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# Manganese-oxide-supported gold catalyst derived from metal–organic frameworks for trace $\text{PCl}_3$ oxidation in an organic system†

Qianyi Zhao,  Qiang Geng  and Guoqiang Huang\*

Polysilicon is widely used in the field of semiconductors and solar energy. Trichlorosilane feedstocks that are used to produce polysilicon in the mainstream production process contain  $\text{PCl}_3$  impurities that have adverse effects on the quality of the polysilicon. Traditional methods for dephosphorization cannot achieve the effect of complete removal, whereas oxidizing  $\text{PCl}_3$  to  $\text{POCl}_3$  in the presence of oxygen for removal *via* adsorption is a promising and appealing route for establishing a dephosphorization process; it has a high phosphorous removal rate due to the strong Lewis-base property of  $\text{POCl}_3$  in comparison with  $\text{PCl}_3$ . In this work, we synthesized an active catalyst with an active interface between Au nanoparticles (NPs) and a manganese-oxide support ( $\text{Mn}_2\text{O}_3$ ) by calcination of a corresponding composite, where Au NPs were embedded uniformly in a metal–organic framework (MOF). The catalyst shows a significantly active catalytic performance for trace  $\text{PCl}_3$  oxidation in an organic system that is an imitation of a trichlorosilane system, with a 99.13% yield of  $\text{POCl}_3$  in an 80 °C and 0.6 MPa reaction environment. The structure–performance–mechanism analysis shows that the possible reaction and catalytic mechanism is  $\text{PCl}_3$  oxidation by interface lattice oxygens, which bridge the Au NPs and the support, in a Mars van Krevelen (MvK) process; this process was promoted by the interaction between the Au NPs and  $\text{Mn}_2\text{O}_3$  in terms of charge transfer and chemical potential changes. This work provides an effective way to dephosphorize trichlorosilane feedstocks in the polysilicon industry and gives guidance for constructing an efficient catalyst *via* the study of the structure and mechanism.

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## 1 Introduction

Phosphorus trichloride ( $\text{PCl}_3$ ) is widely present in the trichlorosilane feedstocks that are used to produce polysilicon in the industrial process *via* an improved Siemens method. In most factories, this method is used as a chief means of polysilicon production.<sup>1</sup> Crude trichlorosilane is obtained *via* a preliminary process and subsequently reduced by hydrogen to polysilicon in a large reduction furnace.<sup>2,3</sup> Existence of trace  $\text{PCl}_3$  in a trichlorosilane feedstock would cause P-doping on the polysilicon, which causes devastating damage to the resistivity and quality of the polysilicon product, on account of the valence shell of phosphorus having one extra electron; this would subsequently have a high probability of becoming a free electron when a silicon atom in the polysilicon crystal structure is replaced by a phosphorus atom.<sup>4–6</sup> Thus, phosphorus is called a donor impurity. According to the Chinese National Standard, electronic-grade polysilicon requires a donor impurity concentration of less than 0.15 ppb, while solar-grade polysilicon

requires there to be less than 1.40 ppb. The two standards are both strict for the purity of trichlorosilane feedstocks. The removal of  $\text{PCl}_3$  from trichlorosilane is therefore imperative.

To date, considerable efforts have been made in the field of trace  $\text{PCl}_3$  removal from trichlorosilane. At present, the methods of removing  $\text{PCl}_3$  in the industrial process mainly include distillation, adsorption and reactive distillation. Distillation makes use of the difference in relative volatility between components to make them separate from each other, and currently plays a dominant part in purification of trichlorosilane.<sup>7</sup> However, the boiling point of  $\text{PCl}_3$  is similar to that of trichlorosilane, such that huge energy consumption is needed in the distillation column to achieve the purification requirement, which will increase the cost and cause environmental pollution.<sup>8</sup> Adsorption and reactive distillation are relatively effective methods for  $\text{PCl}_3$  removal.<sup>9–11</sup> They take advantage of the Lewis base property of  $\text{PCl}_3$ , in that it easily coordinates with a Lewis acid to achieve  $\text{PCl}_3$  removal. Unfortunately, the Lewis base property of  $\text{PCl}_3$  is not very strong, which results in  $\text{PCl}_3$  being unable to interact with a Lewis acid strongly and form stable compound, finally causing an excessive reaction time and insufficient dephosphorization.<sup>12</sup> Part of pentavalent phosphorus compound such as  $\text{POCl}_3$  are strong Lewis base so they can be transformed into stable compound by

School of Chemical Engineering and Technology, Tianjin University, China. E-mail: hgg@tju.edu.cn

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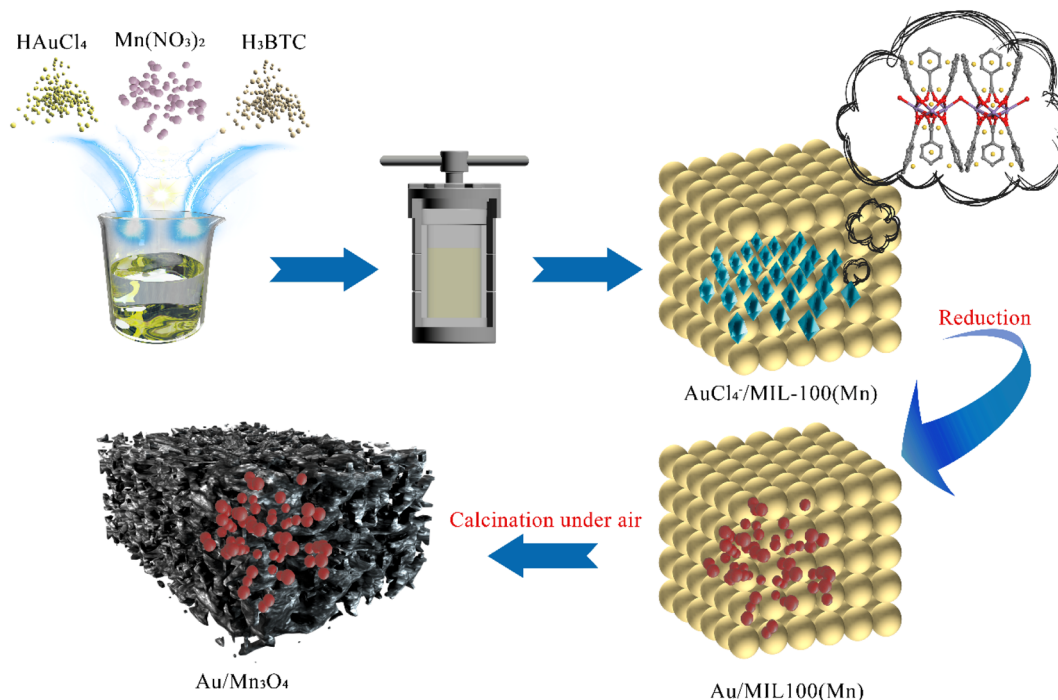


combined with Lewis acid and achieve the requirement of dephosphorization. Therefore, making use of oxidation–reduction reactions to turn  $\text{PCl}_3$  into a pentavalent phosphorus compound is highly desirable. Oxygen ( $\text{O}_2$ ) has been widely used as a relatively ideal oxidizing agent for industrial systems. However, the majority of studies of the  $\text{PCl}_3$  oxidation process under oxygen aim to produce  $\text{POCl}_3$ , such that the reactant  $\text{PCl}_3$  in these processes was present in large quantities, which is not in accordance with the situation of trace  $\text{PCl}_3$  removal from trichlorosilane.<sup>13</sup> Preliminary research has shown that trace  $\text{PCl}_3$  can hardly be oxidized to  $\text{POCl}_3$  under oxygen in the trichlorosilane system. In this situation,  $\text{POCl}_3$  can be formed from trace  $\text{PCl}_3$  and  $\text{O}_2$  in the trichlorosilane system when the reaction is catalyzed by diverse transition-metal oxides. Materials with  $\text{MoO}_3$ ,  $\text{Co}_3\text{O}_4$ ,  $\text{MnO}_2$  and  $\text{Mn}_3\text{O}_4$  as active components have been reported to be developed for catalyzing the oxidation–reduction reaction between  $\text{PCl}_3$  and  $\text{O}_2$  in the trichlorosilane system.<sup>14</sup> It has been confirmed in our preliminary experiment that among these transition-metal oxides,  $\text{Mn}_3\text{O}_4$  has the best catalytic activity. However, the oxidation of  $\text{PCl}_3$  catalyzed by manganese species alone did not exhibit excellent performance. In recent years, metal-oxide-supported Au catalysts that consist of Au nanoparticles (NPs) deposited on metal oxides have attracted enormous interest because of their high catalytic activity for various aerobic oxidation reactions.<sup>15–18</sup> Research has shown that catalysts such as  $\text{Au}/\text{CoO}_x$ ,  $\text{Au}/\text{Fe}_2\text{O}_3$ ,  $\text{Au}/\text{CeO}_2$  and  $\text{Au}/\text{TiO}_2$  exhibit remarkably enhanced activity for oxidation reactions in comparison with isolated Au particles and single metal-oxide supports.<sup>19–23</sup> It has been demonstrated that gold species loading on the support promotes the interaction

between the Au and support, which leads to enhanced reduction of surface lattice oxygen and the activation of gaseous oxygen, thus contributing to reactant oxidation.<sup>22,24,25</sup> Based on the above discoveries, it is logical to assume that catalysts with high activity toward the oxidation of  $\text{PCl}_3$  could be created by supporting Au NPs on the reducible metal oxide  $\text{Mn}_3\text{O}_4$ .

