  composite. As shown in Fig. 5a, pristine  $\text{Ti}_3\text{C}_2$  displays a typical type IV isotherm and H3 hysteresis loop because of capillary condensation steps. The H3 hysteresis loop is also well preserved in the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid. The  $\text{N}_2$  adsorption capacity is improved, implying that the increase of  $\text{N}_2$  adsorption of the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid is attributed to that the CoPc/ $\text{Ti}_3\text{C}_2$  composite has a larger pore size and higher BET specific surface area ( $S_{\text{BET}}$ ) than  $\text{Ti}_3\text{C}_2$ . Besides, the  $S_{\text{BET}}$ 's, pore sizes, and pore volumes of prepared samples are summarized. It can be concluded that the CoPc/ $\text{Ti}_3\text{C}_2$  composite has led to a much larger specific surface area of 126.1  $\text{m}^2 \text{g}^{-1}$ , which is higher than that

of pristine  $\text{Ti}_3\text{C}_2$  (118.5  $\text{m}^2 \text{g}^{-1}$ ). The total pore volume ( $V_{\text{total}}$ ) increased after coupling, which suggests that the pores of  $\text{Ti}_3\text{C}_2$  are not choked by CoPc (Fig. 5b). The high  $S_{\text{BET}}$  and  $V_{\text{total}}$  are expected to create more active sites, facilitate mass transport and promote the utilization of active sites during the reaction.

### 3.2 Selective catalytic oxidation performance of 3,7-dichloro-8-dichloro methyl quinoline to quinclorac

Taking 3,7-dichloro-8-dichloro methyl quinoline (3,7-D-8-DMQ) as the substrate, the catalytic assay of selective oxidation was carried out in acetic acid solution (Fig. 6a). From the experimental results (Fig. 6b), the CoPc-Mn(OAc)<sub>2</sub>-HBr (CoPc-Mn-Br) catalyst produces less quinclorac with the quinclorac yield at 35.6%. However, the optimized CoPc/ $\text{Ti}_3\text{C}_2$ -4.5%-Mn-Br catalyst exhibits 3,7-D-8-DMQ conversion at 95.3%, high selectivity to quinclorac at 91.8% with superior quinclorac yield up to 87.5%. CoPc/ $\text{Ti}_3\text{C}_2$ -4.5%-Mn-Br displays a high catalytic yield for the production of high value-added quinclorac, which is 2.46-fold higher than that of the CoPc-Mn-Br system. These results show that the CoPc/ $\text{Ti}_3\text{C}_2$ -Mn-Br catalytic system is observably more active and robust compared to the CoPc-Mn-Br under the same conditions. Overall, the catalytic performances of the CoPc/ $\text{Ti}_3\text{C}_2$ -4.5%-Mn-Br catalyst are much better than that of the Co-based-Mn-Br catalyst (Table S1†). To the best of our knowledge, this is the first successful example of the CoPc/ $\text{Ti}_3\text{C}_2$ -4.5%-Mn-Br catalyst with excellent catalytic performance in the synthesis of quinclorac. To investigate the origin of the catalytic performance, a series of control experiments were conducted. As shown in Fig. S1,† the quinclorac yield of single CoPc/ $\text{Ti}_3\text{C}_2$ -4.5% or Mn(OAc)<sub>2</sub> or HBr cannot be found under the same conditions. Besides, it is imperative to explore the impact of Mn(OAc)<sub>2</sub> and HBr on the quinclorac yield. Significantly, there is no activity without HBr, suggesting that HBr functions as a chain initiator. The introduction of Mn(OAc)<sub>2</sub> in the CoPc/ $\text{Ti}_3\text{C}_2$ -4.5%-Br system can markedly improve the catalytic activity, indicating the cocatalyst role of Mn(OAc)<sub>2</sub>. For comparison, we performed a series of control experiments without different catalyst components under the

