



# Highly active spinel type $\text{CoCr}_2\text{O}_4$ catalysts for dichloromethane oxidation



Yu Wang, Ai-Pin Jia, Meng-Fei Luo, Ji-Qing Lu\*

Key Laboratory of the Ministry of Education for Advanced Catalysis Materials, Institute of Physical Chemistry, Zhejiang Normal University, Jinhua 321004, China

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## ABSTRACT

A series of spinel type  $\text{CoCr}_2\text{O}_4$  catalysts calcined at different temperatures were prepared and tested for oxidation of dichloromethane ( $\text{CH}_2\text{Cl}_2$ ). These catalysts were active for this reaction. The final products in the reaction were  $\text{CO}_x$ ,  $\text{HCl}$  and  $\text{Cl}_2$ , without the formation of  $\text{Cl}$ -containing organics. The best performance was obtained on a catalyst calcined at  $400^\circ\text{C}$  ( $\text{CoCr}_2\text{O}_4$ -4), with a  $T_{50}$  of  $218^\circ\text{C}$  and a  $T_{90}$  of  $257^\circ\text{C}$ , which was mainly due to the highest surface area of this catalyst ( $91.3\text{ m}^2\text{ g}^{-1}$ ). Detailed quantitative analyses revealed that the catalytic behavior was synergistically governed by surface acidity and reducibility of the catalyst, as evidenced by ammonia temperature-programmed desorption and hydrogen temperature-programmed reduction results, respectively. Kinetic studies revealed similar activation energies ( $124.5$ – $155.5\text{ kJ mol}^{-1}$ ) on these catalysts, implying the reaction might follow the same pathways on these catalysts. More importantly, in situ Fourier transform infrared spectroscopic investigation of the reaction revealed that formate species were the main reaction intermediates, which could be further oxidized to  $\text{CO}$  and  $\text{CO}_2$ .

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## 1. Introduction

Chlorinated volatile organic compounds (CVOCs) are associated with a broad range of industrial processes and they are recognized as a major source of air pollution [1]. The abatement of CVOCs thus becomes a very important topic in catalysis because catalytic combustion offers advantages over thermal destruction, with lower incineration temperature and higher destructive efficiency [2].

The most commonly employed catalysts for CVOCs oxidation include noble metals such as  $\text{Pt}$  [3,4], transition metal oxides such as  $\text{MnO}_x$  [5],  $\text{CrO}_x$  [6] and perovskite-type oxides [7–9], rare earth oxides such as  $\text{CeO}_2$  [10] and zeolite [11] catalysts. Transition metal oxides catalysts are promising in practical applications because of their relatively better performance and resistance to deactivation compared to the other three type catalysts [12].

Among the transition metal oxides, Cr-based oxides are very effective for CVOCs oxidation [13–15]. Various Cr-based catalysts have been reported in literature, including  $\text{CrO}_x/\text{TiO}_2$  [12,16,17],  $\text{CrO}_x/\text{C}$  [18],  $\text{CrO}_x/\text{zeolites}$  [19,20] and  $\text{CrO}_x\text{-CeO}_2/\text{USY}$  [21]. For example, Krishnamoorthy et al. [16] conducted oxidation of 1,2-dichlorobenzene over various supported transition metal oxides

and found that the  $\text{CrO}_x/\text{TiO}_2$  was the most active catalyst compared to other  $\text{TiO}_2$ -supported metal oxides ( $\text{V}_2\text{O}_5$ ,  $\text{MoO}_3$ ,  $\text{Fe}_2\text{O}_3$ , and  $\text{Co}_3\text{O}_4$ ), with a  $T_{90}$  (temperature when 90% conversion obtained) of about  $290^\circ\text{C}$ . Moreover, Huang et al. [21] reported oxidation of various CVOCs (dichloromethane, trichloroethylene and 1,2-dichloroethane) over  $\text{Cr}_2\text{O}_3\text{-CeO}_2\text{-USY}$  catalysts, and they found that  $T_{90}$  for dichloromethane, trichloroethylene and 1,2-dichloroethane were 281, 296 and  $246^\circ\text{C}$ , respectively. Despite of their high activities, however, most of the supported Cr catalysts suffer severe deactivation due to the migration and/or loss of active sites during the reaction. Besides, another obstacle of the Cr catalysts for CVOCs oxidation is the formation of the extremely toxic chromium oxychloride at low temperature [22].

To overcome the disadvantages of the Cr catalysts, a possible strategy is to confine the Cr species in a robust structure, by which may suppress the transformation and leakage of Cr species during the CVOCs oxidation and hence increase the catalyst stability. In this sense, spinel type cobalt chromite oxide ( $\text{CoCr}_2\text{O}_4$ ) might be a potential candidate. In such a  $\text{CoCr}_2\text{O}_4$  structure, Co and Cr cations are bound in a solid matrix, in which the high-spin divalent  $\text{Co}^{2+}$  cations mostly occupy the tetrahedral sites, while the low-spin trivalent  $\text{Cr}^{3+}$  cations occupy the octahedral sites [23]. Indeed, the spinel type  $\text{CoCr}_2\text{O}_4$  oxide has been widely applied in electrochromic coatings [24], protective layers [25], solar absorbers [26], gas sensors [27], electrodes for fuel cells [28], and oxidation of

\* Corresponding author. Tel.: +86 579 82287325; fax: +86 579 82282595.  
E-mail address: [jiqinglu@jznu.cn](mailto:jiqinglu@jznu.cn) (J.-Q. Lu).

different volatile organic compounds [29]. Chen et al. [29] reported methane combustion over a series of Co–Cr catalysts with different Co/Cr ratios and the best performed catalysts was a Co1Cr2 with a Co/Cr molar ratio of 1/2 (a spinel type  $\text{CoCr}_2\text{O}_4$  oxide). The authors concluded that the superior performance of Co1Cr2 was due to the presence of high valent  $\text{Cr}^{6+}$  species, which caused disorders in the spinel structure of cobalt chromites and consequently enhance the adsorption of chemisorbed oxygen species. In application of CVOs oxidation, Kim and Ihm [30] found that spinel type  $\text{CoCr}_2\text{O}_4$  showed superior performance to other supported Cr catalysts such as  $\text{CrO}_x/\text{Al}_2\text{O}_3$  and  $\text{CrO}_x/\text{MCM-41}$  in total oxidation of trichloroethylene, and the authors attributed high  $\text{CO}_2$  selectivity to the presence of abundant surface  $\text{Cr}^{3+}$  species in the  $\text{CoCr}_2\text{O}_4$  oxide.

It can be seen that spinel type  $\text{CoCr}_2\text{O}_4$  oxide could be promising catalyst for CVOs oxidation; however, very limited information has been provided in literature. In addition to the development of highly active catalysts for CVOs oxidation, some important issues such as active sites, reaction mechanisms should also be addressed in order to better understand this catalyst system. Therefore, in the current work, a series of nanocrystalline spinel type  $\text{CoCr}_2\text{O}_4$  oxides were prepared and tested for the total oxidation of  $\text{CH}_2\text{Cl}_2$  as a model reaction. It was found that these catalysts are very active and quite stable for the oxidation of  $\text{CH}_2\text{Cl}_2$ , and the catalytic behaviors were related to some important parameters of the catalysts such as surface acidity and reducibility. Besides, kinetic study as well as in situ infrared spectroscopic investigation provided some insights on the reaction mechanism on these catalysts.

## 2. Experimental

### 2.1. Catalyst preparation

The  $\text{CoCr}_2\text{O}_4$  catalysts were prepared with a sol–gel method as described in a previous work [31]. In a typical procedure, 88 mmol  $\text{Co}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$  (99.0%), 176 mmol  $\text{Cr}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$  (99.0%) and 528 mmol citric acid (99.5%) were dissolved in 200 ml distilled water. The mixture was mildly evaporated at  $90^\circ\text{C}$  until a viscous gel was obtained, which was then heated at  $180^\circ\text{C}$  for 3 h. The final solid was ground and divided into four parts, followed by calcination in static air at different temperatures ( $400$ – $700^\circ\text{C}$ ) for 4 h. The obtained catalysts were designated as  $\text{CoCr}_2\text{O}_4\text{-x}$ , with x referring to calcination temperature.

### 2.2. Catalyst characterizations

Elemental compositions of the catalysts were determined by X-ray fluorescence (XRF) analysis, in an ARL ADVANT'X Intelli Power 4200 scanning X-ray fluorescence spectrometer. The results were analyzed using UniQuant nonstandard sample quantitative analysis software.

The BET surface areas of the catalysts were measured by  $\text{N}_2$  adsorption at liquid-nitrogen temperature ( $77\text{ K}$ ), using a surface area analyzer (Quantachrome Autosorb-1). The catalysts were pretreated at  $120^\circ\text{C}$  for 6 h in vacuum.

The morphologies of the catalysts were observed by a scanning electron microscopy (SEM, Hitachi S-4800) operated at 5.0 kV.

X-ray diffraction (XRD) patterns were recorded with a PANalytical X'Pert PRO MPD powder diffractometer using  $\text{Cu K}\alpha$  radiation. The working voltage was 40 kV and the working current was 40 mA. The patterns were collected in a  $2\theta$  range from  $10^\circ$  to  $80^\circ$ , with a scanning speed of  $0.15^\circ\text{ s}^{-1}$ . Crystallite sizes and lattice parameters of the catalysts were analyzed by full-curve fitting using a JADE 6.5 software.

