Catalytic Ozonation of Herbicide 2,4-D over Cobalt Oxide Supported on Mesoporous Zirconia

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Cobalt oxide was supported on ordered mesoporous zirconia (MZ) by wet impregnation, drying, water washing, and calcinations with cobalt acetate tetrahydrate as the metal precursor for the first time. The material ($CoO_x/MZIW$) was characterized by X-ray diffraction, X-ray photoelectron spectroscopy, Fourier transform infrared spectroscopy, temperature-programmed reduction, and UV-vis diffuse reflectance spectra measurement. The studies showed that the $[Co(H_2O)_6]^{2+}$ interacted strongly with surface hydroxyls of mesoporous zirconia, leading to a highly dispersed CoO_x layer. The CoO_x existed mainly as Co_3O_4 phase. Furthermore, the catalyst was found to be highly effective for the mineralization of 2,4-dichlorophenoxyacetic acid (2,4-D) aqueous solution with ozone. The multivalence oxidation states and high dispersion of CoO_x enhanced the interfacial electron transfer, causing the higher catalytic reactivity. On the basis of all information obtained under different experimental conditions, $CoO_x/MZIW$ enhanced the mineralization of 2,4-D by the formation of the hydroxyl radical (*OH) resulting from the catalytic decomposition of ozone.

1. Introduction

Heterogeneous catalytic ozonation has received increasing attention in recent years because of its potentially higher effectiveness in the degradation and mineralization of refractory organic pollutants and lower negative effect on water quality. It has been developed to overcome the limitations of ozonation processes, such as the formation of byproducts and selective reactions of ozone, which are designed to enhance the production of *OH, known nonselective oxidants. 1,2 Specifically, solid metal oxides are more practical in catalytic ozonation than ionized metals, due to the fact that solid metal oxides subdue bromate formation in the reaction with ozone and are less pH-sensitive than ionized metals. 3,4

So far, metal oxides (e.g., MnO₂, TiO₂, and Al₂O₃) and supported metal oxides have been proposed as effective catalysts for ozonation processes. $^{5-7}$ In many cases, alumina-supported metal oxides of Fe, Ag, Co, Ni, Mn, and Cu have shown high activity for ozone decomposition at ambient temperature. Among transition metal oxides, cobalt oxides, both unsupported and supported on different oxide support materials, are some of the most important catalysts for complete oxidation of volatile organic compounds in air. A large number of papers have reported their high catalytic activity for the elimination of CO, NO_x, 11 and organic compounds 12 in air. However, these catalysts have not yet been applied much for the catalytic ozonation of organic pollutants in water. 13

The catalytic activity of metal oxide depends on the size distribution and morphology of particles. Hence, reducing the diameter of heterogeneous catalysts to nanometer scale may cause the enhanced reactivity of the heterogeneous catalysts at a given amount.¹⁴ Presently, using porous materials as solid nanoreactors, MnO_x nanocrystals with various morphologies have been obtained by a method called "nanocasting". Generally,

inorganic precursors are first introduced into channels of host materials such as mesoporous silicas. Then, MnO_x nanocrystals are produced with a replica of the confined space by subsequent calcinations. Nanoparticles of MnO_x supported on mesoporous silica SBA-15 have been obtained by high dispersion. The structure and dispersion of the supported metal oxide depend primarily on the preparation method, the nature of the support, and the type of precursor itself. Ordered mesoporous materials, with their intrinsically high surface areas, are particularly suitable for this purpose.