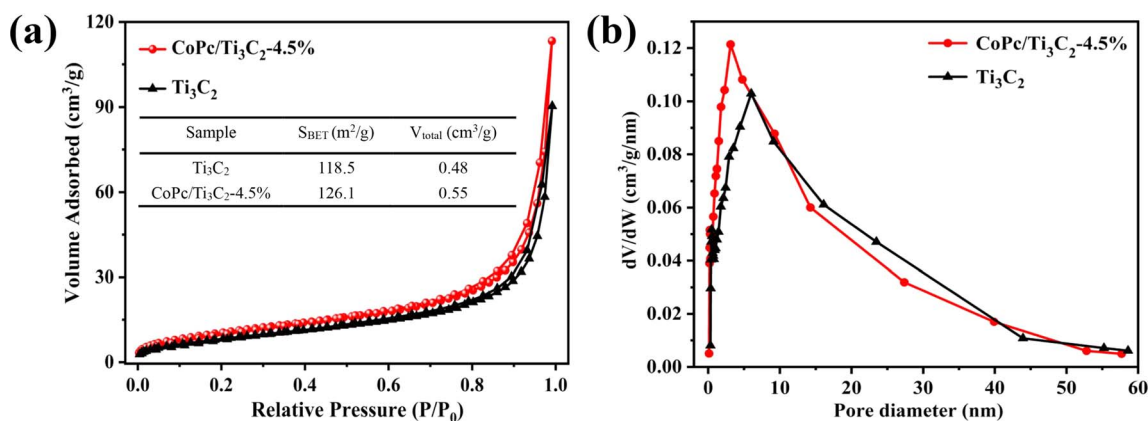


Fig. 5 (a) Nitrogen ( $\text{N}_2$ ) physisorption isotherms and the inset shows the corresponding BET specific surface area ( $S_{\text{BET}}$ ), total pore volume ( $V_{\text{total}}$ ), and (b) pore size distribution curves of  $\text{Ti}_3\text{C}_2$  and the CoPc/ $\text{Ti}_3\text{C}_2$  hybrid.



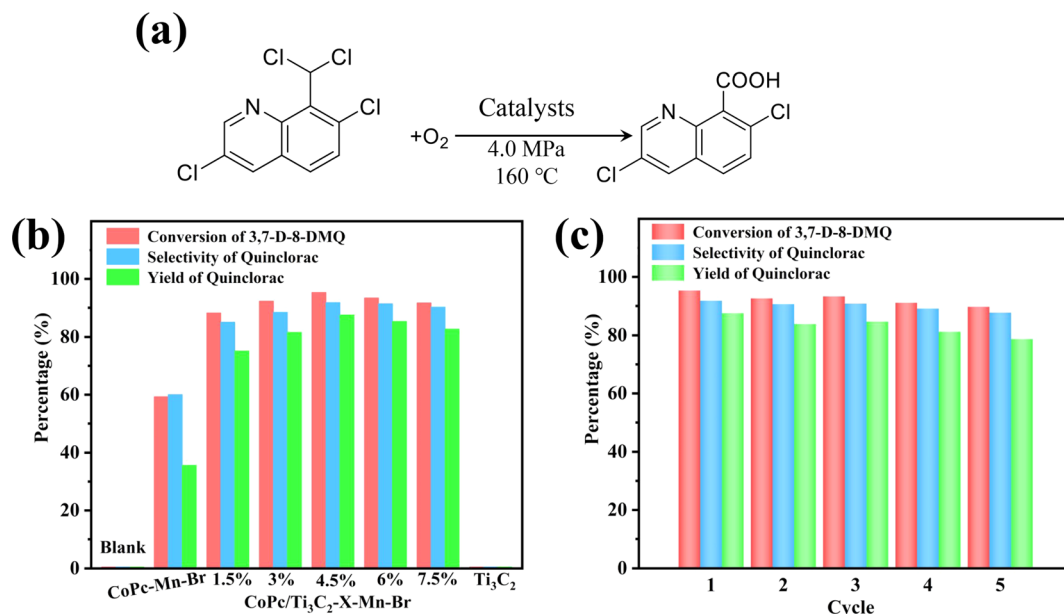


Fig. 6 (a) Chemical reactions for selective catalytic oxidation of 3,7-D-8-DMQ to quinclorac. (b) Conversion, selectivity and yield of catalytic oxidation reaction over CoPc/Ti<sub>3</sub>C<sub>2</sub>-Mn-Br and CoPc-Mn-Br after 6 h. (c) The stability tests over CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br.

same conditions. Moreover, the blank experiments without the presence of the cobalt source show a negligible 3,7-D-8-DMQ conversion, and almost no formation of quinclorac. The acetic acid solvent may have no direct effect on the reaction, and negligible products are obtained without any catalysts. These results confirm that the presence of the cobalt source is responsible for the conversion of 3,7-D-8-DMQ and yield of quinclorac.