Raman spectra were recorded on a Renishaw RM1000 with a confocal microprobe Raman system using an excitation wavelength of 514 nm.

Hydrogen temperature-programmed reduction ( $\text{H}_2$ -TPR) technique was employed to analyze the reducibility of the catalysts. Twenty-five milligrams of the catalyst were placed in a quartz reactor and pretreated in a  $\text{N}_2$  flow ( $20\text{ ml min}^{-1}$ ) at  $300^\circ\text{C}$  for 60 min in order to remove the adsorbed water and carbonates, then the sample was cooled down to  $50^\circ\text{C}$  in a  $\text{N}_2$  flow ( $20\text{ ml min}^{-1}$ ). After that, the sample was heated from  $50$  to  $900^\circ\text{C}$  with a heating rate of  $10^\circ\text{C min}^{-1}$  under a mixture of 5%  $\text{H}_2$ –95%  $\text{N}_2$  ( $20\text{ ml min}^{-1}$ ). The amount of  $\text{H}_2$  consumption was determined by a gas chromatograph with a thermal conductivity detector (TCD), and the thermal conductivity response was calibrated by the reduction of a known  $\text{CuO}$  powder sample.

The surface acidity of the catalyst was measured by ammonia temperature-programmed desorption ( $\text{NH}_3$ -TPD). Fifty milligrams of the catalyst were pretreated in a flow of  $\text{N}_2$  ( $20\text{ ml min}^{-1}$ ) at  $300^\circ\text{C}$  for 0.5 h, and then was cooled down to  $50^\circ\text{C}$ . Afterwards, a  $\text{NH}_3$  flow ( $20\text{ ml min}^{-1}$ ) was introduced to the sample for 15 min, followed by purging at  $80^\circ\text{C}$  for 0.5 h with a  $\text{N}_2$  flow ( $20\text{ ml min}^{-1}$ ) to remove the physisorbed  $\text{NH}_3$ . Then the sample was heated from  $80$  to  $500^\circ\text{C}$  at a rate of  $10^\circ\text{C min}^{-1}$ , and the profile was recorded using a gas chromatograph (TECHTEMP GC 7890II) with a TCD detector.

X-ray photoelectron spectra of the catalysts were obtained on an ESCALAB 250Xi instrument, with a  $\text{Al K}\alpha$  X-ray source ( $1486.6\text{ eV}$ ), under about  $2 \times 10^{-9}\text{ mbar}$  at room temperature and a pass energy of 20 eV. The binding energy (BE) of C1s core level at 284.8 eV was taken as the internal standard.

Temperature-programmed surface reaction (TPSR) was conducted on a home-made reactor connected with a mass spectrometer (MS, Qic-20 Benchtop, HidenAnalytical). Fifty milligrams of the catalyst were pretreated in a flow of air ( $20\text{ ml min}^{-1}$ ) at  $300^\circ\text{C}$  for 0.5 h, and then was cooled down to  $50^\circ\text{C}$ . Then a flow of  $\text{CH}_2\text{Cl}_2$ /air mixture (3000 ppm  $\text{CH}_2\text{Cl}_2$ , total flow rate =  $20\text{ ml min}^{-1}$ ) was introduced to the reactor and the sample was heated from  $50$  to  $400^\circ\text{C}$  at a rate of  $10^\circ\text{C min}^{-1}$ . And  $m/e$  signals of 70, 44, 18, 36.5, 50.5, 49, 60 and 30 were monitored, corresponding to  $\text{Cl}_2$ ,  $\text{CO}_2$ ,  $\text{H}_2\text{O}$ ,  $\text{HCl}$ ,  $\text{CH}_3\text{Cl}$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{COCl}_2$  and  $\text{HCHO}$ , respectively. Due to the same  $m/e$  signals of CO and  $\text{N}_2$  ( $m/e = 28$ ), the formation of CO during the reaction could not be correctly identified.

In situ Fourier transform infrared (FTIR) spectra of the samples were recorded on a NEXUS670 spectrometer equipped with an MCT detector. Self-supported sample wafers (diameter = 16 mm) were prepared from 30 mg of sample by pressing at about 3 MPa. The sample was transferred to a quartz IR cell connected to the closed circulation systems and then pretreated under an air flow ( $90\text{ ml min}^{-1}$ ) at  $400^\circ\text{C}$  for 1 h. After the pretreatment, the sample was cooled to  $50^\circ\text{C}$  and 1 ml of a gas mixture (3000 ppm  $\text{CH}_2\text{Cl}_2$  in air) was introduced to the IR cell and the sample was heated from  $50$  to  $400^\circ\text{C}$  at a ramp of  $10^\circ\text{C min}^{-1}$ . Temperature-dependent FTIR spectra during the reaction were recorded after holding each temperature point for 10 min.

### 2.3. Activity test

Catalytic combustion of  $\text{CH}_2\text{Cl}_2$  was carried out in a conventional fixed-bed reactor (i.d. = 9 mm). One gram of catalyst in 40–60 mesh was diluted into a volume of 2 ml with quartz sand, and then it was loaded in the reactor. A thermal couple was placed in the middle of the catalyst bed to monitor the reaction temperature. The  $\text{CH}_2\text{Cl}_2$  was introduced to the reactor via passing moisture-containing air ( $7\text{ ml min}^{-1}$ ) through a quartz beaker containing liquid  $\text{CH}_2\text{Cl}_2$  kept at  $0^\circ\text{C}$ , which was then

mixed with another moisture-containing air flow ( $493 \text{ ml min}^{-1}$ ) to the reactor. Thus, the concentration of  $\text{CH}_2\text{Cl}_2$  was 3000 ppm (total flow rate =  $500 \text{ ml min}^{-1}$  with a total water vapor concentration of about 12000 ppm, GHSV =  $15000 \text{ h}^{-1}$ ). The conversion of  $\text{CH}_2\text{Cl}_2$  was analyzed by a gas chromatograph (Shimadzu, GC-14C) equipped with an FID detector. The outlet reaction mixture was neutralized by passing through a 0.1 M NaOH aqueous solution.

Conversion of  $\text{CH}_2\text{Cl}_2$  was calculated as follows:

$$X_{\text{CH}_2\text{Cl}_2} = \frac{[\text{CH}_2\text{Cl}_2]_{\text{in}} \text{vol.}\% - [\text{CH}_2\text{Cl}_2]_{\text{out}} \text{vol.}\%}{[\text{CH}_2\text{Cl}_2]_{\text{in}} \text{vol.}\%}$$

where  $[\text{CH}_2\text{Cl}_2]_{\text{in}}$  and  $[\text{CH}_2\text{Cl}_2]_{\text{out}}$  were the  $\text{CH}_2\text{Cl}_2$  concentrations in the inlet and outlet gas (vol.%), respectively.

The kinetic study was performed on the same fixed bed reactor of the catalytic  $\text{CH}_2\text{Cl}_2$  oxidation as mentioned above. The feed gases were measured with mass flow controllers and mixed prior to the reactor inlet. For kinetic measurements, the reactor was operated in a differential mode with the  $\text{CH}_2\text{Cl}_2$  conversion less than 15%. 0.5 g catalyst in 40–60 mesh was diluted with quartz sand to a volume of 2 ml and the reaction conditions were the same as in the above-mentioned fixed-bed testing ( $\text{CH}_2\text{Cl}_2$  concentration was 3000 ppm, total flow rate was  $500 \text{ ml min}^{-1}$ ). Also, the absence of mass transport resistances was checked by Weisz–Prater criterion for internal diffusion and Mears' criterion for external diffusion and the absence of heat transfer was checked by Mears' criterion [32] (See Supplementary Information for detailed calculation). For example, on the  $\text{CoCr}_2\text{O}_4$ -4 catalyst, the calculated values under kinetic conditions are  $2.22 \times 10^{-3}$  for the Weisz–Prater criterion for internal diffusion,  $1.19 \times 10^{-2}$  for the Mears' criterion for external diffusion and  $9.5 \times 10^{-3}$  for the Mears' criterion for heat transfer. Those results ensure plug-flow and isothermal conditions within the catalyst bed.