In the present study, for the first time, CoO_x nanoparticles were highly dispersed on the surface of MZ by the impregnation of the acetate precursor solution, followed by drying at 383 K and washing with water and calcinations at 573-773 K. This was attributed to the strong interaction of the $[Co(H_2O)_6]^{2+}$ with surface hydroxyls of MZ on the basis of different experimental information. 2,4-D is the most widely used herbicide in the world. 16 It has been detected as a major pollutant in ground and surface waters. Moreover, 2,4-D has very poor biodegradability. Most papers describe the degradation of 2,4-D aqueous solution by advanced oxidation processes (AOPs) involving chemical, ¹⁷ photochemical, ¹⁸ or photocatalytic ¹⁹ production of OH. In these processes, the mineralization of 2,4-D needs a prolonged reaction time. Catalytic ozonation has great potential for the mineralization of refractory organics. Therefore, 2,4-D was selected to evaluate the activity and properties of the catalyst with ozone in aqueous solution. The results indicated that the catalyst exhibited higher reactivity for the mineralization of 2,4-D. A preliminary effort to identify a correlation between the surface properties of supported CoO_x and their catalytic activity has been undertaken.

2. Experimental Section

Catalyst Preparation. MZ was prepared via solid-state reaction using the structure-directing method.²⁰ In this synthesis, ZrOCl₂*8H₂O and NaOH were milled into fine powder,

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respectively, and mixed at ambient temperature with the assistance of P123. The mixture was then transferred into an autoclave and kept at 383 K for 48 h. Subsequently, the mixture was washed with deionized water and ethanol, then dried at 383 K overnight and calcined at 623 K for 3 h. Because MZ has ordered mesoporous and a bigger BET surface area (232 m²/g) with sufficient hydroxyl groups, it is a good support for preparing the highly dispersed metal oxide. Different supported CoO_x samples were prepared by the incipient wetness impregnation method with cobalt acetate tetrahydrate [Co(CH₃COO)₂• 4H₂O] as the metal precursor. A 0.33-g quantity of cobalt acetate tetrahydrate was dissolved in 1 mL of distilled water. Then, 2 g of MZ was added to this solution. After impregnation, the sample was dried at 110 °C for several hours. Subsequently, the product was washed with deionized water to remove the excess cobalt, which was not connected directly to the surface of MZ by the hydroxyl group, and dried at 110 °C for several hours and calcined at 350 °C for 2 h, in air. Following this procedure, catalysts with different Co contents were prepared from 1 to 20 wt %. The catalyst with 2 wt % Co exhibited the highest activity. This catalyst, designated CoO_x/MZIW, was used for all of the experiments. As reference, two catalysts, 2 wt % Co supported on mesoporous zirconia and commercial ZrO2 (CZ) (purchased from Shantou Xilong Chemical Factory Guangdong, China), were prepared by the above method without water washing. These two catalysts were designated CoO_x/MZI and CoO_x/CZI.

Catalyst Characterization. Powder X-ray diffraction of the catalyst was recorded on a Scintag-XDS-2000 diffractometer with Cu K α radiation ($\lambda = 1.54059$ Å). UV-vis diffuse reflectance spectra (UV-vis DRS) of the samples were recorded on a UV-vis spectrophotometer (Hitachi UV-3100) with an integrating sphere attachment. The analyzed range was 200-800 nm, and BaSO₄ was the reflectance standard. XPS analysis was performed on an AXIS-Ultra instrument from Kratos, using monochromatic Al Ka radiation (225 W, 15 mA, 15 kV). To compensate for surface charge effects, binding energies were calibrated using a C1s hydrocarbon peak at 284.8 eV. For FTIR measurement, the dry samples were supported on KBr pellets. The infrared spectrum was recorded on a Nicolet FTIR spectrophotometer. The zeta potential of catalysts in KNO_3 (10^{-3} M) solution were measured with a Zetasizer 2000 (Malvern Co., U.K.). Every reading of the instrument was recorded after three consistent readings had been attained. The temperatureprogrammed reduction (TPR) experiments were carried out in a flow of 5% H₂/Ar (30 mL/min) by heating the sample to 900 °C at 10 °C min⁻¹. A mass spectrometer (Hiden) was used for on-line monitoring of the TPR effluent gas. Prior to TPR, the catalyst (0.5 g) was pretreated under a 20% O₂/He mixture at 350 °C for 1 h and cooled to room temperature.