The long-term stability of this catalytic system was investigated. The catalytic stability tests of the CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br are conducted under the optimum reaction conditions of 160 °C, O<sub>2</sub> partial pressure of 4 MPa and reaction time of 6 h for five consecutive cycles (Fig. 6c). After the reaction was completed, the catalysts could be recovered by simple filtration, and used for the next run. The catalytic performance shows very good stability during 30 hours of reaction. After five cycles, there is no observable change in the CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5% composite structure after the cyclic reaction (Fig. S2 and S3†). The as-designed heterogeneous catalyst reveals excellent stability and recyclability in the catalytic application.

### 3.3 Mechanistic investigations

To further investigate the mechanism, the variation of the metallic valence state was characterized by UV-vis spectra. As shown in Fig. 7a, the broad 560 nm band is a typical peak of an octahedral Co(II)-acetato complex. And the new band appearing at 690 nm is assigned to Co(III) complexes, which demonstrates that there is only Co<sup>2+</sup> before the reaction and Co<sup>2+</sup> is partly oxidized to Co<sup>3+</sup> during the reaction. However, for the CoPc-Mn-Br, the new bands corresponding to Co<sup>3+</sup> vanished after 5 h reaction. The Co<sup>3+</sup> cations generated previously are quickly reduced to Co<sup>2+</sup> by Mn<sup>2+</sup>, implying that the redox capability of cobaltous cations would change as the reaction intermediates

coordinated to the cobaltous cation instead of the acetic acid and bromic anions. As shown in Fig. 7b, the Co<sup>3+</sup> peak intensities of CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br are much stronger than that of CoPc-Mn-Br, suggesting that the oxidation from Co<sup>2+</sup> to Co<sup>3+</sup> is promoted in CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br, which is beneficial for the production of free radicals.

To gain insight into the transformation of intermediate species in oxidation reaction, a series of radical trapping experiments were conducted. We find that the typical free radical quenchers, 2,2,6,6-tetramethylpiperidoxyl (TEMPO) and 3,5-di-*tert*-4-butylhydroxytoluene (BHT), significantly suppress the formation of quinclorac (Fig. 7c). These results confirm that the catalytic reaction follows the free radical mechanism. To obtain more details of the chlorine species, a diallyl sulfonamide is submitted to the oxidation reaction of 3,7-D-8-DMQ with a stoichiometric amount of Bu<sub>4</sub>NCl. As expected, a cyclochlorinated compound is isolated, suggesting the formation of Cl<sup>•</sup> in the reaction (Fig. 7d and S4†). According to former literature, chlorine (Cl<sub>2</sub>) or Cl<sup>-</sup> might not be an active intermediate as the characteristic alkenyl dichlorination products are not observed. Thereafter, EPR studies using 5,5-dimethyl-1-pyrroline *N*-oxide (DMPO) as radical scavengers reveal the formation of radicals, as the coupling constants of the signal are consistent with related values of radicals, and the molecular weight is equal to the adduct of DMPO and radicals. As shown in Fig. 7e and f, the special signal with nitrogen hyperfine splitting  $\alpha_N = 15.36$  G and hydrogen hyperfine splitting  $\alpha_H = 27.81$  G is detected, which can be ascribed to DMPO-3,7-dichloroquinoline-CHO<sub>3</sub><sup>•</sup>. The signal peak intensity of CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br is significantly stronger than that of CoPc-Mn-Br. Consequently, with sufficient oxidative driving force from Co<sup>2+</sup> to Co<sup>3+</sup> for free radical activation and formation, CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br possesses more robust capacity to