**Table 1**

Physical properties of spinel  $\text{CoCr}_2\text{O}_4$  catalysts.

| Catalyst                     | $S_{\text{BET}}$ ( $\text{m}^2 \text{g}^{-1}$ ) | Cell parameter (nm) | Average crystallite size (nm) |
|------------------------------|-------------------------------------------------|---------------------|-------------------------------|
| $\text{CoCr}_2\text{O}_4$ -4 | 91.3                                            | 0.8314              | 12.8                          |
| $\text{CoCr}_2\text{O}_4$ -5 | 51.9                                            | 0.8316              | 17.7                          |
| $\text{CoCr}_2\text{O}_4$ -6 | 23.2                                            | 0.8324              | 27.4                          |
| $\text{CoCr}_2\text{O}_4$ -7 | 6.1                                             | 0.8350              | 30.7                          |

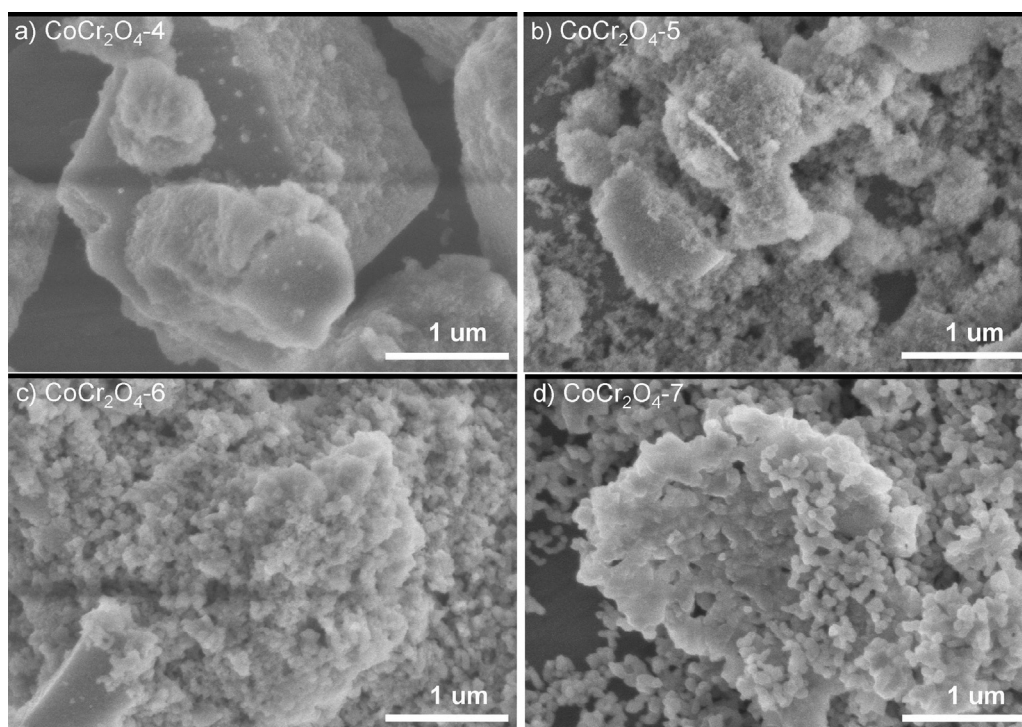
### 3. Results and discussion

#### 3.1. Catalyst characterizations

The bulk compositions of Co and Cr elements in the  $\text{CoCr}_2\text{O}_4$  catalysts are 25.3 and 44.4 wt.%, respectively, as determined by XRF. These values correspond to a Co/Cr molar ratio of 1/2. The surface areas of the catalysts listed in Table 1 decline dramatically with calcination temperature, with the sample calcined at  $400^\circ\text{C}$  ( $\text{CoCr}_2\text{O}_4$ -4) having a surface area of  $91.3 \text{ m}^2 \text{g}^{-1}$  while the sample calcined at  $700^\circ\text{C}$  ( $\text{CoCr}_2\text{O}_4$ -7) having a surface area of  $6.1 \text{ m}^2 \text{g}^{-1}$ . The decline in surface area is due to the sintering of crystallites during the high temperature calcination.

SEM images of the  $\text{CoCr}_2\text{O}_4$  catalysts are shown in Fig. 1. It can be seen that all these samples consist of nano-sized aggregates and no obvious porous structures are detected.

Fig. 2a presents the XRD patterns of the  $\text{CoCr}_2\text{O}_4$  catalysts. All the samples show characteristic diffraction peaks of cubic phase  $\text{CoCr}_2\text{O}_4$  spinel (JCPDS 22-1084) and no other diffraction peaks are detected. Moreover, with increasing calcination temperature, the peaks become more intense. Accordingly, the average crystallite size significantly increase with calcination temperature, ranging from 12.8 nm for the  $\text{CoCr}_2\text{O}_4$ -4 to 30.7 nm for the  $\text{CoCr}_2\text{O}_4$ -7 (Table 1). The growth of the crystallite is in agreement of the decline of surface area. In addition, a slight shift of the diffraction peaks toward lower degrees is observed with increasing



**Fig. 1.** SEM images of (a)  $\text{CoCr}_2\text{O}_4$ -4, (b)  $\text{CoCr}_2\text{O}_4$ -5, (c)  $\text{CoCr}_2\text{O}_4$ -6 and (d)  $\text{CoCr}_2\text{O}_4$ -7 catalysts.

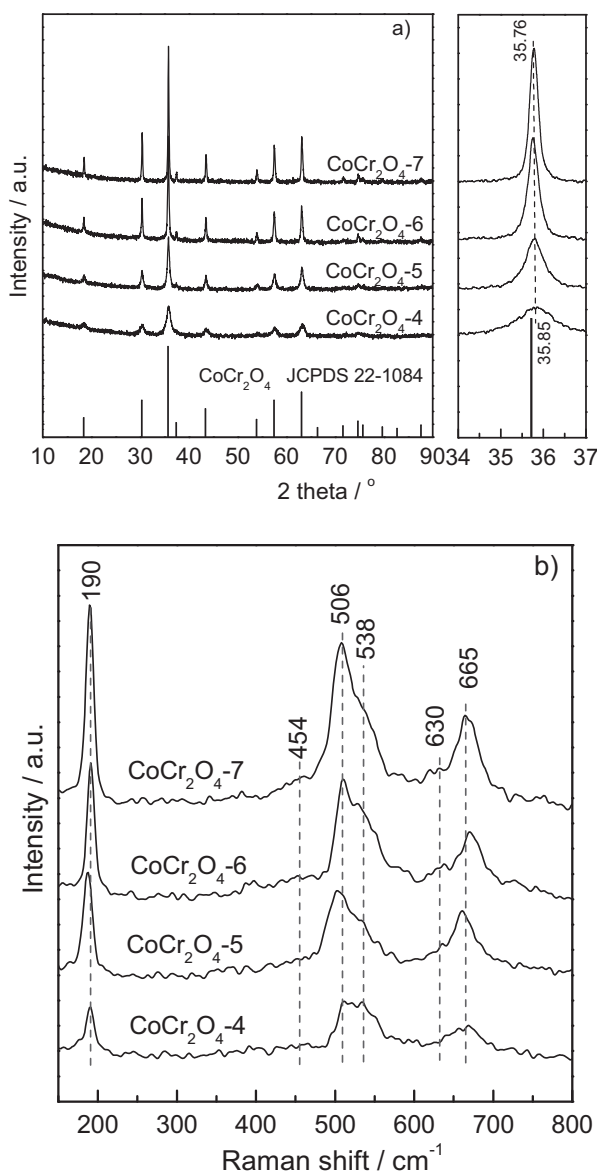


Fig. 2. (a) XRD patterns and (b) Raman spectra of spinel  $\text{CoCr}_2\text{O}_4$  catalysts.

calcination temperature, which consequently results in an increasing cell parameter of the sample (Table 1). The variation of lattice parameters could be due to the disordering of the cations in the spinel structures. At low-temperature calcination,  $\text{Co}^{3+}$  ions could partially replace  $\text{Cr}^{3+}$  in the oxide, which leads to a decline of lattice parameter because the ionic radius of the six-coordinated  $\text{Co}^{3+}$  in low-spin configuration (0.0545 nm) is smaller than that of the six-coordinated  $\text{Cr}^{3+}$  (0.0615 nm) [33]. However, when the sample was calcined at high temperature (e.g. 700 °C), the disordering of the cations in the spinel structure becomes less intense, which leads to a larger lattice parameter.

Raman spectra of the samples are shown in Fig. 2b. All these samples show four distinct bands at 190, 506, 538 and 665  $\text{cm}^{-1}$ , as well as two weak bands at 454 and 630  $\text{cm}^{-1}$ . According to group theory,  $\text{CoCr}_2\text{O}_4$  with purely normal spinel structure has five Raman-active phonon modes ( $\Gamma = 1\text{A}_{1g} + 1\text{E}_g + 3\text{F}_{2g}$ ) [34,35]. All of these modes can be observed at approximately 190  $\text{cm}^{-1}$  ( $\text{F}_{2g}$ ), 454  $\text{cm}^{-1}$  ( $\text{E}_g$ ), 506  $\text{cm}^{-1}$  ( $\text{F}_{2g}$ ), 538  $\text{cm}^{-1}$  (unclear mode), 630  $\text{cm}^{-1}$  ( $\text{F}_{2g}$ ), and 665  $\text{cm}^{-1}$  ( $\text{A}_{1g}$ ). The Raman spectra are similar as those reported in literature [36]. In addition, Chen et al. [36] attributed the band at 538  $\text{cm}^{-1}$  to a Raman-active mode which is due to the

Table 2

Surface compositions of spinel  $\text{CoCr}_2\text{O}_4$  catalysts.

| Catalyst                     | Atomic ratio                            |                                         |                                         |                                               |
|------------------------------|-----------------------------------------|-----------------------------------------|-----------------------------------------|-----------------------------------------------|
|                              | $\text{Co}^{3+}/\text{Co}_{\text{tot}}$ | $\text{Cr}^{3+}/\text{Cr}_{\text{tot}}$ | $\text{Cr}^{6+}/\text{Cr}_{\text{tot}}$ | $\text{O}_{\text{ads}}/\text{O}_{\text{lat}}$ |
| $\text{CoCr}_2\text{O}_4$ -4 | 0.23                                    | 0.81                                    | 0.19                                    | 0.57                                          |
| $\text{CoCr}_2\text{O}_4$ -5 | 0.20                                    | 0.80                                    | 0.20                                    | 0.53                                          |
| $\text{CoCr}_2\text{O}_4$ -6 | 0.17                                    | 0.81                                    | 0.19                                    | 0.52                                          |
| $\text{CoCr}_2\text{O}_4$ -7 | 0.15                                    | 0.81                                    | 0.19                                    | 0.50                                          |

cation disordering that induces a breakdown of the translation symmetry. Moreover, it is found that the intensities of the bands at 190, 506, and 665  $\text{cm}^{-1}$  gradually increase with increasing calcination temperatures. Such trends are due to the changes either in electronic properties [36] or the crystallite sizes [37,38] of the  $\text{CoCr}_2\text{O}_4$  oxides.