Catalytic Activity Measurements. Batch experiments were carried out with a 1.2-L reactor. The reaction temperature was maintained at 20 °C. In a typical experiment, aqueous suspensions of 2,4-D (1 L, 80 mg L⁻¹) and 1.5 g of catalyst powders were placed in the reactor. The suspension was magnetically stirred throughout this experiment. The ozone was generated by a laboratory ozonizer (DHX-SS-IG, Harbin Jiujiu Electrochemistry Technology Co., Ltd., China). Thirty milligrams of gaseous O₃/L oxygen-ozone was bubbled into the reactor through the porous plate of the reactor bottom at a 12 L h⁻¹ flow rate. At given time intervals, samples were withdrawn and filtered through a Millipore filter (pore size 0.45 µm) for analysis. An aliquot of 0.1 M Na₂S₂O₃ was subsequently added to the sample to quench the aqueous ozone remaining in the

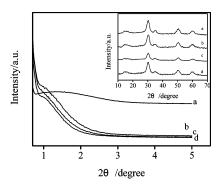


Figure 1. XRD patterns of different catalysts: (a) MZ, (b) 1% CoO_x/ MZIW, (c) 2% CoO_x/MZIW, (d) CoO_x/MZI. The inset shows the highangle peak.

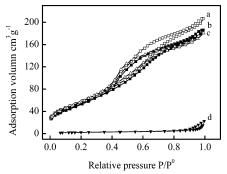


Figure 2. N₂ adsorption—desorption isotherms: (a) CoO_x/MZIW, (b) CoO_x/MZI, (c) MZ, (d) CoO_x/CZI.

reaction solution. The concentration of ozone in the aqueous phase was determined with the indigo method. 2,4-D was measured by high-performance liquid chromatography (HPLC, Alliance 2695) with an Xterra C18 column. Sixty percent acetonitrite with water mobile phase was used. The total organic carbon (TOC) of the solution was analyzed with a Phoenix 8000 TOC analyzer.

3. Results and Discussion

XRD and N₂ Adsorption. The XRD patterns of different samples are shown in Figure 1. The low-angle XRD pattern of the MZ sample showed a broader primary diffraction peak, indicating that the prepared MZ had an ordered mesostructure.²¹ The inorganic framework was periodic because it consisted of nanocrystallites in the tetragonal phase according to the XRD peaks at high angles shown in the inset of Figure 1. After the introduction of Co, the low-angle XRD peaks of MZ shifted toward higher angles, although MZ still had a periodic mesoporous framework. This was possibly attributed to the formation of Co nanoparticles in the pores. No XRD diffraction peaks of CoO_x were observed in these samples. However, the peak intensity of MZ was lowered with the increase of the loading amount of CoO_x for $CoO_x/MZIW$ (curves b and c), while it was not changed by the addition of CoO_x for CoO_x/MZI. The results indicated that the Co ions were incorporated into the structure of MZ for CoO_x/MZIW, whereas the Co ions were not incorporated into the crystal phase of MZ for CoOx/MZI. The XRD patterns of CoO_x/CZI and CZ were same (Supporting Information, Figure S1). No XRD diffraction peaks of CoO_x were observed in CoO_x/CZI because of its low content.

The N₂ adsorption-desorption isotherms are presented in Figure 2 for MZ, CoO_x/MZIW and CoO_x/MZI. They are typically type-IV with hysteresis loops, which means that the materials had a mesoporous structure. The introduction of CoO_x

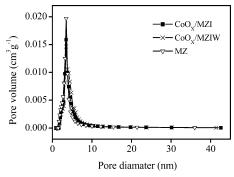


Figure 3. Pore size distribution for CoO_x/MZIW, CoO_x/MZI, and MZ.