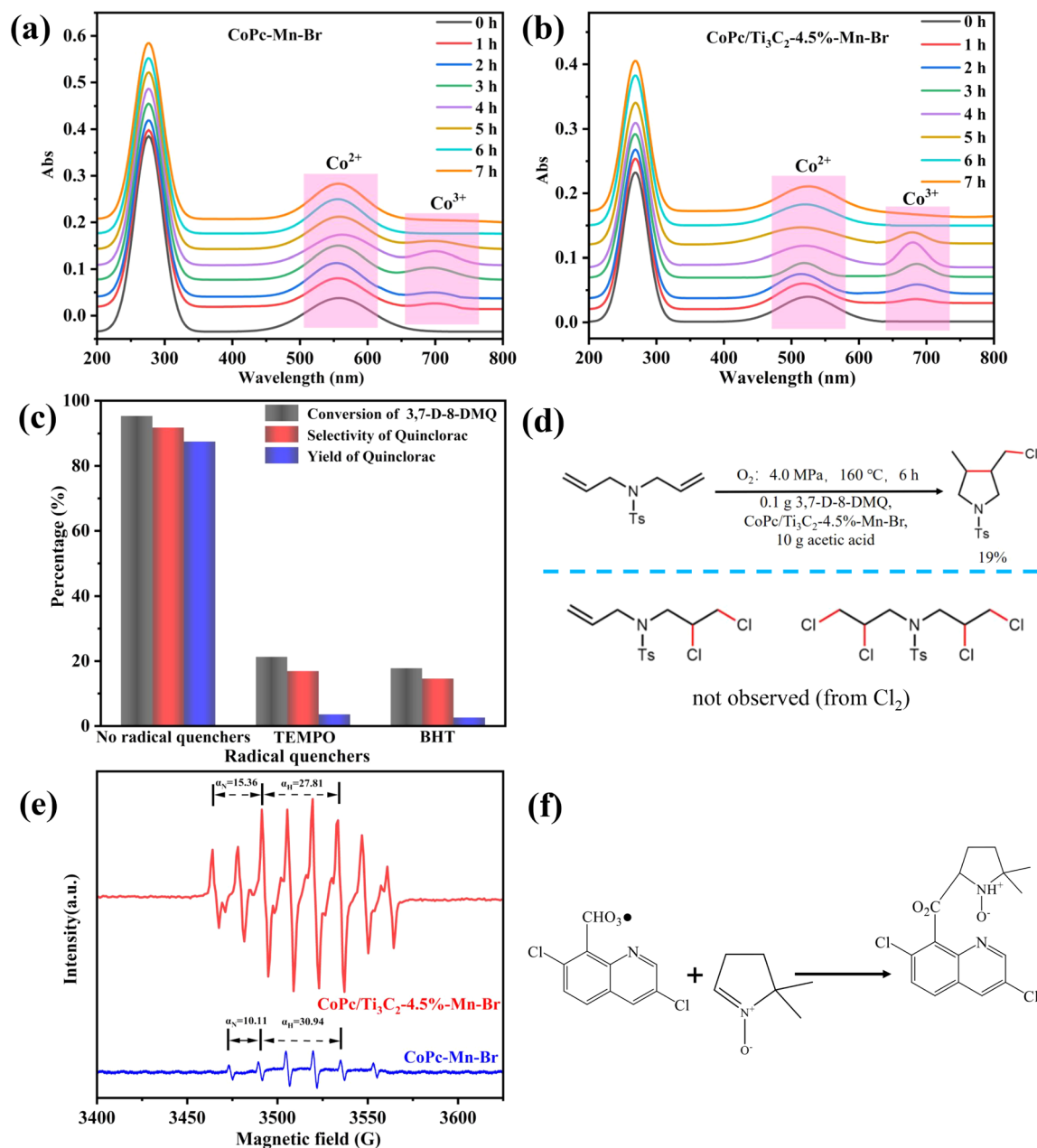


Fig. 7 The UV-vis spectra as a function of reaction time: (a) CoPc-Mn-Br and (b) CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br. (c) Radical quenching experiments of CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br. (d) Cl<sup>•</sup> trapping. (e) EPR spectra of CoPc-Mn-Br and CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br. (f) Chemical reactions for intermediate radicals and DMPO.

carry out the conversion of 3,7-D-8-DMQ to the key intermediate radicals as compared with CoPc-Mn-Br, which is consistent with previous results.