The oxidation states of the elements in the catalysts and surface compositions are analyzed by XPS, as shown in Fig. 3 and Table 2. For the Co 2p<sub>3/2</sub> spectra (Fig. 3a), a broad and overlapped band at binding energies (BEs) of 772–794 eV is observed in all samples, which could be further deconvoluted into three components with BEs centered at 780.8, 782.9 and 787.3 eV. The peaks with BEs at 780.8 and 782.9 eV could be assigned to  $\text{Co}^{2+}$  and  $\text{Co}^{3+}$ , respectively [39–41]. And the presence of  $\text{Co}^{2+}$  in the catalysts is also confirmed by the satellite peak at 787.3 eV [36]. The surface composition analyses also indicate that the concentration of  $\text{Co}^{3+}$  cations progressively decreases with increasing calcination temperature (23%  $\text{Co}^{3+}$  in the  $\text{CoCr}_2\text{O}_4$ -4 and 15% in the  $\text{CoCr}_2\text{O}_4$ -7, Table 2), which suggests that calcination temperature exerts influences on the oxidation states of the Co species in the samples. The higher  $\text{Co}^{3+}$  content in the low-temperature calcined sample (e.g.  $\text{CoCr}_2\text{O}_4$ -4) implies that some of these  $\text{Co}^{3+}$  cations could replace  $\text{Cr}^{3+}$  cations in the B site of the  $\text{AB}_2\text{O}_4$  spinel structure, which is consistent with the smaller lattice parameter in the  $\text{CoCr}_2\text{O}_4$ -4 sample (Table 1).

In Fig. 3b, the peak of Cr 2p<sub>3/2</sub> could also be resolved into three components at BEs at 575.7, 576.9 and 579.2, assigning to  $\text{Cr}(\text{OH})_3$  or  $\text{Cr}_2\text{O}_3$  [42],  $\text{Cr}^{3+}$  (occupied octahedral sites) [29,43,44] and  $\text{Cr}^{6+}$  [30,45], respectively. The occupied octahedral  $\text{Cr}^{3+}$  sites are exposed at the surface of the spinel samples and regarded as active sites for the oxidation reaction [29,43,44]. The presence of  $\text{Cr}^{6+}$  in spinel  $\text{CoCr}_2\text{O}_4$  oxides was reported by Sloczynski et al. [45], as the authors found that surface of Cr-containing spinel oxides was enriched with  $\text{Cr}^{6+}$ , and the location of  $\text{Cr}^{6+}$  was surface-only. Besides, Kim and Ihm [30] found that the  $\text{Cr}^{6+}$  was stable on the surface in  $\text{H}_2$  environment and was inactive for the TCE oxidation. Surface compositions of the Cr species in the catalysts (Table 2) reveal that the concentrations of various Cr species in these catalysts remain almost constant regardless of the calcination temperature.

As for the O1s spectra (Fig. 3c), the asymmetric peak could be deconvoluted to one main peak at BE of 530.4 eV and one weak peak at BE of 531.4 eV, which could be assigned to lattice oxygen ( $\text{O}_{\text{lat}}$ ) and adsorbed oxygen ( $\text{O}_{\text{ads}}$ ), respectively [46]. The  $\text{O}_{\text{ads}}$  peaks can be associated with a wide variety of species such as surface chemisorbed oxygen, hydroxyl and oxygen ions in low coordination situation and oxygen-containing surface contamination [47]. Moreover, the areal ratio of  $\text{O}_{\text{ads}}/\text{O}_{\text{lat}}$  reflects the proportion of the two different oxygen species, and a higher ratio indicates larger amount of chemisorbed oxygen species [48]. As shown in Table 2, the  $\text{O}_{\text{ads}}/\text{O}_{\text{lat}}$  ratio is relatively higher on the  $\text{CoCr}_2\text{O}_4$ -4 catalyst compared to those on the other catalysts, implying higher amount of surface oxygen on the  $\text{CoCr}_2\text{O}_4$ -4 catalyst.

Surface acidity plays an important role in CVOCs oxidation as they provide sites for the chemisorption of CVOCs molecules [3]. Thus, surface acidities of the catalysts are measured by  $\text{NH}_3$ -TPD,

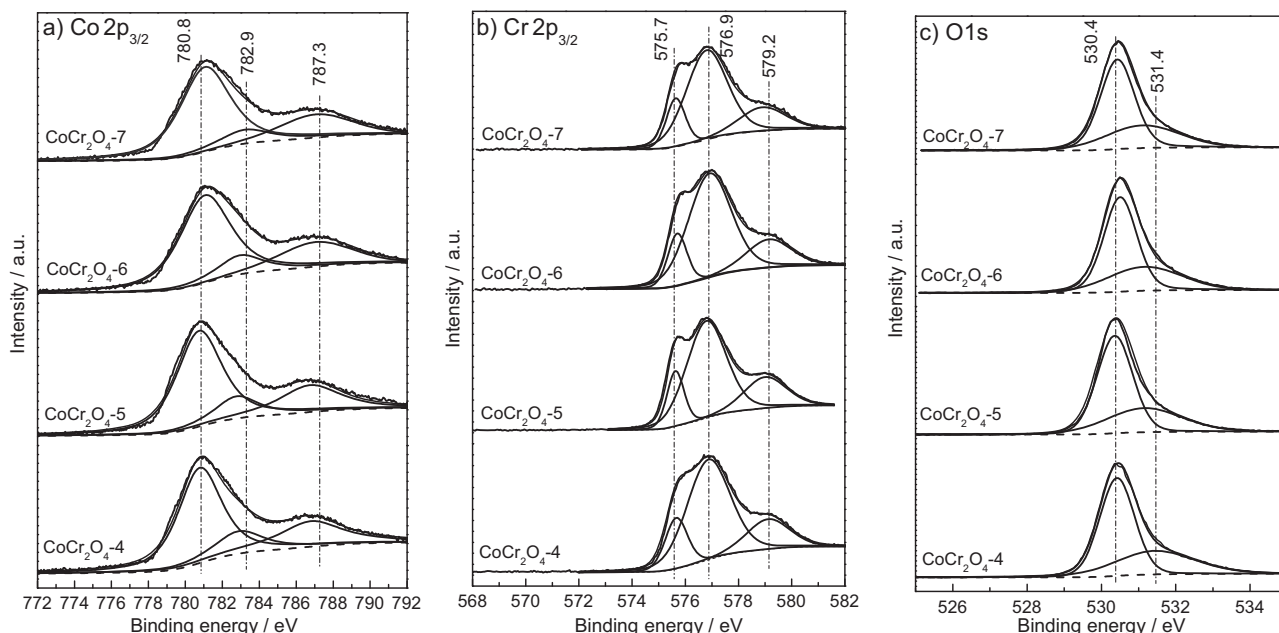


Fig. 3. Co2p, Cr2p and O1s XPS spectra of spinel  $\text{CoCr}_2\text{O}_4$  catalyst.

and the results are shown in Fig. 4a. One weak peak at less than  $200^\circ\text{C}$  and an intense peak at about  $210$ – $260^\circ\text{C}$  are detected in the catalysts, indicating the presence of surface acidic sites in the catalysts. The amounts of  $\text{NH}_3$  desorption on these samples are also quantified, which are  $0.43$ ,  $0.17$ ,  $0.10$  and  $0.06 \text{ mmol g}_{\text{cat}}^{-1}$  for the  $\text{CoCr}_2\text{O}_4$ -4,  $\text{CoCr}_2\text{O}_4$ -5,  $\text{CoCr}_2\text{O}_4$ -6 and  $\text{CoCr}_2\text{O}_4$ -7, respectively. The declined values clearly suggest that calcination temperature could remarkably influence the surface acidity of the cobalt chromites [49]. However, considering that the surface areas of these catalysts are very different, and the amount of surface acidic sites should be normalized based on surface area. It is found that

the normalized values of the catalysts are essentially constant ( $4.28$ – $4.92 \mu\text{mol m}^{-2}$ ).