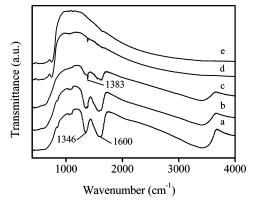


Figure 4. FTIR spectra of different samples: (a) MZ, (b) CoO_x/MZI, (c) CoO_x/MZIW, (d) CoO_x/ CZI, (e) CZ.

did not change the distribution of the pore diameters (shown in Figure 3), indicating the high dispersion of CoO_x . However, the BET surface area of $CoO_x/MZIW$ was 206 m²/g, smaller than that of MZ, while the BET surface area of CoO_x/MZI was 193 m²/g, a significant reduction. As a reference, CZ was also characterized by N_2 adsorption—desorption. Its BET surface area was 8.83 m²/g. No mesopores were detected in the sample (curve d) in Figure 2.

FTIR Measurements. The surface properties of a catalyst are very important for the degradation of organic pollutants in heterogeneous catalytic ozonation.²² Therefore, different catalysts were further characterized with FTIR. As shown in Figure 4 for MZ (Figure 4a), there were three OH absorption bands: the first one was at 3500-3200 cm⁻¹ corresponding to the stretching of OH groups of adsorbed water. The second one at 1600 cm⁻¹ was assigned to the bending vibration of adsorbed water.23 The third one was at 1346 cm-1, attributed to the deformation vibration of Zr-OH.²⁴ CoO_x/MZI and CoO_x/MZIW exhibited similar FTIR spectra of MZ except for a new absorption peak at 1386 cm⁻¹, assigned to OH deformation vibrations of hydrated CoO_x (Figure 4b and c). However, for CoO_y/MZIW, the intensity of the new peak was much stronger, and these bands at 1346 and 1600 cm⁻¹ became weaker than that for MZ. Oppositely, these bands did not significantly change in the CoO_x/MZI sample. The results indicated that CoO_x strongly interacted with surface hydroxyls of MZ in CoO_x/ MZIW, which agreed with the XRD results. The water washing process enhanced the diffusing of Co(II) on the surface of MZ, leading the strong interaction. There were no OH groups on CZ (Figure 4e), except the bands corresponding to a weak OH stretching vibration (3400 cm⁻¹). CoO_x/CZI also exhibited a weak peak at 1386 cm⁻¹, assigned to OH deformation vibrations of hydrated CoO_x.

XPS and UV–Vis DRS Analysis. The dispersion of CoO_x on different samples was further studied by the distribution of

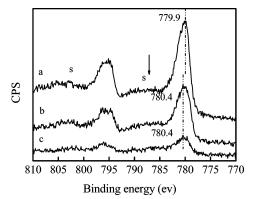


Figure 5. XPS Co 2p spectra of different supported CoO_x: (a) CoO_x/CZI, (b) CoO_x/MZIW, (c) CoO_x/MZI.

TABLE 1: Co Concentration in the Whole Particle and the Surface Phase of Different Catalysts

sample	Co in the bulk (wt %)	Co on the surface (wt %)
CoO _x /MZIW	1.9	4.2
CoO_x/MZI	1.7	1.7
CoO_x/CZI	2.3	7.6