On the basis of the above experimental results and analyses, a plausible mechanism for the synthesis of quinclorac over the CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br catalyst is proposed (Fig. 8). Taken together, these experiments provide support for the surmised radical generation (Br<sup>•</sup> and Cl<sup>•</sup>) reaction pathways. Initially, Co<sup>2+</sup> and Mn<sup>2+</sup> are oxidized to Co<sup>3+</sup> and Mn<sup>3+</sup>. Co<sup>3+</sup> and Mn<sup>3+</sup> react with Br<sup>-</sup> ions to generate Br<sup>•</sup>. Br<sup>•</sup> as the initiator of free radical chain reactions reacts with the substrates, Co<sup>2+</sup> and

Mn<sup>2+</sup> in the presence of oxygen to produce 3,7-dichloro-8-aldehyde quinoline and Cl<sup>•</sup>. Then Co<sup>3+</sup> and Mn<sup>3+</sup> react with 3,7-dichloro-8-aldehyde quinoline to produce the key intermediate species – 3,7-dichloro-8-carbonyl quinoline free radicals. Subsequently, the 3,7-dichloro-8-peroxycarboxylic acid quinoline is generated sequentially under the action of oxygen, Co<sup>2+</sup> and Mn<sup>2+</sup>. 3,7-Dichloro-8-peroxycarboxylic acid quinoline would combine with the generated 3,7-dichloro-8-aldehyde quinoline to form the quinclorac product. More detailed mechanistic studies are still ongoing in our lab, and this