Reducibility of the catalysts is measured by  $\text{H}_2$ -TPR technique, as shown in Fig. 4b. There are two reduction peaks at about  $250^\circ\text{C}$  ( $\alpha$  peak) and  $750^\circ\text{C}$  ( $\beta$  peak), which are attributed to the reduction of surface  $\text{Cr}^{6+}$  to  $\text{Cr}^{3+}$  species [31] and reduction of  $\text{Cr}^{3+}$  to lower oxidation states [29], respectively. The  $\text{H}_2$  consumptions of the  $\alpha$  peak of the catalysts are calculated to be  $0.734$ ,  $0.403$ ,  $0.186$  and  $0.051 \text{ mmol g}_{\text{cat}}^{-1}$  for the  $\text{CoCr}_2\text{O}_4$ -4,  $\text{CoCr}_2\text{O}_4$ -5,  $\text{CoCr}_2\text{O}_4$ -6 and  $\text{CoCr}_2\text{O}_4$ -7 catalysts, respectively. Note that the nominal  $\text{H}_2$  consumption of the catalyst is about  $2.04 \text{ mmol g}_{\text{cat}}^{-1}$  if we assume the

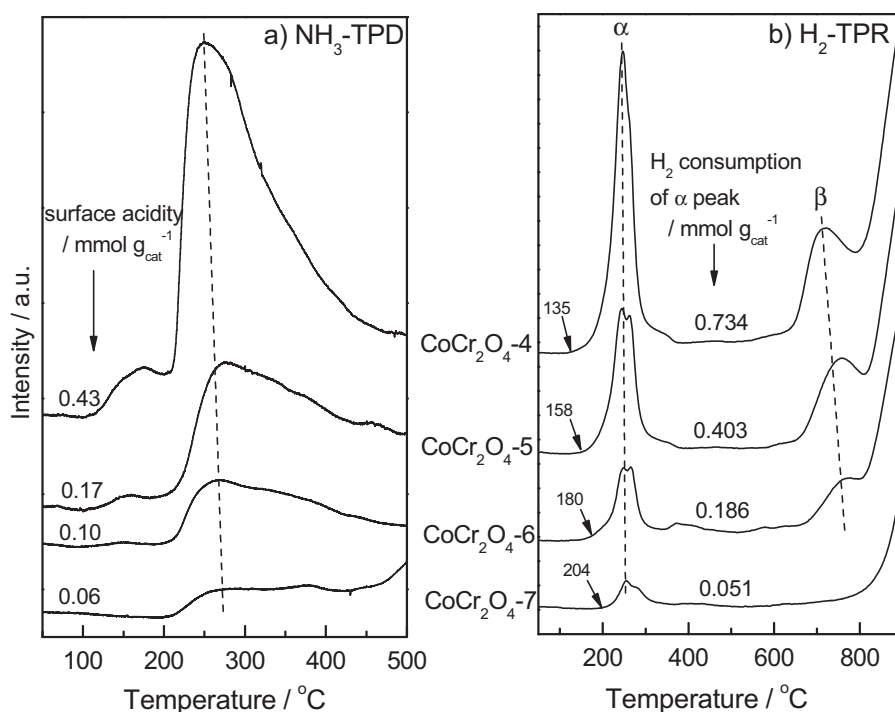


Fig. 4. (a)  $\text{NH}_3$ -TPD and (b)  $\text{H}_2$ -TPR profiles of spinel  $\text{CoCr}_2\text{O}_4$  catalysts.

$\text{Cr}^{6+}$  contents in these catalysts are about  $1.76 \text{ mmol g}_{\text{cat}}^{-1}$  based on the XPS results (Table 2). Thus, the actual  $\text{H}_2$  consumption values ( $0.734\text{--}0.051 \text{ mmol g}_{\text{cat}}^{-1}$ ) are much lower than the nominal ones, indicating that only partial  $\text{Cr}^{6+}$  cations could be reduced. Moreover, the higher  $\text{H}_2$  consumption of the  $\text{CoCr}_2\text{O}_4\text{-4}$  suggests more facile reduction of the low-temperature calcined sample probably because of its smaller nano-crystallites compared to others [50] even though the compositions of these catalysts are the same. The easier reduction of the  $\text{CoCr}_2\text{O}_4\text{-4}$  sample is also verified by the fact that the onset temperature of the  $\alpha$  peak is much lower on the  $\text{CoCr}_2\text{O}_4\text{-4}$  ( $135^\circ\text{C}$ ) compared to  $\text{CoCr}_2\text{O}_4\text{-7}$  ( $204^\circ\text{C}$ ). The assignment of the  $\alpha$  peak to the reduction of surface species is also evidenced by an analysis method of the TPR results developed by Zimmer et al. [51]. The authors proposed an equation:

$$\frac{N_{\text{total}}}{M_{\text{sample}}} = a + b \times \frac{A_{\text{sample}}}{M_{\text{sample}}}$$

where  $N_{\text{total}}$  is the  $\text{H}_2$  consumption of one certain peak,  $M_{\text{sample}}$  is the weight of the sample,  $A_{\text{sample}}$  is the surface area of the sample, and  $a$  and  $b$  are constants. If  $N_{\text{total}}/M_{\text{sample}}$  is plotted against  $A_{\text{sample}}/M_{\text{sample}}$  and a linear correlation could be established, then it could be concluded that the certain peak is due to the reduction of the surface. In the current work, the  $\text{H}_2$  consumptions of  $\alpha$  peak in these samples are plotted according to above equation. A perfect linear regression of the data yields an intercept  $a = 5.84 \mu\text{mol m}^{-2}$  and a slope  $b = 7.79 \mu\text{mol m}^{-2}$  (see Fig. S1 in Supplementary Information), which indicates that the  $\alpha$  peak in Fig. 4b is dominated by the reduction of the surface  $\text{Cr}^{6+}$  species.

### 3.2. Catalytic testing

The catalytic activities of the catalysts for  $\text{CH}_2\text{Cl}_2$  oxidation are measured as a function of the reaction temperature and the light-off curves are shown in Fig. 5a. The overall activities follow the order of  $\text{CoCr}_2\text{O}_4\text{-4} > \text{CoCr}_2\text{O}_4\text{-5} > \text{CoCr}_2\text{O}_4\text{-6} > \text{CoCr}_2\text{O}_4\text{-7}$ . To better compare the activities of these catalysts,  $T_{50}$  (the temperature at which a conversion of 50% is obtained) values are also listed in Fig. 5a. The  $T_{50}$  values are 217, 231, 259 and  $278^\circ\text{C}$  for the  $\text{CoCr}_2\text{O}_4\text{-4}$ ,  $\text{CoCr}_2\text{O}_4\text{-5}$ ,  $\text{CoCr}_2\text{O}_4\text{-6}$  and  $\text{CoCr}_2\text{O}_4\text{-7}$ , respectively. It is worth pointing out that the spinel  $\text{CoCr}_2\text{O}_4$  catalysts in the current work are very active for  $\text{CH}_2\text{Cl}_2$  oxidation, and these catalysts are superior to other reported catalyst systems such as  $\text{CrO}_3/\text{C}$  [52],  $\text{Mn}/\text{H-ZSM-5}$  [53],  $\text{Pd}/\text{CrZr}$  [54],  $\text{CrAlO}$  [55] and  $\text{K-Pt}/\text{Al}_2\text{O}_3$  [4] (see Table S1 in Supplementary Information). For example, the reaction rate on the  $\text{CoCr}_2\text{O}_4\text{-4}$  catalyst at  $250^\circ\text{C}$  is  $3.897 \text{ mmol}_{\text{CH}_2\text{Cl}_2} \text{ g}_{\text{cat}}^{-1} \text{ h}^{-1}$ , which is much higher than the rate obtained on a  $18\text{CrAlO}$  catalyst ( $0.241 \text{ mmol}_{\text{CH}_2\text{Cl}_2} \text{ g}_{\text{cat}}^{-1} \text{ h}^{-1}$ ) reported in our previous work [55].