Co. The Co concentrations on the whole particle and the surface of the particle were measured by the analysis of XPS and ICP (induced coupled plasma spectrometer). Table 1 summarizes the analysis results for different samples. The Co concentration on the surface phase (4.2 wt %) was much higher than that (1.9 wt %) in the bulk for CoO_x/MZIW. This result indicated that most of the cobalt oxides were dispersed on the surface of MZ. For CoO_x/MZI, the surface Co concentration (1.7 wt %) was equal to that (1.7 wt %) in bulk. Conversely, most of the CoO_x was loaded on the surface of CZ by impregnation and calcination; the surface Co concentration (7.6 wt %) was much higher than that in the bulk (2.3 wt %). This was attributed to nonporosity, a smaller surface area and insufficient surface hydroxyls of CZ. The rich surface hydroxyl groups of the support played an important role in the loading and dispersion of CoO_x by the preparation method of impregnation, drying, water washing, and calcination. It was found that little cobalt was loaded on the surface of CZ, and the dosages of Co were almost completely washed out with water. These phenomena did not occur when MZ acted as the support. These results suggested that CoO_x was anchored on the surface of MZ by the interaction of [Co(H₂O)₆]²⁺ with the surface hydroxyls of the support. In the preparation method, the Co concentrations on the whole particle increased with increasing of the dosage of Co and reached to the maximum value around 2% at the dosage of Co 5%; then the Co concentration did not change with increasing Co dosage, even at dosages of 10% and 20%. During impregnation and subsequent drying in air, the complex reacted with both the acidic and basic hydroxyls of MZ to form the (H₂O)_rCo^{2+/3+}-O-Zr link. The excess cobalt, which was not connected directly to the surface of MZ, was almost completely washed out with water in the washing step. Thus, monolayerdispersed CoO_x was formed by calcination. In the preparation method of impregnation, drying, and calcination, the Co concentrations on the whole particle always increased with increasing dosages of Co. Therefore, the multilayer CoO_x was supported on the surface of MZ. To confirm the metallic state of the cobalt incorporated in these samples, these samples were characterized by XPS and UV-vis DRS. The cobalt 2p binding energies were similar for the CoOx/MZIW and CoOx/MZI samples (Figure 5). The binding energies of 780.4 and 795.5 eV were for 2p_{3/2} and 2p_{1/2} transitions, respectively. Unfortu-

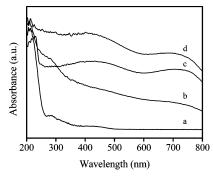


Figure 6. UV-vis diffuse reflectance spectra of different samples: (a) MZ, (b) CoO_x/MZIW, (c) CoO_x/MZI, (d) CoO_x/CZI.

nately, the absolute binding energy of the 2p_{3/2} peak is not always very helpful in identifying the cobalt chemical environment because of the small variation of the binding energy from Co²⁺ to Co³⁺.^{25,26} However, the 2p_{3/2} to 2p_{1/2} separation and satellite structure have elements that are useful in characterizing the cobalt chemical environment. Octahedral Co²⁺ cations, as found in CoO for example, have a very intense, characteristic satellite at ~787 eV; the approximate position is marked as "S" in Figure 5. Moreover, the fwhm (full width at half-maximum) of the 2p_{3/2} main peak is 3.4 eV, which is consistent with the reference data for Co₃O₄.^{27,28} Therefore, there were two types of cobalt (octahedral Co²⁺ and tetrahedral Co³⁺) contributing to this spectral region. Because UV-vis DRS can provide information regarding the oxidation state of cobalt and its chemical environment, the UV-vis spectra of different samples are shown in Figure 6. The pure MZ spectrum only shows a strong absorption edge with a maximum at about 210 nm (curve a), which is attributed to charge-transfer transitions from the 2p level of O to the 4d level of eightfold or sevenfold coordinated Zr⁴⁺.²⁹ Both CoO_x/MZI and CoO_x/CZI exhibited similar UV-vis spectra (curves c and d). Two broad bands at about 425 and 730 nm were observed, which indicated the presence of octahedral Co. According to the literature, 30 the spectra of CoO_x/MZI and CoO_x/CZI are almost identical to the spectra for the bulk Co₃O₄. CoO_x/MZIW exhibited absorption bands in the region of 250-450 nm (curve b), typically assigned to charge-transfer bands,³¹ reflecting the interaction of Co(II) ions with mesoporous ZrO2. The results agreed with that of XRD, indicating the strong interaction between CoO_x and MZ in the sample. Also, CoO_x/MZIW yielded two weak absorption bands at 425 and 750 nm, indicating that Co₃O₄ formed in a very small amount. Comparing the three UV-vis spectra, CoO_x was more highly dispersed on the surface of MZ for CoO_x/ MZIW.