To obtain a catalyst with superior catalytic performance, the selection of the synthesis method is consequently a key step. Common supported-Au catalyst preparation methods mainly include impregnation, deposition–precipitation, and sol-immobilization.<sup>26–30</sup> But generating catalysts with a uniform distribution of Au NPs and strong interactions between the Au and support, which is beneficial to the catalytic activity, is not straightforward when using these traditional methods.<sup>31–33</sup> Liao *et al.* engineered an active catalyst for the CO oxidation reaction with an even distribution of Au–Pd alloy on the support and an active interface between the metal and cobalt oxide supports *via* calcination of a composite of NPs encapsulated in a metal–organic framework (MOF).<sup>23,34</sup> This method provides an effective way to synthesize a catalyst with an even distribution of metal NPs on the metal-oxide support and an active interface between the metal and support material for improved catalytic performance.<sup>35</sup>

Motivated by the above, a  $\text{PCl}_3$ -oxidation catalyst with high catalytic activity was prepared in this study, wherein Au NPs were loaded on a  $\text{Mn}_3\text{O}_4$  support *via* the synthetic strategy developed by Liao and co-workers. In this strategy, the metal ions used as a metal-NP precursor were encapsulated into the pores of a MOF during the self-assembly process of the MOF, and subsequently reduced to metal NPs embedded in the



**Scheme 1** Synthesis of the Mn-MIL-100-derived Au-decorated manganese-oxide support applied for trace  $\text{PCl}_3$  oxidation under oxygen in an organic system.

framework of the MOF. The metal-oxide support is formed through the calcination of the MOF composite, which consists of the metal-organic framework and metal NPs homogeneously distributed on it. In this work, as graphically illustrated in Scheme 1, Au ions are trapped in the framework of Mn-MIL-100 (MIL stands for Matériaux de l'Institut Lavoisier), which is based on a trimeric building unit composed of manganese metal nodes and a trimesic acid ( $\text{H}_3\text{BTC}$ ) linker.<sup>36</sup> Then the Au NPs are formed in the framework of Mn-MIL-100 by reducing the Au ions. The MOF composites were subsequently calcined under air and converted into the final structure of Au NPs embedded in manganese oxide ( $\text{Mn}_3\text{O}_4$ ) cages ( $\text{Au}_x/\text{Mn}_3\text{O}_4$ ;  $x$ ) represents the mass fraction of Au, ( $x$ ) = 4.13–10.52).

In the presence of  $\text{O}_2$ , trace  $\text{PCl}_3$  oxidation over the  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalyst was carried out under different conditions to observe the catalytic activity. Considering the toxicity and corrosivity of trichlorosilane, the reaction was conducted in an organic system consisting of  $\text{PCl}_3$  and *n*-hexane, which was chosen instead of trichlorosilane due to the similarity in physical properties between trichlorosilane and *n*-hexane. As expected, the  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalyst exhibits enhanced catalytic performance in comparison with just  $\text{Mn}_3\text{O}_4$ . Au NPs have thus been demonstrated to be an effective loaded metal to facilitate the catalytic activity of the single metal oxide. In order to optimize the catalytic activity of the  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalyst and determine the optimal reaction conditions, a systematic study towards the influence of reaction and catalyst parameters, such as the mass fraction of Au in the  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalyst, reaction temperature, reaction pressure and catalyst dosage, on the conversion rates of  $\text{PCl}_3$  was performed. For the purpose of understanding the effects of the catalyst properties on the catalytic activities and investigating the catalytic mechanism of  $\text{PCl}_3$  oxidation on the  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalyst, a thorough characterization of the  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalyst and its precursors and a DFT study of the mechanism of  $\text{PCl}_3$  oxidation on the  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalyst has also been carried out.

Herein, an  $\text{Au}/\text{Mn}_3\text{O}_4$  catalyst was synthesized, derived from a metal-organic framework, and has significantly active catalytic performance for  $\text{PCl}_3$  oxidation in an organic system. The physicochemical characterization of the as-prepared catalyst was investigated in detail, revealing the structure–performance relationships between the  $\text{POCl}_3$  yield and the electronic and geometric effects contributed by the Au NPs. Meanwhile, the optimal process conditions and reaction mechanism were studied for better industrial applications.

## 2 Materials and methods

### 2.1 Materials

Trimesic acid ( $\text{H}_3\text{BTC}$ ), lauric acid and chloroauric acid were purchased from Heowns. Manganese(II) nitrate tetrahydrate ( $\text{Mn}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ ), sodium borohydride, *n*-hexane and methanol were purchased from Aladdin. Ammonium molybdate, vitamin C, L-antimony potassium tartrate and sulfuric acid, used to detect the content of pentavalent phosphorus, were purchased from Macklin.  $\text{PCl}_3$  was purchased from Adamas-Beta. Pure oxygen was supplied by Air Liquid.

### 2.2 Synthesis of $\text{Au}_x/\text{Mn}_3\text{O}_4$ catalysts

Mn-MIL-100 was synthesized *via* a modified solvothermal method.<sup>37</sup> Briefly, 0.72 mmol of  $\text{Mn}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ , 2.85 mmol of  $\text{H}_3\text{BTC}$  and 13.5 mmol lauric acid were dissolved in 75 mL of methanol at a stirring speed of 300 rpm for 40 min. The as-prepared solution was then transferred into an autoclave (100 mL) and maintained at 125 °C for 10 h. The resultant brown suspension was then centrifuged at 12 000 rpm and the sediment was collected. The collected particles were subsequently washed with methanol 2 times, followed by vacuum drying at 60 °C for 4 h. The manganese-oxide cages used as the control group were produced by calcination of Mn-MIL-100 under air for 4 h.

For the  $\text{Au}/\text{Mn-MIL-100}$  composites, in which Au NPs are embedded in the framework of Mn-MIL-100, various masses of chloroauric acid were melted into the manganese(II) nitrate tetrahydrate-containing solution at a stirring speed of 300 rpm for 40 min. The as-prepared solution was then transferred into an autoclave (100 mL) and maintained at 125 °C for 10 h. The resultant brown suspension was then centrifuged at 12 000 rpm and the sediment was collected. The collected particles were washed with methanol 2 times, then re-suspended in 60 mL of methanol. 0.2 g of sodium borohydride was added into the brown suspension and stirring was maintained for 2 h at room temperature. The suspension was centrifuged and the collected particles were washed with methanol twice, followed by vacuum drying for further use. The acquired solid powder was further calcined under air at 500 °C to obtain catalysts supported by manganese oxide.

### 2.3 Characterization

The morphology of all samples was observed through scanning electron microscopy (SEM) conducted on an FEI Apreo S instrument. The X-ray diffraction (XRD) patterns were acquired using a Rigaku SmartLab SE X-ray diffractometer.  $\text{N}_2$  adsorption–desorption isotherms were measured at −196 °C with a Micromeritics ASAP2460 sorption analyzer. The Au content measurement of all samples was performed using inductively coupled plasma atomic emission spectroscopy (ICP-AES) on a Agilent 725 instrument. X-ray photoelectron spectroscopy (XPS) was performed using a Thermo ESCA LAB Xi+ spectrometer to analyze the state of the gold and manganese species in the samples, and the results were calibrated using the C 1s peak at 284.8 eV. Fourier-transform infrared (FT-IR) spectra were collected on a Thermo Nicolet iS20 spectrometer. Thermogravimetric analysis (TGA) of the samples, at temperatures ranging from 25–800 °C, was performed using a Henven HTG-4 analyzer.