It should be noticed that for CVOs oxidation, high conversion is not the only criterion for good catalyst systems, and selectivities of the products are more important. The desired products in Cl-containing CVOs oxidation should be  $\text{CO}_2$  and  $\text{HCl}$  or  $\text{Cl}_2$ . Therefore, TPSR over the catalysts were carried out and the product distributions are also analyzed on these spinel catalysts by MS, and the results are shown in Fig. 5b. As shown in Fig. 5b, the detectable final products over the  $\text{CoCr}_2\text{O}_4\text{-4}$  and  $\text{CoCr}_2\text{O}_4\text{-7}$  catalysts are  $\text{CO}_2$ ,  $\text{HCl}$ ,  $\text{Cl}_2$ , as well as very small amount of  $\text{HCHO}$ . No other Cl-containing organic compounds are detected. The product distributions are quite different from those obtained on  $\text{Al}_2\text{O}_3$ -based catalysts, on which Cl-containing organic compounds such as  $\text{CH}_3\text{Cl}$  are always formed [56]. However, the formation of  $\text{CO}$  during the reaction could not be ruled out because of its same  $m/e$  signal ( $m/e = 28$ ) as that of  $\text{N}_2$ . Besides, it can be seen that the onset temperatures of the two catalysts are different ( $260^\circ\text{C}$  for the  $\text{CoCr}_2\text{O}_4\text{-4}$  and  $300^\circ\text{C}$  for the  $\text{CoCr}_2\text{O}_4\text{-7}$ ), and concentrations of the products are much larger on the  $\text{CoCr}_2\text{O}_4\text{-4}$  compared to those on the  $\text{CoCr}_2\text{O}_4\text{-7}$ , judging from the peak areas of the products. These

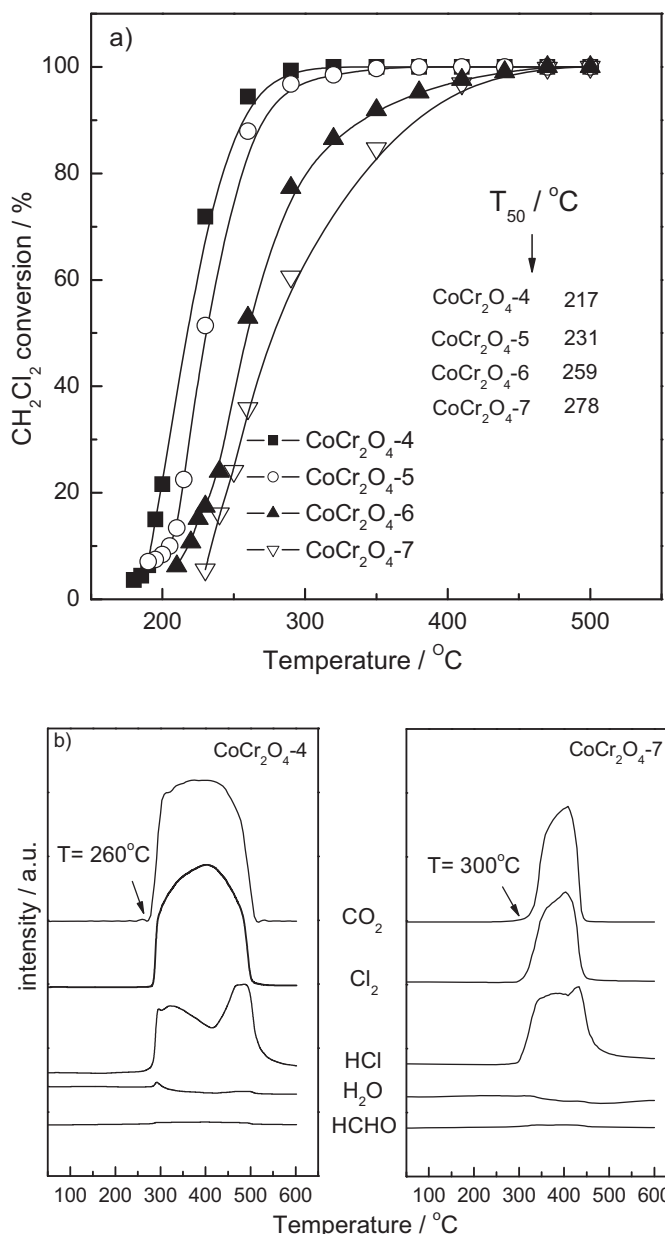


Fig. 5. (a) Light-off curves and (b) product distributions of  $\text{CH}_2\text{Cl}_2$  oxidation over  $\text{CoCr}_2\text{O}_4\text{-4}$  and  $\text{CoCr}_2\text{O}_4\text{-7}$  catalysts.

results again confirm that the  $\text{CoCr}_2\text{O}_4\text{-4}$  catalyst is more active than the  $\text{CoCr}_2\text{O}_4\text{-7}$ .

The stability of the catalysts is also investigated. As can be seen in Fig. 6, the catalysts are quite stable during 10 h reaction. Note that in order to obtain similar  $\text{CH}_2\text{Cl}_2$  conversions (about 80–90%) over these catalysts, different reaction temperatures were used (for the  $\text{CoCr}_2\text{O}_4\text{-4}$  and  $\text{CoCr}_2\text{O}_4\text{-5}$  the reaction temperature was  $250^\circ\text{C}$ , while for the  $\text{CoCr}_2\text{O}_4\text{-6}$  and  $\text{CoCr}_2\text{O}_4\text{-7}$  the reaction temperature was  $350^\circ\text{C}$ ). The bulk compositions and surface areas of the spent catalysts are very close to those of the fresh samples. For example, the spent  $\text{CoCr}_2\text{O}_4\text{-4}$  has a surface area of  $93.1 \text{ m}^2 \text{ g}^{-1}$  and a Cr content of 44.2 wt.%, which are close to the values of the fresh  $\text{CoCr}_2\text{O}_4\text{-4}$ . Moreover, comparison of the catalysts properties (XRD and XPS results) before and after reaction reveals that the catalysts remain unchanged after reaction (see Figs. S2 and S3 and Table S2 in Supplementary Information). The stable  $\text{CoCr}_2\text{O}_4$  catalysts are also probably related to the spinel structure, in which the Co and Cr species are confined in a relatively robust matrix. Another

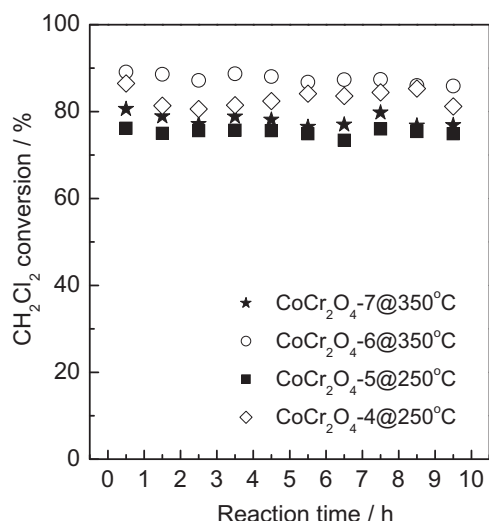


Fig. 6. Stability of spinel  $\text{CoCr}_2\text{O}_4$  catalysts at different reaction temperatures.

noteworthy point is the possible formation of toxic chromium oxychloride such as  $\text{Cr}(\text{ClO})_2$  during the reaction. In the current work, since the Cr content and the physical/chemical properties of the spent sample do not change, it is quite safe to conclude that there is no such chromium oxychloride formed during the reaction. Besides, chromium oxychloride has a color of dark red, which has not been observed in the current work.

It is generally recognized that surface acidity and catalyst reducibility are two important parameters for CVOs oxidation because the surface acidic sites provide centers for CVOs molecule chemisorption, while the enhanced reducibility of the catalyst is beneficial to the activation of oxygen species and lattice oxygen of the oxides could often directly participate in the reaction [3]. The activity of the catalyst is usually determined by synergistic effects of these two parameters [4,10]. Thus, relationship between catalyst reducibility, surface acidity and catalytic performance is plotted as shown in Fig. 7. As shown in Fig. 7a, the specific reaction rate in unit of  $\mu\text{mol g}_{\text{cat}}^{-1} \text{s}^{-1}$  (data taken from

Fig. 5a at the reaction temperature of  $230^\circ\text{C}$ ) is proportional to both the surface acidic sites (data taken from Fig. 4a) and the surface reducibility (data taken from Fig. 4b), which strongly suggests that the overall activity of the catalyst is directly related to these two parameters. However, considering that these catalysts have very different surface areas, these variables (reducibility, surface acidity and reaction rate) should be normalized based on surface area, and the correlation is shown in Fig. 7b. It is found that the areal reaction rates are essentially constant, ranging from  $0.82\text{--}1.09 \times 10^{-8} \text{ mol m}^{-2} \text{s}^{-1}$ . These rates are in agreement with the essentially constant surface acidity ( $4.31\text{--}4.92 \text{ mmol}_{\text{NH}_3} \text{m}^{-2}$ ) and reducibility ( $7.76\text{--}8.04 \text{ mmol}_{\text{H}_2} \text{m}^{-2}$ ). These results indicate that the enhanced activity of the  $\text{CoCr}_2\text{O}_4\text{-4}$  is mainly due to its higher surface area, as the catalyst with high surface area certainly contains more active sites compared to that with low surface area.

### 3.3. Mechanistic investigation

For spinel type oxides ( $\text{AB}_2\text{O}_4$ ) in catalytic reactions, it is recognized that the octahedral B-site cations are the active sites because these cations are exposed at the surface while the tetrahedral A-site cations are usually inactive [29,43]. In the case of  $\text{CoCr}_2\text{O}_4$  system, Cr species in B site play very essential roles in the reaction, as reported in previous work [30]. In the current work, the Cr species in the  $\text{CoCr}_2\text{O}_4$  catalysts provide surface acidic sites for  $\text{CH}_2\text{Cl}_2$  chemisorption and active oxygen species (either adsorbed oxygen or lattice oxygen) as confirmed by the  $\text{NH}_3$ -TPD (Fig. 4a) and  $\text{H}_2$ -TPR (Fig. 4b) results, which are two important parameters governing the catalytic activity.