TPR Measurements. The H₂-TPR profiles of different catalysts are shown in Figure 7. According to literature,³² unsupported Co₃O₄ contains one or two peaks close to each other in the temperature range of 200-400 °C. The reduction process is assumed to occur in two steps. One at lower temperature is attributed to the reduction of Co³⁺ to Co²⁺, while the other one at slightly higher temperature is assigned to the reduction of Co²⁺ to Co⁰. In the TPR curves of CoO_x/CZI (curve a), two peaks were observed, one at 300 °C corresponding to the reduction of Co₃O₄ to Co⁰, ³³ and the other one at 705 °C assigned to the reduction of the dispersed Co²⁺ on the surface of CZ.³³ For CoO_x/MZI (curve b), the two reduction peaks at 315 and 370 °C were observed for Co₃O₄. ^{34,35} The peak at 485 °C was possibly attributed to the good dispersion of Co³⁺.36 The CoO_x/MZIW showed two reduction peaks (curve c). One centered at about 565 °C could be assigned to the well-dispersed Co³⁺:³⁶ the other one, centered at about 705 °C, suggested that

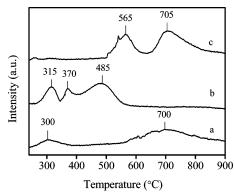


Figure 7. TPR profiles of different samples: (a) CoO_x/CZI, (b) CoO_x/ MZI, (c) CoO_x/MZIW.

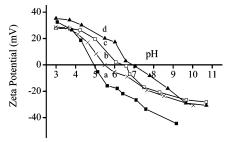


Figure 8. Plot of the zeta potential as a function of pH for different catalyst suspensions in the presence of KNO₃ (10⁻³ M). (a) MZ, (b) CoO_x/MZIW, (c) CoO_x/CZI, (d) CoO_x/MZI.

well-dispersed Co²⁺ was uniformly distributed on the support surface. 32 The results revealed that Co was highly dispersed on the support for CoO_x/MZIW.

Surface Zeta Potential. The activity of solid catalyst in aqueous solution is relative to its surface charge properties. Figure 8 shows the changes of zeta potential with pH of the solution. The upward shifts of isoelectric point from the 4.9 value for MZ used as support were observed after the impregnation of Co. The isoelectric point of CoO_x/MZIW, CoO_x/MZI, and CoO_x/CZI were 5.5, 6.4, and 6.9 respectively. Previous studies³⁷ showed that the negatively charged surface has a strong reactivity toward ozone. This phenomenon is likely due to the electrophilic characteristics of ozone, which has a high affinity for molecular sites with a strong electronic density. CoOx/MZIW possibly had the highest activity.

Ozonation of 2,4-D in the Presence of Supported CoO_x. The catalytic activity of different catalysts was evaluated by the degradation of 2,4-D with ozone at pH 7. As shown in Figure 9, only 28% of TOC was removed at a reaction time of 40 min in the presence of ozone alone. However, the TOC removal increased with the addition of different catalysts. About 50% of TOC was removed at a reaction time of 40 min in CoO_x/ MZI and CoO_x/CZI suspensions with ozone (curves c and d). The two catalysts exhibited almost the same catalytic ozonation activity. CoO_x/MZIW-2 (from Co(NO₃)₂) showed slightly higher activity than them; about 70% of TOC was removed (curve e). The CoO_x/MZIW exhibited the highest activity; the TOC content of the 2,4-D solution was greatly reduced, by 90%, in CoO_x/ MZIW suspension with ozone (curve f). Forty-three percent of TOC was removed in Co²⁺ 1 mg/L (highest concentration detected in $CoO_x/MZIW$ during the reaction at pH = 7) solution with ozone, which is almost the same as that one in the homogeneous catalytic reaction at pH = 4 although the released Co²⁺ concentration is 2 mg/L. The catalytic contribution of homogeneous Co²⁺ was almost same at different pH. The results indicated that the catalytic activity depended predominantly on

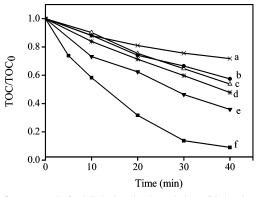


Figure 9. Removal of TOC during the degradation of 2,4-D in aqueous dispersions of various catalysts with ozone: (a) O₃, (b) Co²⁺ 1 mg/L, (c) CoO_x/CZI, (d) CoO_x/MZI, (e) CoO_x/MZIW-2, (f) CoO_x/MZIW. (pH = 7, catalyst: 1.5 g/L, gaseous ozone concentration: 30 mg/L).