### 2.4 $\text{PCl}_3$ oxidation over $\text{Au}_x/\text{Mn}_3\text{O}_4$

The oxidation of trace  $\text{PCl}_3$  under an oxygen environment in an organic system was carried out in a 100 mL autoclave. A trace amount of  $\text{PCl}_3$  was injected into 100 mL of *n*-hexane under nitrogen to obtain the reaction liquid. Then a given amount of reaction liquid and catalyst were added into the batch reactor. After purging with oxygen three times, the reactor was



pressurized to a specific pressure using oxygen and then heated at a desired temperature under suitable stirring. After a set reaction time, the reaction was finished and the reactor was cooled down to room temperature. The liquid in the reactor after reaction was then subjected to ultrasonic treatment for 20 min and the solid catalyst was removed from the system by centrifugation. 1 mL of the residual liquid was then taken and extracted using 100 mL of deionized water. The chromogenic agent was made up of sulfuric acid solution (5 mol L<sup>-1</sup>), 3 wt% ammonium molybdate solution, 5.4 wt% vitamin C solution and 0.136 wt% L-antimony potassium tartrate solution with a volume ratio of 3 : 2 : 2 : 1. The content of phosphoric acid in aqueous phase formed from the hydrolyzation of POCl<sub>3</sub> in the above residual liquid was determined by spectrophotometry, using ammonium molybdate as the active component in the chromogenic agent, on a ultraviolet spectrophotometer (BINGLIN, λ = 891 nm). The POCl<sub>3</sub> content in the system after the reaction was quantified using an ammonium molybdate standard curve and the corresponding conversion relation of the concentration. The yield of POCl<sub>3</sub> was defined as the moles of POCl<sub>3</sub> generated after reaction divided by the moles of PCl<sub>3</sub> added initially, as shown in eqn (1):

$$\text{Yield (\%)} = \frac{\text{moles of POCl}_3 \text{ generated}}{\text{moles of PCl}_3 \text{ in feed}} \times 100 \quad (1)$$

## 2.5 Computational details

All the calculations are performed in the framework of density functional theory (DFT) with the projector augmented plane-wave method, as implemented in the Vienna *ab initio* simulation package.<sup>38</sup> The generalized gradient approximation proposed by Perdew, Burke, and Ernzerhof is selected for the exchange–correlation potential.<sup>39</sup> The long-range van der Waals interaction is described by the DFT-D3 approach.<sup>40</sup> The cut-off energy for the plane wave is set to 450 eV. The energy criterion is set to 10<sup>-5</sup> eV in the iterative solution of the Kohn–Sham

equation. A vacuum layer of 15 Å is added perpendicular to the sheet to avoid artificial interaction between the periodic images. The Brillouin zones for energy calculation of Mn<sub>2</sub>O<sub>3</sub> and Au<sub>10</sub>@Mn<sub>2</sub>O<sub>3</sub> are sampled with a Gamma-centered *k*-mesh of 3 × 3 × 1. All the structures are relaxed until the residual forces on the atoms have declined to less than 0.03 eV Å<sup>-1</sup>.

## 3 Results and discussion

### 3.1 Characterization of Au<sub>x</sub>/Mn<sub>3</sub>O<sub>4</sub> catalysts

Different weights of the Au precursor were added into the system during the process of Mn-MIL-100 synthesis, simultaneously to the trimesic acid, manganese nitrate, and lauric acid selected as the precursors, so that the Au precursor could be encapsulated into the pores of Mn-MIL-100. Then the Au precursor was reduced into metal NPs by addition of NaBH<sub>4</sub>, forming MOF composites in which Au NPs were embedded in the framework of Mn-MIL-100 (Au<sub>x</sub>/MIL-100; (*x*) represents the mass fraction of Au). As shown in Fig. S1,† Mn-MIL-100 consists of smooth and uniform octahedral particles with an average size of *ca.* 500 nm, matching well with its trimeric building units. Scanning electron microscopy (SEM) images of Au/Mn-MIL-100 are shown in Fig. 1A, showing that the structure of Mn-MIL-100 turns out to be slightly collapsed once the Au NPs are embedded in it and Au/Mn-MIL-100 exists in an irregular octahedron configuration, which indicates that the loading of Au NPs causes minor disruption to the structure of Mn-MIL-100. Fig. 1B shows the element distributions of C, O, Mn, and Au species *via* scanning electron microscopy with energy dispersive X-ray spectroscopy (SEM-EDX) of Au/Mn-MIL-100, confirming the successful loading and even distribution of Au species inside the Mn-MIL-100.

Fig. 2A shows the X-ray diffraction (XRD) pattern of Au/Mn-MIL-100, which is in accordance with that of Mn-MIL-100 to a great extent, but the peak intensities decrease with the enhancement in the Au mass fraction, verifying that Au NPs do have some effect on the octahedral configuration of Mn-MIL-

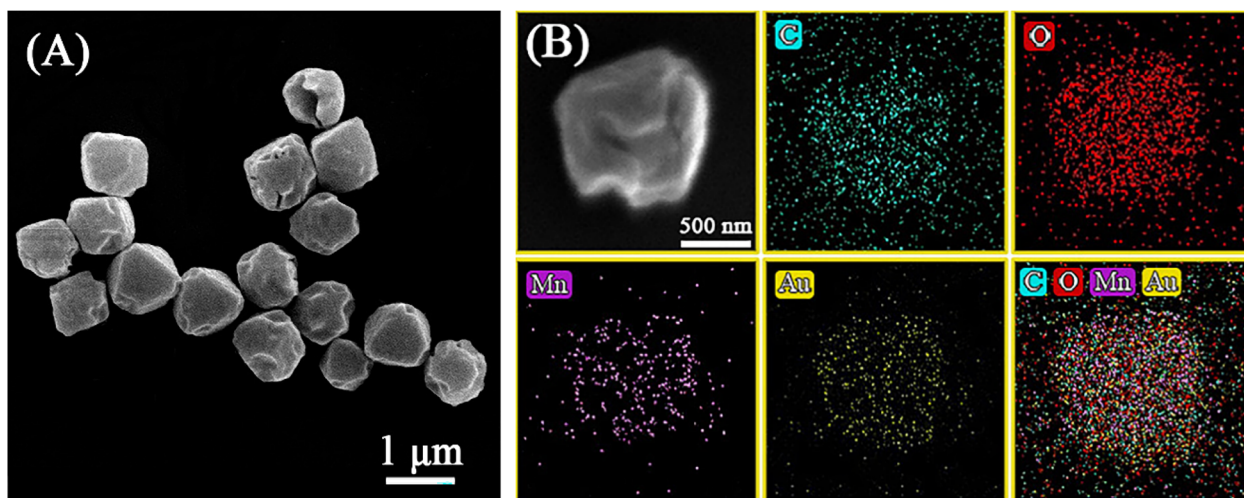


Fig. 1 (A) SEM image, and (B) element distributions of C, O, Mn and Au species *via* SEM-EDX of Au/Mn-MIL-100.



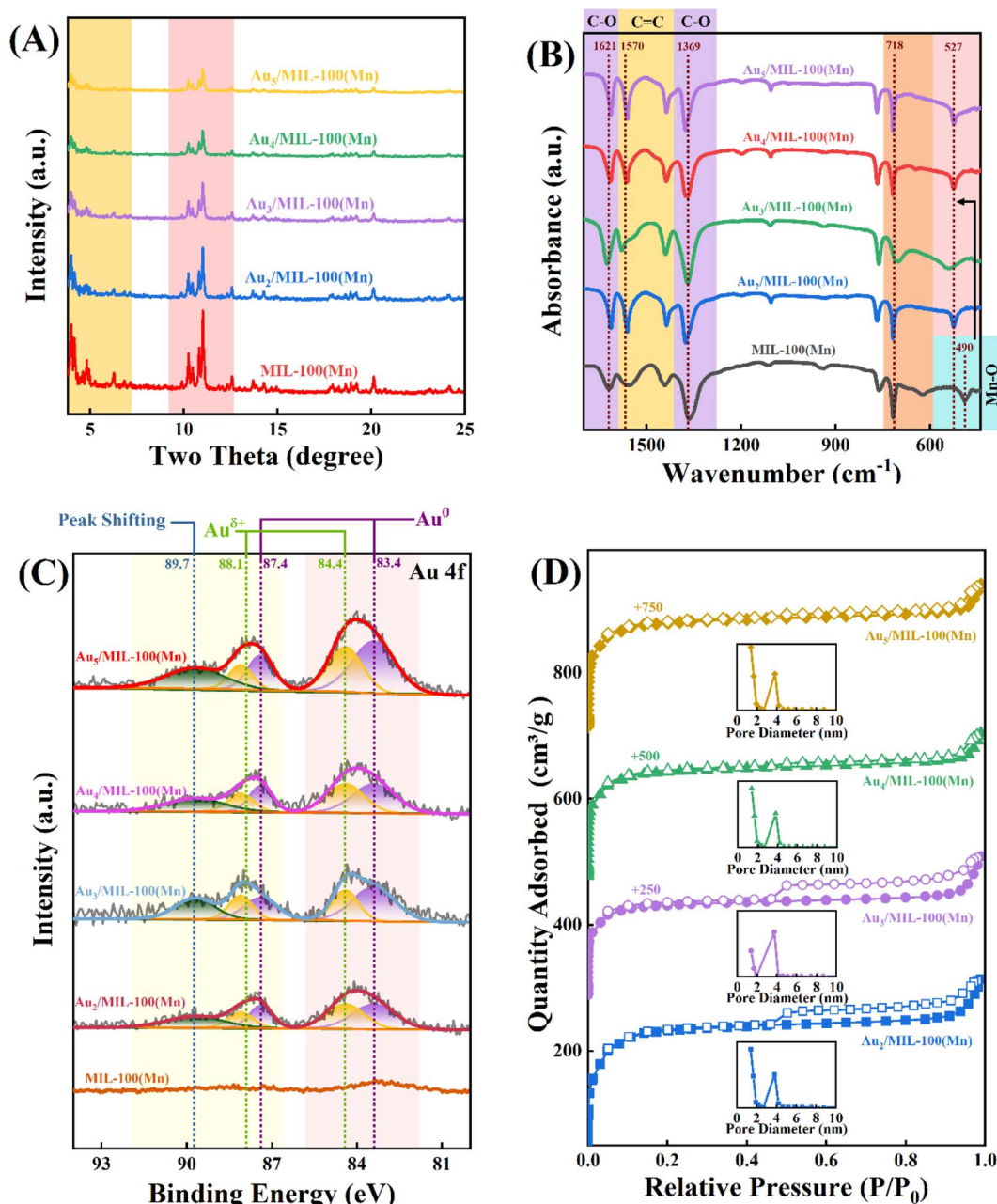


Fig. 2 (A) XRD patterns, (B) FT-IR spectra, (C) Au 4f XPS spectra, and (D) N<sub>2</sub> adsorption–desorption isotherms (with inset pore-size distributions) of Au/Mn-MIL-100.