Since the  $\text{CoCr}_2\text{O}_4$  catalysts employed in the current work have very different activities, it is possible that reaction pathways might differ on these catalysts. To clarify this point, kinetic studies were performed on these catalysts and the Arrhenius plots are shown in Fig. 8. Note that the  $\text{CH}_2\text{Cl}_2$  conversions are low (typically less than 20%) to ensure a differential reaction mode on the catalyst (see Table S3 for the detailed reaction rates and the verification of the absence of mass and heat transfer limits in Supplementary Information). It is found that the apparent activation energies obtained on these catalysts are  $124.5\text{--}155.5 \text{ kJ mol}^{-1}$ , which are comparable

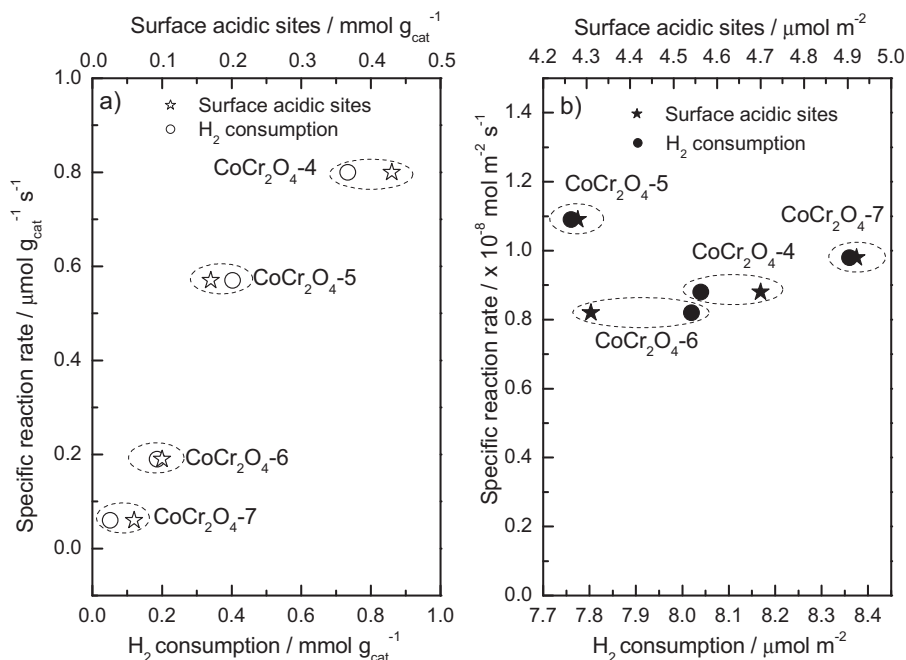


Fig. 7. Correlation between (a) overall and (b) normalized surface acidity, catalyst reducibility and specific reaction rate of  $\text{CoCr}_2\text{O}_4$  catalysts.

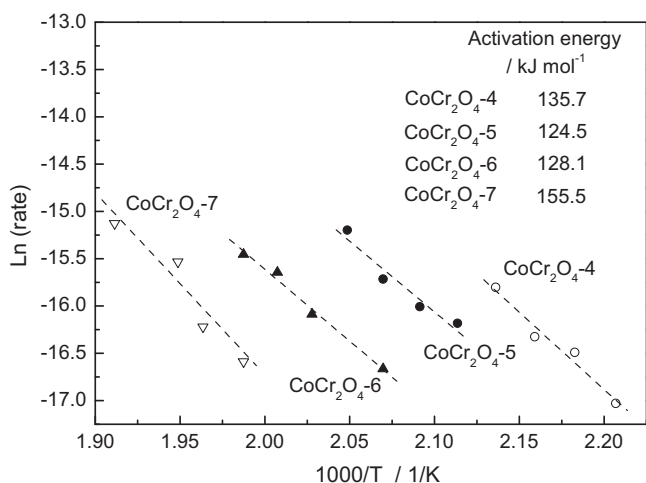


Fig. 8. Arrhenius plots of  $\text{CH}_2\text{Cl}_2$  oxidation over spinel  $\text{CoCr}_2\text{O}_4$  catalysts.

to the values reported on  $\text{Pt}/\text{Al}_2\text{O}_3$  catalysts [4]. The similar activation energies on these spinel catalysts suggest that the reaction pathways are likely unchanged in spite of their different overall activities.

In order to further understand the catalytic behaviors of these  $\text{CoCr}_2\text{O}_4$  catalysts, in situ FTIR experiments were conducted and the results are shown in Fig. 9 (FTIR spectra of the  $\text{CoCr}_2\text{O}_4$ -5 and  $\text{CoCr}_2\text{O}_4$ -6 are shown in Fig. S4 in Supplementary Information). For the  $\text{CoCr}_2\text{O}_4$ -4 catalyst (Fig. 9a), after it is exposed to the air +  $\text{CH}_2\text{Cl}_2$  mixture at  $50^\circ\text{C}$ , bands at  $3006$ ,  $2985$ ,  $2365$ ,  $2332$ ,  $1610$   $\text{cm}^{-1}$  are observed. The bands at  $3006$  and  $2985$   $\text{cm}^{-1}$  are attributed to asymmetric vibration ( $\nu_{\text{as}}$ ) of  $-\text{CH}_2$  group in  $\text{CH}_2\text{Cl}_2$  molecule [56]; the bands at  $2365$  and  $2332$   $\text{cm}^{-1}$  are attributed to gas phase  $\text{CO}_2$ ; the band at  $1610$   $\text{cm}^{-1}$  is assigned to  $\text{H}_2\text{O}$  adsorbed on the catalyst surface, which gradually disappears at elevated temperature. When the catalyst is heated, the bands at  $3006$  and  $2985$   $\text{cm}^{-1}$  gradually decline in intensities with increasing temperature, implying either desorption or consumption of the  $\text{CH}_2\text{Cl}_2$  molecules. Meanwhile, the disappearance of  $\text{CH}_2\text{Cl}_2$  is accompanied by increasing intensities of the bands at  $2365$  and  $2332$   $\text{cm}^{-1}$  ( $\text{CO}_2$ ), and evolution of new bands at  $2178$ ,  $2120$ ,  $1540$ ,  $1443$  and  $1355$   $\text{cm}^{-1}$ . The weak bands at  $2178$  and  $2120$   $\text{cm}^{-1}$  are due to the formation of  $\text{CO}$ , while the bands at  $1540$ ,  $1433$  and  $1355$   $\text{cm}^{-1}$  are assigned to asymmetric vibration ( $\nu_{\text{as}}$ ),  $\delta$  ( $\text{CH}_2$ ) and symmetric vibration ( $\nu_{\text{s}}$ ) of formate species ( $-\text{COOH}$ ), respectively [4,11,57,58]. Similar observations are also detected on the  $\text{CoCr}_2\text{O}_4$ -5 (Fig. S4),  $\text{CoCr}_2\text{O}_4$ -6 (Fig. S4) and  $\text{CoCr}_2\text{O}_4$ -7 (Fig. 9b) catalysts.

The in situ FTIR results on the  $\text{CoCr}_2\text{O}_4$  catalysts reflect two facts. One fact is that  $\text{CH}_2\text{Cl}_2$  oxidation over the spinel  $\text{CoCr}_2\text{O}_4$  catalyst may follow similar pathways as over  $\text{Al}_2\text{O}_3$  catalyst [56], with the formate species being the main intermediate. The formate species could be decomposed to  $\text{CO}$ , which is further oxidized to  $\text{CO}_2$ . The other fact is the formation of  $\text{CO}$  during the reaction. The formation of  $\text{CO}$  is not detected in the TPSR process (Fig. 5b), its presence is clearly evidenced by the FTIR results (Fig. 9). Although the formation of  $\text{CO}$  is not desirable in CVOCs oxidation, it may provide a clue of the development of better catalyst system in the future. For example, the spinel oxides could be used as supports for some metals such as  $\text{Cu}$  and  $\text{Pt}$ , which are known to be very active in  $\text{CO}$  oxidation.

To conveniently compare the IR results on these catalysts, normalized intensities of the characteristic bands at  $1540$   $\text{cm}^{-1}$  (formate species),  $2121$   $\text{cm}^{-1}$  ( $\text{CO}$ ) and  $2356$   $\text{cm}^{-1}$  ( $\text{CO}_2$ ) as a function of temperature were plotted and shown in Fig. 10. For each catalyst, the intensity of formate species reaches a maximum at