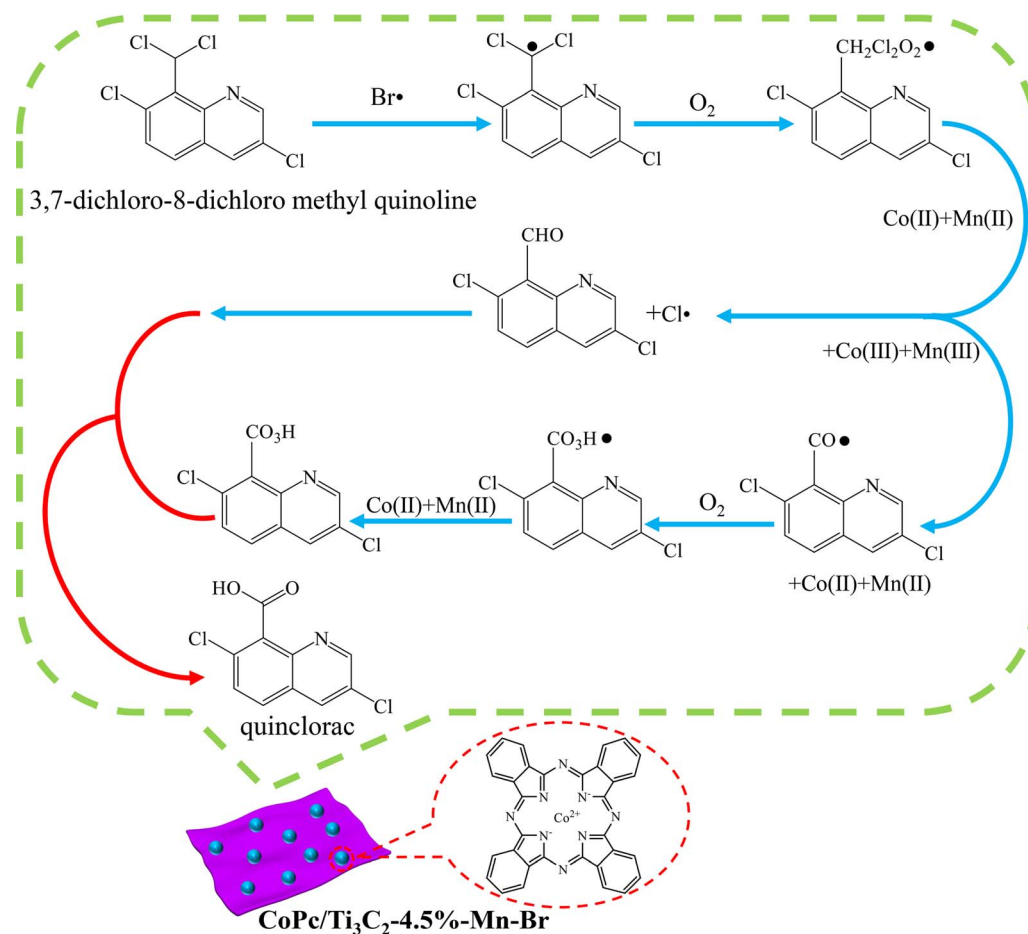


Fig. 8 Proposed mechanistic scheme of selective oxidation synthesis of quinclorac.

encouraging result could enlighten more multifunctional synergistic catalysis strategies for selective oxidation reactions.

## 4. Conclusions

In summary, we have successfully developed a non-noble metal, highly efficient and recyclable CoPc/Ti<sub>3</sub>C<sub>2</sub>-4.5%-Mn-Br catalyst system for selective oxidation of 3,7-D-8-DMQ to quinclorac. With this novel catalytic system, quinclorac can be directly obtained from 3,7-D-8-DMQ using O<sub>2</sub> as a convenient and clean oxygen source. It is worth noting that the Co oxidation states of the CoPc/Ti<sub>3</sub>C<sub>2</sub> hybrid play a vital role in the catalytic activity and stability during the reaction. Based on the exploration of intermediate radicals, we propose a reasonable catalytic mechanism. The excellent catalytic performance of introducing a CoPc/Ti<sub>3</sub>C<sub>2</sub> heterogeneous structure is attributed to the stronger capacity for stabilizing the accumulated redox cycle of Co<sup>2+</sup>/Co<sup>3+</sup>. This study provides a promising strategy for the development of highly effective selective catalytic systems. We believed that this work would advance the excited quinoline chemistry and open more green paths to future oxidation catalysts for large-scale and practical application.

## Author contributions

Simeng Zhu: investigation, formal analysis, validation and writing – original draft. Peng Liu: writing – review & editing and formal analysis. Xinlin Hong: supervision, project administration and resources.

## Conflicts of interest

There are no conflicts of interest to declare.

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## References

- Z. Chen, J. Song, X. Peng, S. Xi, J. Liu, W. Zhou, R. Li, R. Ge, C. Liu, H. Xu, X. Zhao, H. Li, X. Zhou, L. Wang, X. Li, L. Zhong, A. I. Rykov, J. Wang, M. J. Koh and K. P. Loh, *Adv. Mater.*, 2021, **33**, 2101382.