100. The disruption is aggravated with the enhancement in the Au mass fraction, but the as-prepared Au/Mn-MIL-100 basically maintained the well-constructed framework of MIL-100, with metal NPs embedded in it. The electronic properties of the Au element were explored *via* X-ray photoelectron spectroscopy (XPS), as exhibited in Fig. 2C. The Au-free Mn-MIL-100 sample shows no Au-related signal in Au 4f XPS, as expected, while obvious bands are observed for the four Au/Mn-MIL-100 samples, whose intensities progressively increase with the rise in the Au mass fraction. Considering the complicated micro-environment of Au species and the properties of Au 4f, each spectrum was deconvoluted into five peaks at 83.4, 84.4, 87.4,

88.1 and 89.7 eV. The subordinate peaks at 83.4 and 87.4 eV are associated with the 4f<sub>7/2</sub> and 4f<sub>5/2</sub> photoelectrons of Au<sup>0</sup> with a zero valence state and the peaks at 84.4 and 88.1 eV are attributed to Au<sup>δ+</sup> with a positive charge, further proving that Au NPs were successfully embedded in the framework of Mn-MIL-100 and an interaction was simultaneously generated in terms of charge transfer between the Au NPs and Mn-MIL-100.<sup>41–44</sup> In addition, the peak at 89.7 eV exhibits a slight shift to higher energy compared to the characteristic peaks of Au<sup>δ+</sup> and Au<sup>0</sup>; this was ascribed to Au\*, which represents Au species that replaced a small portion of the metal sites in the trimeric building units of Mn-MIL-100 during the self-assembly process



between Mn ions and H<sub>3</sub>BTC. The replacement results in sparsity of the electron cloud density around these Au species owing to the higher electronegativity of the O element compared to that of the Au element and leads to the higher binding energy of Au\*.<sup>45</sup> The Au\*/(Au\* + Au<sup>0</sup> + Au<sup>δ+</sup>) average molar ratio (0.14) is relatively low, indicating that the damage to the structure of Mn-MIL-100 owing to the exchange of Au ions for metal sites during the self-assembly process occurs to a very low degree. Fig. 2B shows the Fourier-transform infrared (FTIR) spectra of Au/Mn-MIL-100. The band at 490 cm<sup>-1</sup>, which could be associated with the marked Mn–O band in Mn-MIL-100, is shifted to a higher wavenumber (*i.e.*, 527 cm<sup>-1</sup>) for the Au/Mn-MIL-100 materials.<sup>46</sup> This is attributed to free electrons, which were lost by Au species in the process of transition from Au<sup>0</sup> to Au<sup>δ+</sup>, moving around the Mn–O bond and increasing its stretching vibration frequency. The peaks at 718 cm<sup>-1</sup> are assigned to the benzene ring of the ligand, while the peaks at 1369 and 1621 cm<sup>-1</sup> are ascribed to the C–O stretching of the BTC anions.<sup>36,47</sup> The peaks centered at 1570 cm<sup>-1</sup> can be attributed to the C=C stretching of benzene.<sup>48</sup> The N<sub>2</sub> adsorption/desorption isotherms of Au/Mn-MIL-100 are exhibited in Fig. 2D, displaying typical type I isotherms, which could be attributed to the micropores of Mn-MIL-100. An H4 hysteresis loop could also be observed in some of the isotherms, revealing the presence of internal voids inside the Mn-MIL-100. One of the possible causes for the formation of these internal voids is the substitution of Mn ions in Mn-MIL-100 by Au ions, as mentioned above. The trimeric units that Mn-MIL-100 was

based on were constructed through coordination bonds between the d<sup>2</sup>sp<sup>3</sup> hybrid orbitals of the Mn ions and the lone pairs of the six O atoms supplied by H<sub>3</sub>BTC. Since the orbital hybridization mode of the Au ions is distinct from that of the Mn ions, some of the trimeric units would collapse and be transformed to the coordination structure between the Au ions and O atoms, leading to irregular pore construction and internal voids in Au/Mn-MIL-100. Furthermore, Pang *et al* indicated that the reduction of chloroauric acid would generate a large number of hydrogen ions and expose Au/Mn-MIL-100 to an acidic environment.<sup>49</sup> Liao *et al.* figured out that another cause for the formation of internal voids is that metal ligands such as H<sub>3</sub>BTC would protonate rather than coordinate with Mn ions in an acidic environment; Mn ions located in the core of Mn-MIL-100 consequently move outward to the environment and compensate the coordinated Mn ions located in the shell of Mn-MIL-100, which promotes the formation of voids.<sup>23</sup>

Au/Mn<sub>3</sub>O<sub>4</sub> catalysts were obtained through further calcination of Au/Mn-MIL-100 under air. As shown in Fig. 3A, the morphology evolution during pyrolysis was evaluated *via* SEM images for different calcination temperatures. When the temperature reaches 400 °C, the MOF structure of Au/Mn-MIL-100 converts to small clusters without a fixed form. When the temperature continues to increase, blocky clusters of porous structure, with an average particle size of 500 nm, start to appear at 500 °C and further transform into big clusters at 600 °C and 700 °C. The size of the clusters calcined at 600 °C was five times larger than that of clusters calcined at 500 °C, while the

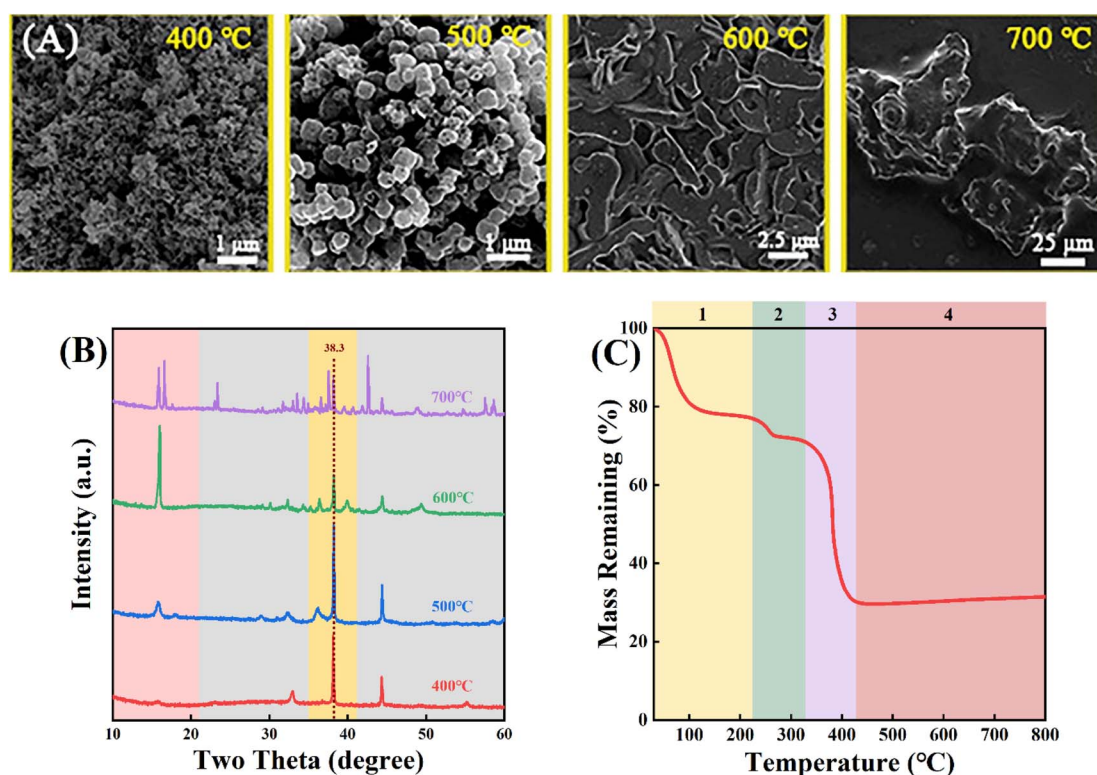


Fig. 3 (A) SEM images and (B) XRD patterns of Au<sub>2</sub>/Mn-MIL-100 calcined at different temperatures (400, 500, 600 and 700 °C) under air. (C) Weight loss of Au<sub>2</sub>/Mn-MIL-100 calcined under air.