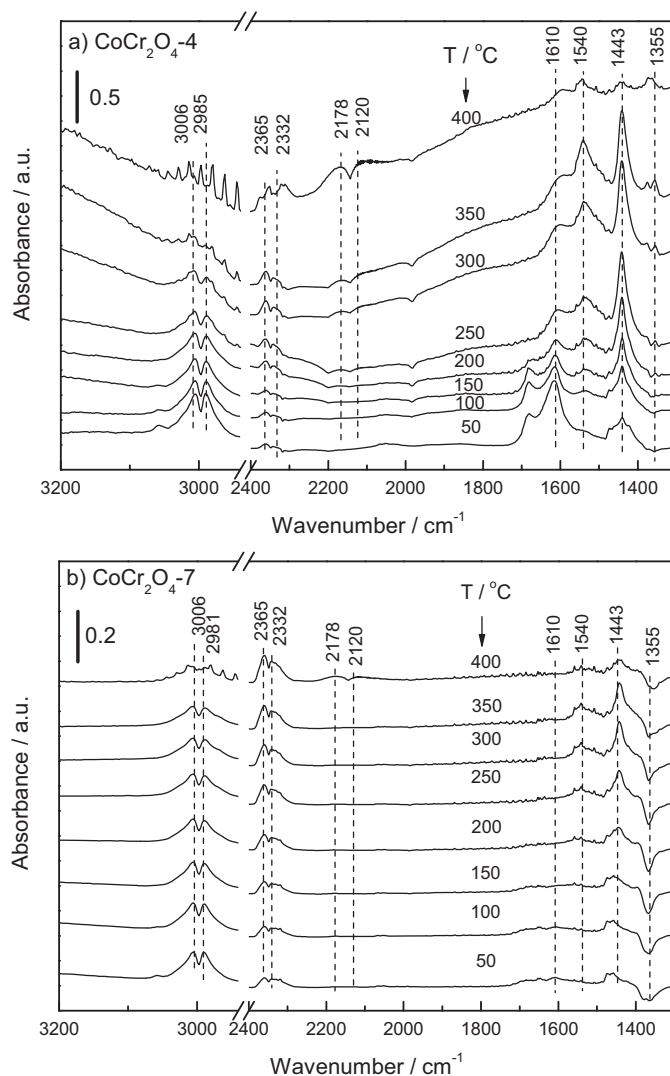
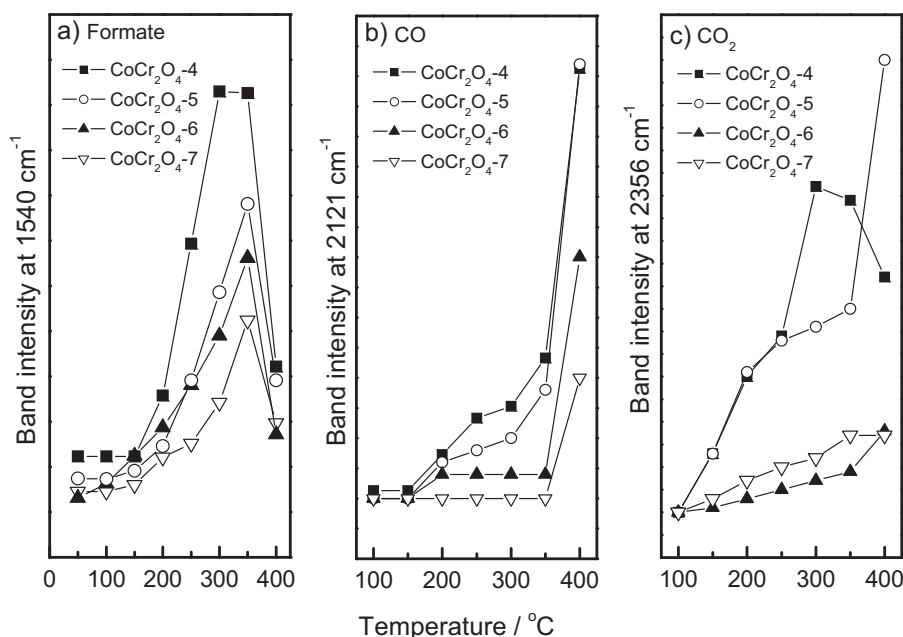


Fig. 9. In situ FTIR spectra of surface reaction of  $\text{CH}_2\text{Cl}_2$  + air over (a)  $\text{CoCr}_2\text{O}_4$ -4 and (b)  $\text{CoCr}_2\text{O}_4$ -7 catalysts.

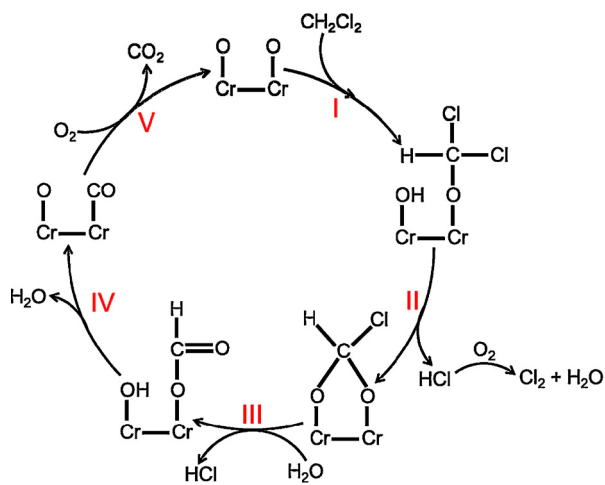
about  $350^\circ\text{C}$  and then declines (Fig. 10a). Meanwhile, the intensities of the bands at  $2120$  and  $2365$   $\text{cm}^{-1}$  progressively increase (Fig. 10b and c), and a rapid increase of the intensity is observed at about  $350^\circ\text{C}$ . These results suggest that the formate species could possibly accumulate on the catalyst surface at low temperature, and then be decomposed to  $\text{CO}_x$  at higher temperature. For the catalysts calcined at different temperatures, the intensities of the bands at  $1540$   $\text{cm}^{-1}$  (formate species),  $2121$   $\text{cm}^{-1}$  ( $\text{CO}$ ) and  $2356$   $\text{cm}^{-1}$  ( $\text{CO}_2$ ) all follow the order of  $\text{CoCr}_2\text{O}_4$ -4 >  $\text{CoCr}_2\text{O}_4$ -5 >  $\text{CoCr}_2\text{O}_4$ -6 >  $\text{CoCr}_2\text{O}_4$ -7. This comparison suggests that the  $\text{CoCr}_2\text{O}_4$ -4 is more active than the others, which is in good agreement with the overall activities of the catalysts (Fig. 5a).

Based on the FTIR spectra and reported mechanisms of  $\text{CH}_2\text{Cl}_2$  oxidation over various oxides [56,59], possible reaction pathways of  $\text{CH}_2\text{Cl}_2$  oxidation over the  $\text{CoCr}_2\text{O}_4$  catalyst could be demonstrated as shown in Scheme 1. The reaction pathways mainly consist of two sequential cycles: formation of  $\text{HCHO}$  and oxidation of  $\text{HCHO}$  to  $\text{CO}$  and  $\text{CO}_2$ . The reaction steps are described as follows:

**STEP 1.** A  $\text{CH}_2\text{Cl}_2$  molecule could chemisorb on two adjacent  $\text{Cr}-\text{O}$  sites which serve as acidic sites to form a  $\text{Cr}-\text{OH}$  specie and a  $\text{Cr}-\text{OCHCl}_2$  specie.



**Fig. 10.** Intensities of characteristic bands as a function of temperature over different catalysts. (a) Band at 1540 cm<sup>-1</sup> (formate species); (b) band at 2121 cm<sup>-1</sup> (CO); (c) band at 2356 cm<sup>-1</sup> (CO<sub>2</sub>).



**Scheme 1.** Possible reaction pathways of CH<sub>2</sub>Cl<sub>2</sub> oxidation over spinel CoCr<sub>2</sub>O<sub>4</sub> catalyst.

**STEP 2.** The Cr–OH and Cr–OCHCl<sub>2</sub> species decompose to HCl and a chloromethoxyl species (HCO<sub>2</sub>Cl). HCl could further react with O<sub>2</sub> via Deacon reaction (2HCl + 1/2 O<sub>2</sub> → Cl<sub>2</sub> + H<sub>2</sub>O), which could explain the formation of Cl<sub>2</sub> during the reaction.

**STEP 3.** The chloromethoxyl species reacts with H<sub>2</sub>O to form a HCl molecule and a Cr–OH and species a Cr–COOH (formate) species. Note that H<sub>2</sub>O comes from the moisture-containing feed gas. The chloromethoxyl species could also react with H<sub>2</sub>O to form HCl + HCHO molecules. The formation of HCHO is evidenced in TPSR results (Fig. 5b).

**STEP 4.** The Cr–OH and Cr–COOH species react to form a H<sub>2</sub>O molecule and a chemisorbed CO (Cr–CO) and a Cr–O site.

**STEP 5.** The Cr–CO could further react with O<sub>2</sub> to form a CO<sub>2</sub> molecule and thus complete the reaction cycle.

#### 4. Conclusion

Although spinel type oxides have been widely investigated and CoCr<sub>2</sub>O<sub>4</sub> oxide was characterized in detail in literature [30], some new information has been obtained in the current work. Firstly, CoCr<sub>2</sub>O<sub>4</sub> catalysts were found to be very active for CH<sub>2</sub>Cl<sub>2</sub> oxidation, which has not been reported before. The best catalytic performance is obtained on the catalyst calcined at 400 °C (CoCr<sub>2</sub>O<sub>4</sub>-4). Secondly, quantitative analyses of the catalytic results in the current work established correlation between the specific reaction rate and some important catalyst properties such as surface area, reducibility and surface acidity. The enhanced activity on this catalyst could be attributed to its highest surface area compared to those calcined at higher temperatures, as it provides more surface acidic sites and active oxygen species than the other catalysts. Thirdly, kinetic study revealed similar activation energies on these catalysts, implying the same reaction pathways of this reaction. Moreover, some reaction intermediates such as formate species have been detected by in situ FTIR, based on which a possible reaction mechanism was proposed.

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#### Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.apcatb.2014.10.044>.

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