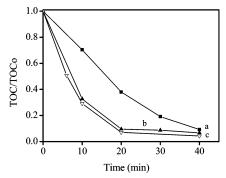


Figure 10. Effect of 2,4-D concentrations on TOC removal in CoO_x/MZIW suspension at pH 7: (a) 80 mg/L, (b) 20 mg/L, (c) 10 mg/L (catalyst: 2 g/L, gaseous ozone: 30 mg/L).

the dispersion of Co on the support. The highest dispersion of Co on MZ was obtained by the preparation processes of impregnation, drying, washing with water, and then calcination, leading to the highest activity. The small surface area and insufficient surface hydroxyls of CZ led to the lower Co dispersion on its surface and the lower activity. The nature of the support, the preparation method, and the Co source played a decisive role in the fabrication of the active phases for Co. The effect of 2,4-D concentrations on the TOC removal was investigated in CoO_x/MZIW suspension with ozone (Figure 10). The TOC of the reaction solution was almost completely removed within 20 min at the 2,4-D concentration of 10 mg/L, while about 94% of TOC was removed within the same reaction time at the 2,4-D concentration of 20 mg/L. Even when the 2,4-D concentration was 80 mg/L, more than 90% TOC disappeared at 40 min. The results demonstrated that CoO_x/ MZIW is a highly efficient catalyst for the mineralization of 2,4-D in the ozonation process. Figure 11 presents the temporal variations of the TOC content of the 2,4-D solution at various concentrations of CoO_x/MZIW. Clearly, the TOC removal rate increased with an increase in the concentration of CoO_x/MZIW, and reached to the maximum at the catalyst concentration of 2 g/L. The results indicated that there were the maximum active centers for the generation of OH at an optimum catalyst concentration. A higher concentration of CoOx/MZIW led to the aggregation of the catalyst, causing a decrease in the catalytic activity. The effect of pH on the TOC removal is shown in Figure 12. In the tested pH range, the activity of the catalyst did not change greatly. The removal rate is also similar both at pH 7 and pH = 10 with ozone alone. The removal rate of TOC at pH 4 was slightly higher than those at other pH levels. The isoelectric point of CoO_x/MZIW was 5.5 pH units. At pH <

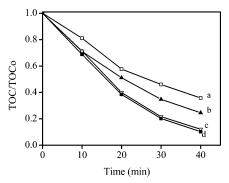


Figure 11. Effect of CoO_x/MZIW concentrations on TOC removal at pH 7: (a) 1 g/L, (b) 1.5 g/L, (c) 2 g/L, (d) 3 g/L.

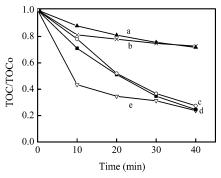


Figure 12. Effect of initial pH on TOC removal of 2,4-D solution in ozone alone: (a) pH = 7, (b) pH = 10, and $CoO_x/MZIW$ catalytic ozonation: (c) pH = 10, (d) pH = 7, (e) pH = 4 (catalyst: 1.5 g/L, gaseous ozone concentration: 30 mg/L).

5.5, the surface of the catalyst was positively charged, while it was negatively charged at pH > 5.5. The more 2,4-D adsorbed onto the catalyst at pH = 4, the more TOC was removed. Alternatively, ozone reacts readily with the negatively charged surface because of its electrophilic characteristics. A higher efficiency of the catalytic ozonation was still obtained in the range of pH 7-10 although 2,4-D hardly had any adsorption on the surface of the catalyst under these conditions. This result indicated that more free OH was generated in the solution by the catalytic transformation of ozone, causing the oxidation of 2,4-D in solution. As shown in Figure S2 (Supporting Information), CoO_x/MZIW did not exhibit any obvious loss of activity when it was further reused for eight cycles. The concentration of Co2+ leached out into solution from CoOx/MZIW decreased with increasing cycle times. Moreover, after eight successive cycles, the concentration of Co in the bulk of the catalyst was 1.8 wt %, slightly less than that one 1.9 wt % in the fresh sample.