size of the clusters calcined at 700 °C was about forty times larger than that of clusters calcined at 600 °C, indicating that the materials would gradually aggregate when the calcination temperature rises from 500 to 700 °C. To fully recognize the chemical reactions and phase transitions involved during the pyrolysis of Au/Mn-MIL-100, thermogravimetric analysis-mass spectrometry (TGA-MS) was conducted under air. As shown in Fig. 3C, the pyrolysis process in the temperature range from 0–800 °C could be divided into four phases: (i) ethanol that permeated into the pores of Au/Mn-MIL-100 evaporates (0 °C <  $T$  < 220 °C); (ii) the H<sub>3</sub>BTC ligand loses some of its benzene rings, leaving the Mn atom bonded to the two oxygen atoms of a carboxy group (220 °C <  $T$  < 320 °C); (iii) the H<sub>3</sub>BTC ligand loses all of its benzene rings, leading to bond-breaking between Mn and O and the release of CO<sub>2</sub> (320 °C <  $T$  < 430 °C); (iv) Au/Mn<sub>3</sub>O<sub>4</sub> compounds form, which are stable within 800 °C.<sup>50</sup> The XRD results of the Au/Mn<sub>3</sub>O<sub>4</sub> materials obtained by calcination at different temperatures are shown in Fig. 3B. The intensity of the peak at 38.3° that is ascribed to Au NPs first increases and then decreases with the rise in calcination temperature, revealing an increase in size of the Au particles with the calcination temperature increasing from 500 to 600 °C; then the Au particles and Mn<sub>3</sub>O<sub>4</sub> support were gradually encapsulated in a carbon matrix when the temperature continued to increase. Thus, different calcination temperatures were used to optimize the interface between the metal NPs and oxide with 500 °C being selected for further testing.

The mass fraction of Au( $x$ ) in the calcined sample was quantified *via* ICP-MS. Fig. 4A shows SEM images of the as-prepared Au<sub>4.13</sub>/Mn<sub>3</sub>O<sub>4</sub> calcined at 500 °C. The elemental distribution of O, Mn, and Au species (Fig. 4B) shows overlap of the four species, indicating the formation of manganese oxide and a uniform distribution of Au NPs on the Mn<sub>3</sub>O<sub>4</sub> support. In addition, the sparse elemental mapping of C shows that most of the C element from the H<sub>3</sub>BTC linker escaped in the form of CO<sub>2</sub> during the pyrolysis process, revealing that the quantity of carbon matrix that the catalysts prepared by calcination at 500 °C were embedded in was at a relatively low level. To confirm the advantages of the method for the preparation of Au<sub>*x*</sub>/Mn<sub>3</sub>O<sub>4</sub>

catalysts derived from Mn-MIL-100, the Au-Mn<sub>3</sub>O<sub>4</sub> material was synthesized *via* an impregnation method for comparison. The SEM images of the Au-Mn<sub>3</sub>O<sub>4</sub> (Fig. S2A†) show that its morphology is particles with a relatively large size, and the corresponding selected area element distribution analysis (Fig. S2B†) shows a situation of local accumulation of the Au element, indicating that the Au element was dispersed on the material unevenly, further demonstrating that the synthesis method using MOFs as a precursor could promote distribution uniformity. The XRD patterns of Au<sub>*x*</sub>/Mn<sub>3</sub>O<sub>4</sub> in Fig. 5A show that the diffraction peaks of Au/Mn<sub>3</sub>O<sub>4</sub> derived from Mn-MIL-100 are in accordance with the standard values of Mn<sub>3</sub>O<sub>4</sub> (JCPDS No. 24-0734) and the intensity of the characteristic Au peak located at 38.3° progressively increases with the rise in Au mass fraction, further confirming the successful synthesis of the Mn<sub>3</sub>O<sub>4</sub> support and the increase in size of the Au NPs embedded in the metal oxide with the increase in the Au mass fraction.<sup>51</sup> The N<sub>2</sub> adsorption-desorption isotherm (Fig. 5B) can be designated as a type-II isotherm, revealing a mesoporous and macroporous structure of the as-prepared Au<sub>*x*</sub>/Mn<sub>3</sub>O<sub>4</sub> catalysts. The specific areas, total pore volumes and pore sizes of the Au<sub>*x*</sub>/Mn<sub>3</sub>O<sub>4</sub> catalysts are summarized in Table 1. The average specific surface area of Au<sub>*x*</sub>/Mn<sub>3</sub>O<sub>4</sub> is 8.0 m<sup>2</sup> g<sup>−1</sup>, which is significantly reduced compared to that of Au/Mn-MIL-100 (*i.e.* 736 m<sup>2</sup> g<sup>−1</sup>) due to the pyrolysis of its precursor. Although MIL-100, with a high specific surface area, has been extensively researched because of its high catalytic activity for various reactions, in this study its high specific area was taken advantage of to make the distribution of Au NPs uniform, and it was converted into a manganese-oxide support, which was the catalytically active ingredient for the PCl<sub>3</sub> oxidation reaction afterwards.<sup>52,53</sup> Thus, Mn-MIL-100 was used as a template to separate the Au clusters rather than as a material for the catalyst. During the process of pyrolysis, the H<sub>3</sub>BTC linker was sacrificed to make the Mn ions move close to each other and generate the interface with the Au NPs. XPS measurements were carried out to figure out the electronic properties of the Mn and Au elements in the Au<sub>*x*</sub>/Mn<sub>3</sub>O<sub>4</sub> catalysts. In the Mn 2p spectra, as presented in Fig. 5C, four peaks with electron binding energies of 641.4, 643.3, 653.0

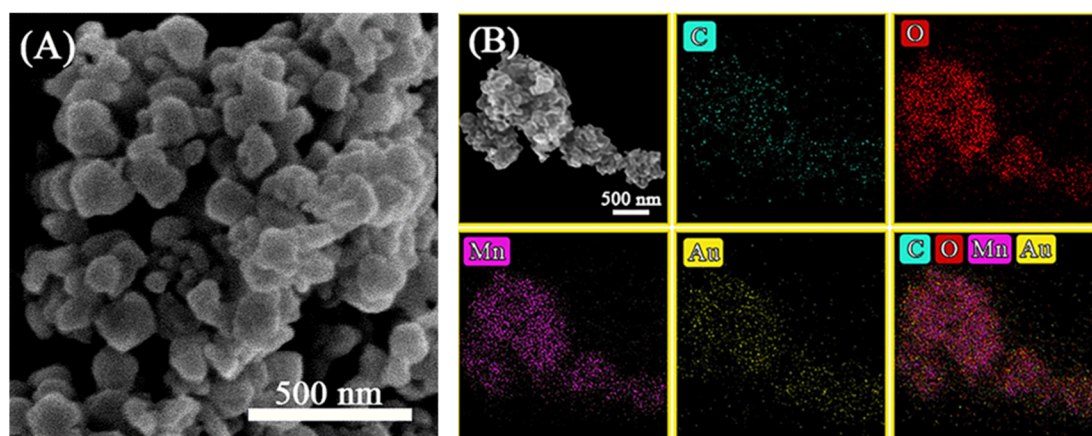


Fig. 4 (A) SEM image, and (B) element distribution of C, O, Mn and Au species *via* SEM-EDX of Au<sub>*x*</sub>/Mn<sub>3</sub>O<sub>4</sub>.