- 2 H. Hwang, J. Kim, J. Jeong and S. Chang, *J. Am. Chem. Soc.*, 2014, **136**, 10770–10776.
- 3 R. Sharma, R. Kumar, I. Kumar and U. Sharma, *Eur. J. Org. Chem.*, 2015, **2015**, 7519–7528.
- 4 Z. He, H. Liu, Q. Qian, L. Lu, W. Guo, L. Zhang and B. Han, *Sci. China: Chem.*, 2017, **60**, 927–933.
- 5 N. J. Pachupate and P. D. Vaidya, *J. Environ. Chem. Eng.*, 2018, **6**, 883–889.
- 6 P. L. Arias, J. A. Cecilia, I. Gandarias, J. Iglesias, M. López Granados, R. Mariscal, G. Morales, R. Moreno-Tost and P. Maireles-Torres, *Catal. Sci. Technol.*, 2020, **10**, 2721–2757.
- 7 X. Tian, L. Song, K. Farshadfar, M. Rudolph, F. Rominger, T. Oeser, A. Ariafard and A. S. K. Hashmi, *Angew. Chem., Int. Ed.*, 2019, **59**, 471–478.
- 8 A. K. Bains, V. Singh and D. Adhikari, *J. Org. Chem.*, 2020, **85**, 14971–14979.
- 9 P. E. Hande, M. Mishra, F. Ali, S. Kapoor, A. Datta and S. J. Gharpure, *ChemBioChem*, 2020, **21**, 1492–1498.
- 10 Z. Zhang, Z. Yuan, D. Tang, Y. Ren, K. Lv and B. Liu, *ChemSusChem*, 2014, **7**, 3496–3504.
- 11 S. J. Wang, Z. Wang, Y. Tang, J. Chen and L. Zhou, *Org. Lett.*, 2020, **22**, 8894–8898.
- 12 L. Qi, J. Chen, B. Zhang, R. Nie, Z. Qi, T. Kobayashi, Z. Bao, Q. Yang, Q. Ren, Q. Sun, Z. Zhang and W. Huang, *ACS Catal.*, 2020, **10**, 5707–5714.
- 13 B. Prabagar, Y. Yang and Z. Shi, *Chem. Soc. Rev.*, 2021, **50**, 11249–11269.
- 14 C. Teja and F. R. N. Khan, *Chem.–Asian J.*, 2020, **15**, 4153–4167.
- 15 N. A. Harry, S. M. Ujwaldev and G. Anilkumar, *Org. Biomol. Chem.*, 2020, **18**, 9775–9790.
- 16 Y. Kodama, S. Ariyasu, M. Karasawa, Y. Aiba and O. Shoji, *Catal. Sci. Technol.*, 2023, **13**, 6146–6152.
- 17 S. G. Makarov, S. Y. Ketkov and D. Wöhrle, *Chem. Commun.*, 2020, **56**, 5653–5656.
- 18 K. Trangwachirachai, I. T. Kao, W. H. Huang, C. L. Chen and Y. C. Lin, *Catal. Sci. Technol.*, 2023, **13**, 5248–5258.
- 19 Q. Huang, M. Zhao, Y. Yang, Y. N. Niu and X. F. Xia, *Org. Chem. Front.*, 2021, **8**, 5988–5993.
- 20 X. Deng, P. Zhao, X. Zhou and L. Bai, *Chem. Eng. J.*, 2021, **405**, 126979.
- 21 Z. Hou, L. Dai, Y. Liu, J. Deng, L. Jing, W. Pei, R. Gao, Y. Feng and H. Dai, *Appl. Catal., B*, 2021, **285**, 119844.
- 22 J. Xu, Y. Li, T. Ding and H. Guo, *Chem.–Asian J.*, 2021, **16**, 3114–3117.
- 23 W. An, S. H. Lee, D. Kim, H. Oh, S. Kim, Y. Byun, H. J. Kim, N. K. Mishra and I. S. Kim, *J. Org. Chem.*, 2021, **86**, 7579–7587.
- 24 D. Parmar, A. K. Dhiman, R. Kumar, A. K. Sharma and U. Sharma, *J. Org. Chem.*, 2022, **87**, 9069–9087.
- 25 J. Chen, Y. Guo, J. Chen, L. Song and L. Chen, *ChemCatChem*, 2014, **6**, 3174–3181.
- 26 Z. Hao, X. Zhou, Z. Ma, C. Zhang, Z. Han, J. Lin and G. L. Lu, *J. Org. Chem.*, 2022, **87**, 12596–12607.
- 27 N. Zargari, Y. Kim and K. Jung, *Green Chem.*, 2015, **17**, 2736–2740.
- 28 Y. Ding, Y. Huang, W. Sun, F. Li, S. Zhang, S. Fang, M. Wang and X. Hua, *Microchem. J.*, 2022, **183**, 107982.
- 29 Y. Ding, T. Guo, Z. Li, B. Zhang, F. E. Kühn, C. Liu, J. Zhang, D. Xu, M. Lei, T. Zhang and C. Li, *Angew. Chem., Int. Ed.*, 2022, **61**, e202206284.