Mechanism Discussion. In order to confirm this reaction mechanism, the involved active species were investigated in the catalytic ozonation of 2,4-D. tert-Butanol is a strong radical scavenger that has the reaction rate constant of $6 \times 10^8 \,\mathrm{M}^{-1}\mathrm{s}^{-1}$ with hydroxyl radicals and only $3 \times 10^{-3} \,\mathrm{M}^{-1} \,\mathrm{s}^{-1}$ with ozone. It can terminate radical chain reactions by generating inert intermediates. Thus, tert-butanol was adopted as the indicator for the radical-type reaction. At pH = 7, 2,4-D was hardly adsorbed on the surface of CoO_x/MZIW. The oxidation reaction of 2,4-D occurred mainly in solution. The adsorption ratio of tert-butanol was 5.2% on the surface of $CoO_x/MZIW$ at pH = 7. Therefore, tert-butanol could trap *OH both in solution and on the surface of the catalyst. As shown in Figure S3 (Supporting Information), the addition of tert-butanol markedly reduced the ozonation of 2,4-D at pH 7 in the presence of CoO_x/ MZIW. This result indicated that 'OH was the main active

species in the catalytic ozonation reaction. CoO_x/MZIW accelerated more ozone transformation into OH. Meanwhile, a series of ozone decomposition experiments were carried out in the presence of CoO_x/MZIW. As shown in Figure S4 (Supporting Information), only 50% O₃ was decomposed within 15 min without catalyst (curve a), while the ozone decomposition rate was markedly enhanced in the presence of different catalysts. MnO_x/MZIW exhibited the highest activity (curve d); O₃ was almost completely decomposed within 5 min. MnO_x/CZI showed the least activity (curve b). The ozone decomposition occurred in parallel with the TOC removal in different catalyst suspensions. These results indicated that the supported CoO_x could catalytically decompose O₃ into OH. According to the previous work,³⁸ the mechanism of ozone decomposition on the catalyst consists of mainly redox steps: adsorption of ozone on the catalyst, and desorption of the adsorbed intermediates. Consquently, the faster the catalyst undergoes oxidation and reduction, the faster the rate of the decomposition reaction. The multiple oxidation states of Co in MnO_x/MZIW enhanced the electron transfer, resulting in higher activity. According to the mechanism, the surface Co on the support is active component. Thus, the turnover frequency (TOF) of CoO_x/MZIW, CoO_x/ MZI, and CoO_x/CZI obtained was 0.265, 0.176, and 0.137 s⁻¹, respectively, based on Table 1. Therefore, the dispersion and oxidation state of the supported CoO_x were crucial factors for the high efficiency of the catalytic ozonation.

4. Conclusions

 CoO_x was highly dispersed on the surface of MZ at the monolayer by the impregnation of the cobalt acetate tetrahydrate solution, followed by drying, water washing, and calcination. The supported CoO_x existed mainly as Co_3O_4 phase in $CoO_x/MZIW$, CoO_x/MZI , and CoO_x/CZI . The strong interaction of Co species with MZ was verified in the structure of $CoO_x/MZIW$. The surface hydroxyls and water washing played an important role in the high dispersion of Co on the surface of MZ. The $CoO_x/MZIW$ exhibited the highest activity for the mineralization of 2,4-D with ozone. The multivalence oxidation states and high dispersion of CoO_x in the catalyst enhanced the interfacial electron transfer, leading to high reactivity.

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Supporting Information Available: XRD of CZ and CoO_x/CZ , recycle experiment, effect of *tert*-butanol on 2,4-D removal and ozone decomposition. This material is available free of charge via the Internet at http://pubs.acs.org.

References and Notes

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