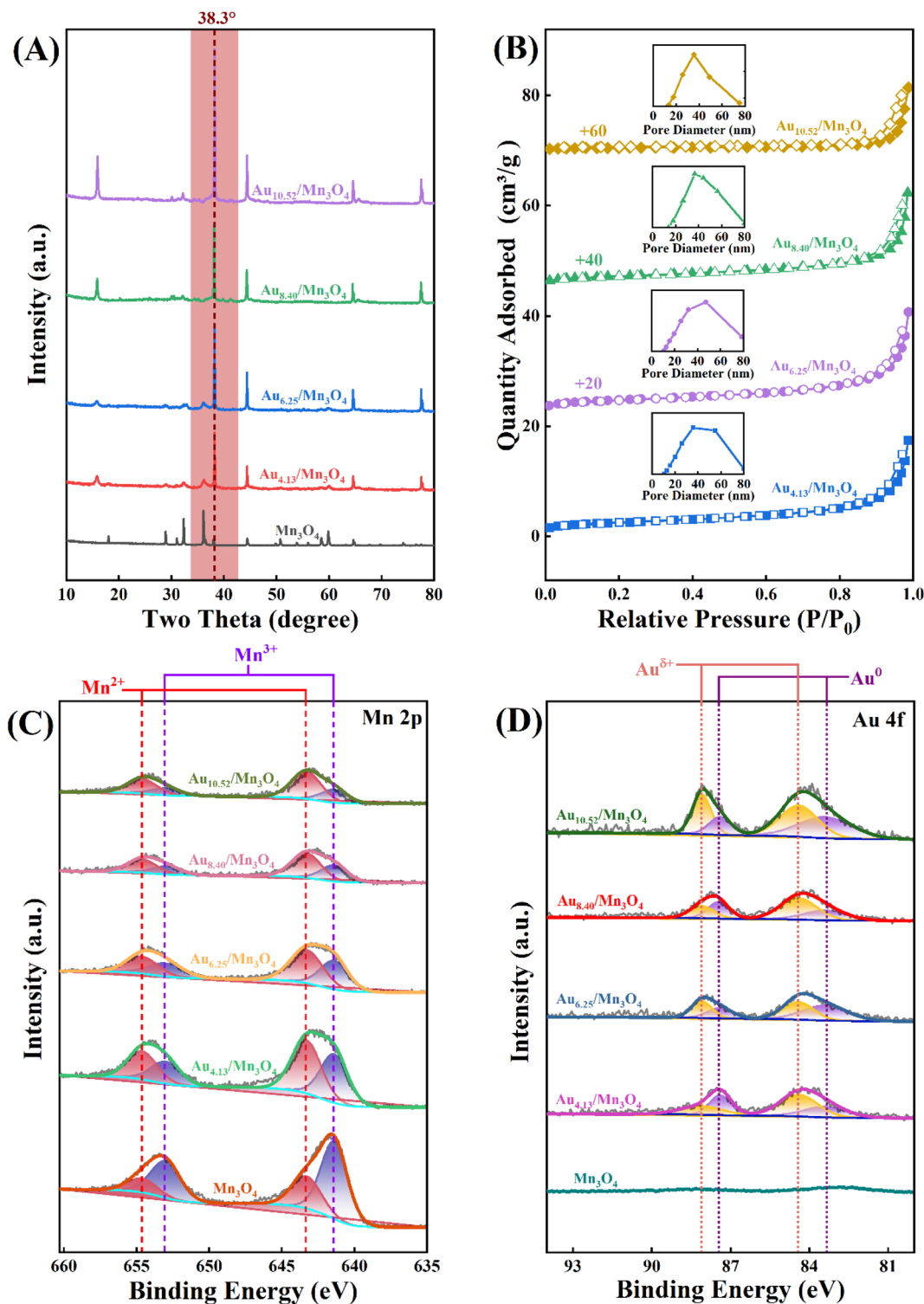


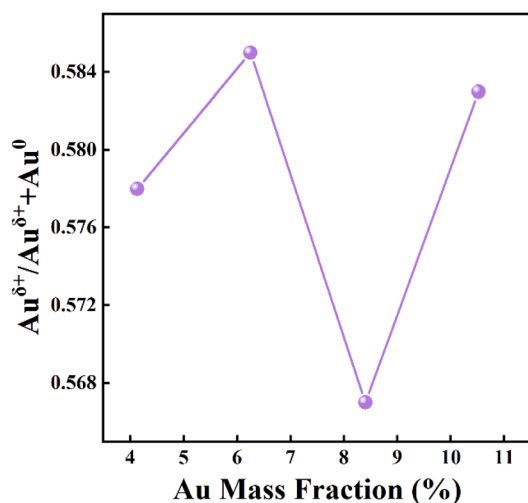
Fig. 5 (A) XRD patterns, (B)  $\text{N}_2$  adsorption-desorption isotherms (with inset pore-size distributions), (C) Mn 2p XPS spectra, and (D) Au 4f XPS spectra of  $\text{Au}_x/\text{Mn}_3\text{O}_4$ .

and 654.6 eV are observed after deconvolution and can be attributed to  $\text{Mn}^{2+}$  and  $\text{Mn}^{3+}$ , further confirming the formation of the  $\text{Mn}_3\text{O}_4$  support.<sup>54</sup> In the Au 4f spectra exhibited in Fig. 5D, the Au species in  $\text{Au}_x/\text{Mn}_3\text{O}_4$  show binding energies of 83.4, 84.4, 87.4 and 88.1 eV, which can be ascribed to  $\text{Au}^0$  with a zero

valence state and  $\text{Au}^{\delta+}$  with a minor positive charge, indicating the interaction in terms of charge transfer between the Au NPs and  $\text{Mn}_3\text{O}_4$  support at their generated interface. The  $\text{Au}^{\delta+}/(\text{Au}^0 + \text{Au}^{\delta+})$  molar ratio was obtained from the analysis of the corresponding peak area. An increasing-decreasing trend is

**Table 1** Summary of specific surface areas, total pore volumes and pore sizes of catalysts with different mass fractions of Au

	Specific surface area (m <sup>2</sup> g <sup>-1</sup> )	Pore volume (cm <sup>3</sup> g <sup>-1</sup> )	Pore size (nm)
Au <sub>4.13</sub> /Mn <sub>3</sub> O <sub>4</sub>	8.7	0.026	13
Au <sub>6.25</sub> /Mn <sub>3</sub> O <sub>4</sub>	9.3	0.027	13
Au <sub>8.40</sub> /Mn <sub>3</sub> O <sub>4</sub>	7.7	0.026	14
Au <sub>10.52</sub> /Mn <sub>3</sub> O <sub>4</sub>	6.4	0.046	31

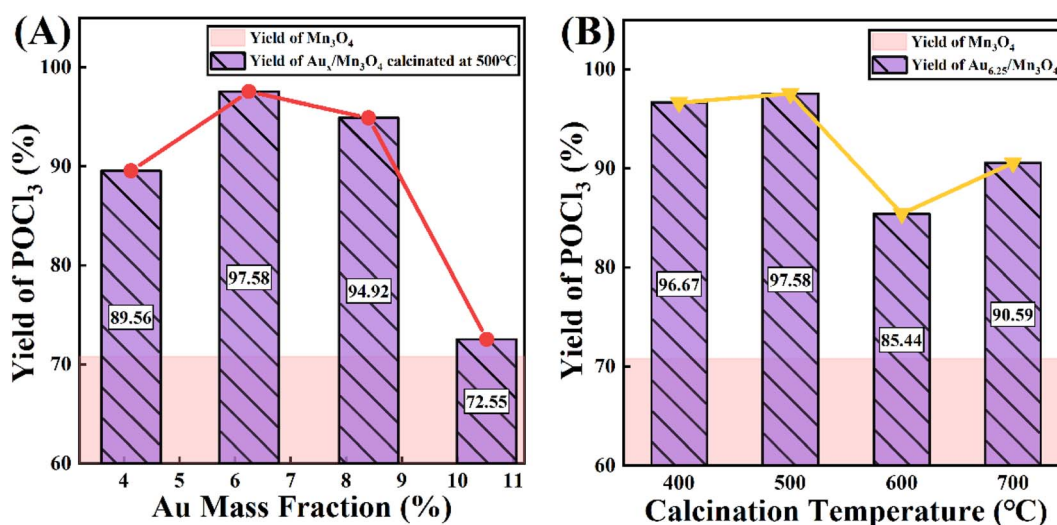


**Fig. 6** The influence of Au mass fraction on the  $\text{Au}^{\delta+}/(\text{Au}^{\delta+} + \text{Au}^{\delta0})$  molar ratio of the  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalysts.

observed for the  $\text{Au}^{\delta+}/(\text{Au}^{\delta+} + \text{Au}^{\delta0})$  molar ratio with the increase in the Au mass fraction (Fig. 6), reaching the maximum value of 0.585 when the Au mass fraction is 6.25. It can be inferred that although the amounts of  $\text{Au}^{\delta+}$  and  $\text{Au}^{\delta0}$  increase with the elevation in the total Au content, the degree of increase in the  $\text{Au}^{\delta+}$  amount is less than the increase in the  $\text{Au}^{\delta0}$  amount when the Au mass fraction exceeds 6.25.

### 3.2 $\text{POCl}_3$ oxidation over $\text{Au}_x/\text{Mn}_3\text{O}_4$ catalysts

Trace  $\text{POCl}_3$  was oxidized to  $\text{POCl}_3$  by  $\text{O}_2$  over the as-prepared  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalysts; this was done in an autoclave, where  $\text{O}_2$  was introduced into the system and used for pressure control. The yield of  $\text{POCl}_3$  after the reaction was chosen for evaluating the catalytic activity of the catalysts. Fig. 7A presents the  $\text{POCl}_3$  yield of the  $\text{Au}/\text{Mn}_3\text{O}_4$  catalysts with different Au mass fractions. It can be clearly seen that the  $\text{POCl}_3$  yield was dramatically enhanced from 70.58% to 89.56% once Au was loaded on the support, revealing that the presence of Au species gave a significant improvement in the activity of the catalysts, verifying that the catalytic performance of the as-synthesized catalysts derived from MOFs was greatly enhanced in comparison to that of the traditional catalyst  $\text{Mn}_3\text{O}_4$ . The yield of  $\text{POCl}_3$  then showed a trend of first increasing and then decreasing with the rise in Au content, where  $\text{Au}_{6.25}/\text{Mn}_3\text{O}_4$  gave the maximum  $\text{POCl}_3$  yield of 97.58%. It is widely accepted that a close relationship between the metal loading on a support and its catalytic performance can be ascribed to the geometric and electronic effects on the interface of the metal NPs and their support. In terms of the geometric effects, a higher Au content creates more active sites at the beginning, but the size of the Au NPs embedded in the metal oxide becomes too large as the Au content increases, leading to a decrease in the number of active sites instead. From the aspect of the electronic effects, the trend of the  $\text{Au}^{\delta+}/(\text{Au}^{\delta+} + \text{Au}^{\delta0})$  molar ratio is in accordance with the trend of the catalytic performance with the rise in Au content, revealing that the catalytic activity was mainly influenced by the relative content of  $\text{Au}^{\delta+}$ . To summarize, the origin of the influence of Au content on the  $\text{POCl}_3$  yield was intrinsically derived from the synergistic results of geometric and electronic effects. Fig. 7B shows the yield of  $\text{POCl}_3$  over  $\text{Au}_x/\text{Mn}_3\text{O}_4$  catalysts calcined at different temperatures (400, 500, 600 and 700 °C), where the best catalytic activity was obtained with the catalyst calcined at 500 °C. This can be attributed to the aggregation of catalyst particles and the encapsulation of Au NPs into the



**Fig. 7** The influence of (A) Au mass fraction and (B) calcination temperature on catalytic performance of  $\text{Au}/\text{Mn}_3\text{O}_4$  in terms of the yield of  $\text{POCl}_3$  in the  $\text{PCl}_3$  oxidation reaction.



carbon matrix as the calcination temperature exceeds 500 °C, leading to a significant reduction in active sites.

Subsequently, four process parameters, namely agitation speed, reaction temperature, reaction pressure and catalyst dosage, were probed to obtain optimum process conditions that enable a maximum yield of  $\text{POCl}_3$  (Fig. 8). The yield of  $\text{POCl}_3$  (97.58%) at an agitation speed of 500 rpm was slightly higher than that at 400 rpm and then showed inconspicuous changes with the increase in agitation speed (Fig. 8A). In terms of the reaction temperature (Fig. 8B), an increasing-decreasing trend was observed for the  $\text{POCl}_3$  yield as the reaction temperature continued to rise, reaching a maximum yield of 99.13% at 80 °C. For the influence of reaction pressure on the reaction, as presented in Fig. 8C, the yield of  $\text{POCl}_3$  was remarkably enhanced when the pressure increased to 0.6 MPa from 0.5 MPa, reaching 99.13%, which was the maximum value across the whole pressure range. With regard to the catalyst dosage (Fig. 8D), the optimal yield of  $\text{POCl}_3$  was obtained at a dosage of 40 mg. In

conclusion, after comprehensively considering  $\text{POCl}_3$  yield and economic efficiency, the optimal technological parameters were selected as follows: an agitation speed of 500 rpm, reaction temperature of 80 °C, reaction pressure of 0.6 MPa and catalyst dosage of 40 mg.

### 3.3 Reaction mechanism

According to the results shown in Fig. 7 and 8, we hypothesize that the possible explanation for the higher  $\text{PCl}_3$  oxidation activity of Au NPs supported on  $\text{Mn}_3\text{O}_4$  is the supply of oxygen from the lattice of the metal-oxide support in a Mars van Krevelen (MvK) process. Fig. 9 shows the reaction sequence of  $\text{PCl}_3$  oxidation over the Au/ $\text{Mn}_3\text{O}_4$  catalysts, which includes a MvK-style supply of oxygen. First,  $\text{PCl}_3$  is adsorbed close to the interface between the Au NPs and metal-oxide support. Then lattice oxygen of  $\text{Mn}_3\text{O}_4$  forms a new bond with phosphorus, transforming  $\text{PCl}_3$  into  $\text{POCl}_3$  and forming an oxygen anion vacancy on the interface at the same time. O atoms produced by

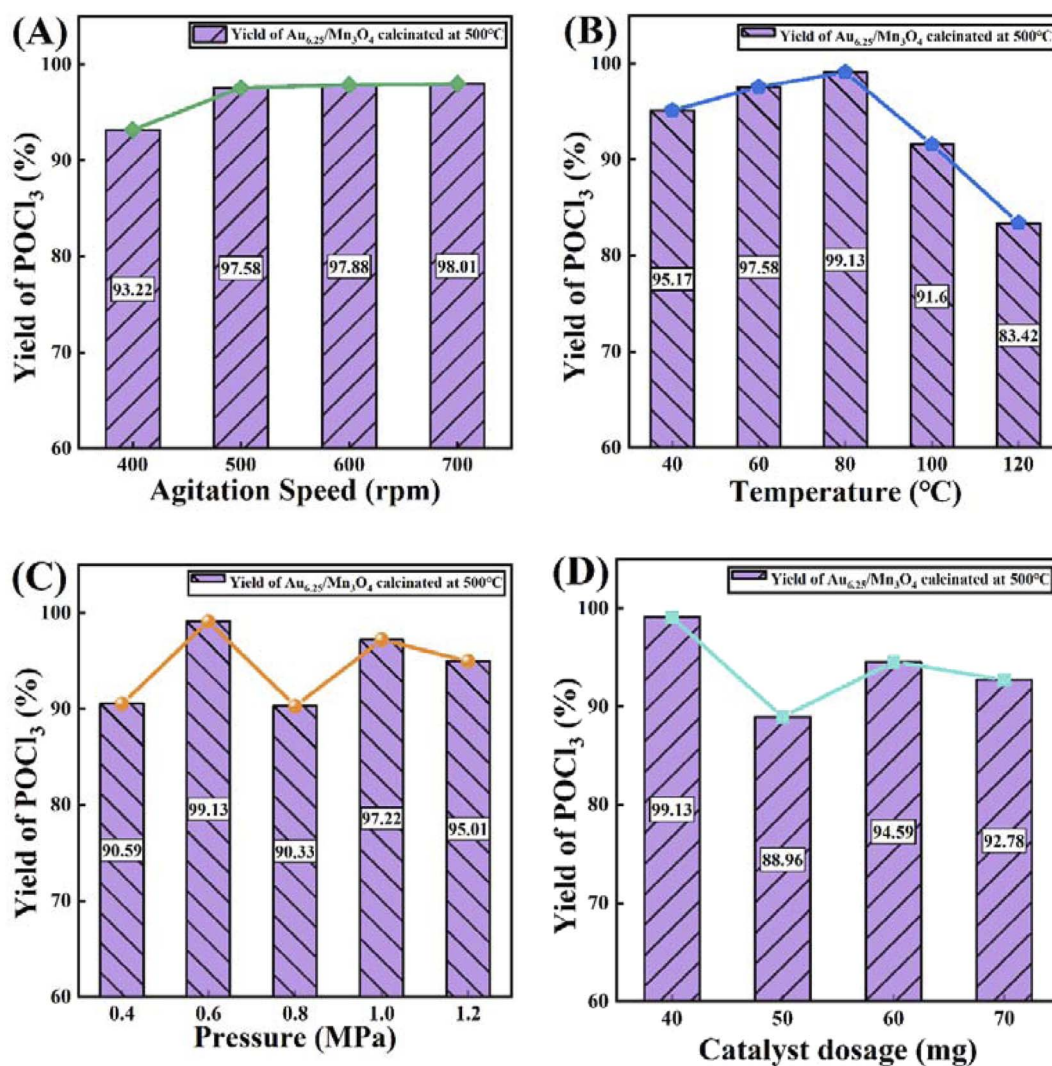


Fig. 8 The influence of (A) agitation speed, (B) reaction temperature, (C) reaction pressure and (D) catalyst dosage on catalytic performance of  $\text{Au}_{6.25}/\text{Mn}_3\text{O}_4$  in terms of the yield of  $\text{POCl}_3$  in the  $\text{PCl}_3$  oxidation reaction.

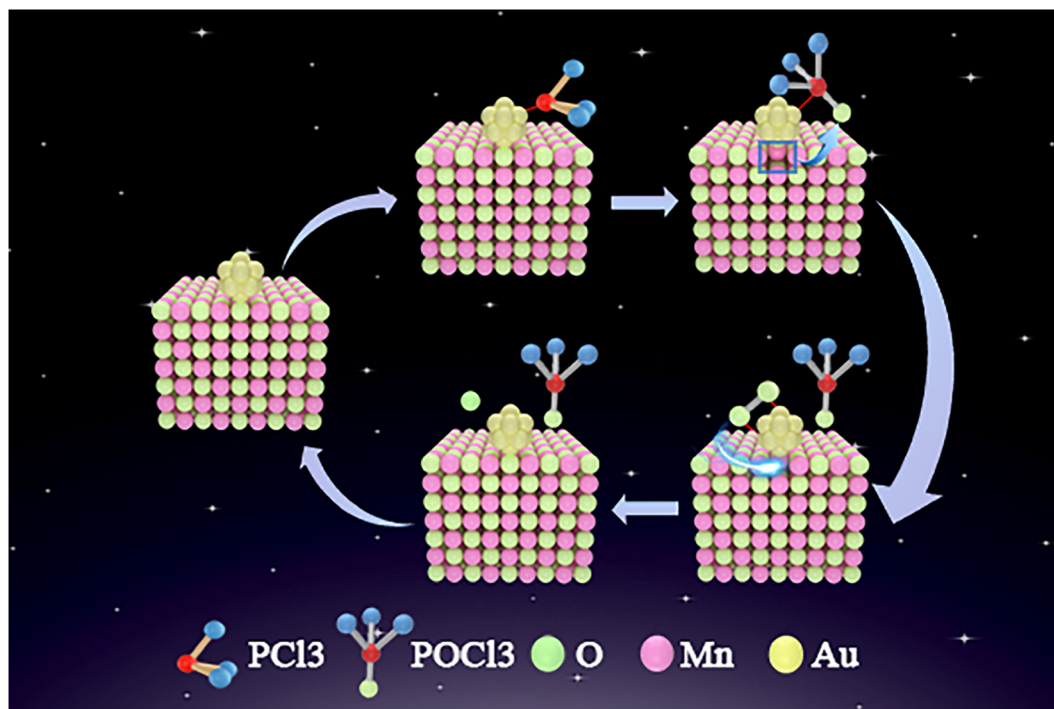
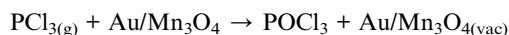


Fig. 9 MvK process for oxidation of  $\text{PCl}_3$  over the interface between the Au NPs and  $\text{Mn}_3\text{O}_4$ .

the dissociation of  $\text{O}_2$  that was adsorbed on the interface between the Au NPs and the support compensate the oxygen vacancies; thus, the final result is that  $\text{PCl}_3$  is oxidized by oxygen over the Au/ $\text{Mn}_3\text{O}_4$  catalysts, which maintain their properties after the reaction.<sup>16,20</sup>

To demonstrate this hypothesis, DFT calculations were performed to analyze the thermodynamics of the oxidation of  $\text{PCl}_3$  by the oxygen from the lattice of the support, generating Au/ $\text{Mn}_3\text{O}_4$  with an oxygen vacancy and  $\text{POCl}_3$ .<sup>55</sup> The equation is as follows:



The oxygen atoms of the Au/ $\text{Mn}_3\text{O}_4$  catalysts can be divided into two groups. One is the interface oxygen atoms ( $\text{O}_{\text{int}}$ ) that bridge the Mn atoms with Au; the other is the perimeter oxygen atoms ( $\text{O}_{\text{peri}}$ ) that bridge two Mn atoms at the periphery of Au. In order to figure out which group of oxygen atoms is effective in the above process, the variation in Gibbs free energy ( $\Delta G$ ) for the reactions involving  $\text{O}_{\text{int}}$  and  $\text{O}_{\text{peri}}$ , respectively, were calculated as shown in Fig. 10.

The values of  $\Delta G$  suggest that  $\text{PCl}_3$  oxidation by lattice oxygens from clean  $\text{Mn}_3\text{O}_4$  is non-spontaneous from 100 to 800 K. Similarly,  $\text{PCl}_3$  oxidation by  $\text{O}_{\text{peri}}$  is difficult thermodynamically owing to the positive variation in the Gibbs free energy. The variation in  $\Delta G$  exhibits a trend of slightly decreasing with an increase in temperature, but the trend of overall has a flat manner. The situation is quite different when the  $\text{O}_{\text{int}}$  atoms are used to participate in the  $\text{PCl}_3$  oxidation reaction with a MvK-style supply of oxygen. The lattice oxygen atoms at the

interface between the Au NPs and  $\text{Mn}_3\text{O}_4$  have negative free energies from 100–800 K, revealing that the lattice oxygen atoms of Au/ $\text{Mn}_3\text{O}_4$  that are located near the periphery of Au would not participate in the  $\text{PCl}_3$  oxidation reaction, whereas the lattice oxygen atoms that bridge the Au cluster with the metal-oxide support show great potential for the reaction. An insignificant descending trend in the Gibbs free energy variation for  $\text{O}_{\text{int}}$  can also be observed as the temperature increases, indicating that the spontaneity of the  $\text{PCl}_3$  oxidation reaction using  $\text{O}_{\text{int}}$  as the

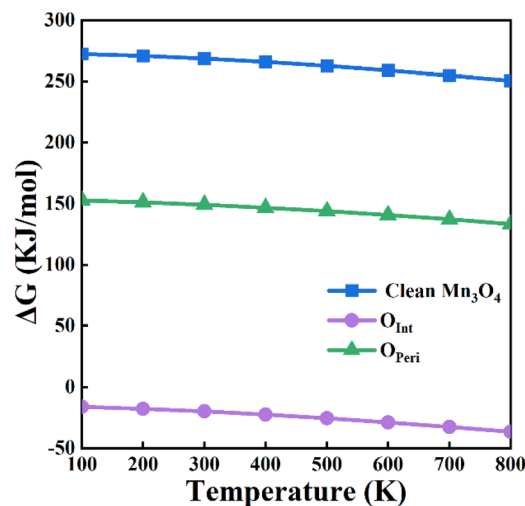


Fig. 10 Variation in  $\Delta G$  for the formation of  $\text{POCl}_3$  using the lattice oxygens of  $\text{Mn}_3\text{O}_4$  and using the interface and perimeter lattice oxygens of the  $\text{Au}_{10}/\text{Mn}_3\text{O}_4$  model system.



reactant becomes more intense, but the effect is not that remarkable. The comparison of  $\Delta G$  for the reaction between the lattice oxygen from clean  $\text{Mn}_3\text{O}_4$  and  $\text{O}_{\text{int}}$  also suggested that the bridging lattice oxygens between the Au clusters and Mn species were activated by the loading of Au; this indicates the generation of an interaction that could change the chemical potential of the lattice oxygen between the Au and metal-oxide support, ensuring the reaction proceeded and improving the catalytic activity of the Au/ $\text{Mn}_3\text{O}_4$  catalyst.

The corresponding relationship between the trend in the  $\text{Au}^{\delta+}/(\text{Au}^0 + \text{Au}^{\delta+})$  molar ratio and the increase in the Au mass fraction, and the trend in the  $\text{POCl}_3$  yield with the increase in the Au mass fraction, indicated that  $\text{Au}^{\delta+}$  has a certain supporting effect on the  $\text{PCl}_3$  oxidation reaction. The MvK mechanism gives an insight into the possible role of  $\text{Au}^{\delta+}$ , in that it promotes the dissociation of  $\text{O}_2$  adsorbed on the Au-support interface. The amount of lattice oxygen on the interface between the Au NPs and  $\text{Mn}_3\text{O}_4$  is restricted, such that  $\text{PCl}_3$  oxidation would be unable to happen when the rate of  $\text{PCl}_3$  oxidation is faster than that of lattice oxygen supplementation. The rate of lattice oxygen supplementation depends on the rate of oxygen adsorption and dissociation. Owing to the system being exposed to the oxygen-rich and constantly stirred environment, we suppose that a constant supply of oxygen is introduced to the interface of catalyst. The strong interaction in terms of charge transfer between the Au NPs and oxide support promotes the generation of  $\text{Au}^{\delta+}$ , which subsequently accelerates the dissociation of oxygen with as much oxidation as possible of every  $\text{PCl}_3$  adsorbed on the Au NPs, resulting in the enhancement in  $\text{PCl}_3$  yield.

The Au/ $\text{Mn}_3\text{O}_4$  catalyst engineered *via* the MOF-mediated method consists of Au NPs deposited on a manganese-oxide support, which is the traditional, industrial, active catalytic component for trace  $\text{PCl}_3$  oxidation in an organic system. Compared with the traditional  $\text{Mn}_3\text{O}_4$  catalyst, Au NPs loaded on the support were strongly capable of capturing the trace reactant  $\text{PCl}_3$  and activating the lattice oxygen of the metal oxide, enabling  $\text{PCl}_3$  to be directly oxidized by the lattice oxygen bridging the oxide support and the Au atom that it adsorbs on. The synthesis method, which derives the catalyst from the corresponding MOF composite, could significantly enhance the uniformity of Au NP distribution and induce the formation of an active interface, promoting an increase in the number of reaction active sites and the supplementation of lattice oxygen vacancies in comparison to conventional metal-loading methods. Thus, the as-synthesized catalyst exhibits remarkable progress in catalytic activity compared with the catalysts used in industry or synthesized by conventional methods, with a yield of  $\text{POCl}_3$  higher than that for  $\text{Mn}_3\text{O}_4$  by 28.55 percentage points. In addition, it is inevitable that the cost and the process complexity of the catalyst engineered in this study are higher than for the traditional catalyst, and could be further investigated and modified.

## 4 Conclusion

To sum up, a high-performance catalyst, Au/ $\text{Mn}_3\text{O}_4$ , for trace  $\text{PCl}_3$  oxidation reaction in an organic system was successfully

synthesized by using an effective strategy wherein the catalyst was derived from MOFs. This method provides an efficient way to synthesize catalysts with a uniform distribution of Au NPs on a metal-oxide support and strong interaction between them, which could promote the catalytic activity of the as-synthesized catalyst. DFT calculations suggested that a possible catalytic mechanism is the supply of lattice oxygen as a  $\text{PCl}_3$  oxidant, which is from interface oxygens bridging the Au NPs and Mn, in a MvK process. Structure-performance analysis revealed that the interaction between the Au and support, in terms of charge transfer and chemical potential change, dominantly determined the yield of  $\text{PCl}_3$ . This work provides an effective method, an active catalyst and the best process parameters for the dephosphorization of trichlorosilane feedstocks in the polysilicon industry.

## Conflicts of interest

There are no conflicts of interest to declare.

## Acknowledgements

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