The Effects of Mn\textsuperscript{2+} Precursors on the Structure and Ozone Decomposition Activity of Cryptomelane-Type Manganese Oxide (OMS-2) Catalysts

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Supporting Information

ABSTRACT: The effects of Mn\textsuperscript{2+} precursors on the structure and ozone decomposition activity of cryptomelane-type manganese oxide (OMS-2) catalysts were investigated under high-humidity conditions. The OMS-2 catalysts were synthesized using a hydrothermal approach. Characterization of OMS-2 was carried out using X-ray diffraction (XRD), scanning electron microscopy (SEM), \textit{N\textsubscript{2}} physical adsorption, Raman spectroscopy, X-ray absorption fine structure (XAFS), \textit{H\textsubscript{2}} temperature-programmed reduction (\textit{H\textsubscript{2}}-TPR), and inductively coupled plasma (ICP) spectroscopy. The OMS-2-Ac synthesized using MnAc\textsubscript{2} as a Mn\textsuperscript{2+} precursor showed the best catalytic activity for ozone decomposition (~80\%) under RH = 90\% and space velocity of 600000 h\textsuperscript{−1} and is a promising catalyst for purifying waste gases containing ozone under high-humidity conditions. Acetate groups could prevent the aggregation of manganese oxide particles, which may introduce more crystalline defects. On the basis of the characterization results, it is supposed that the greater surface area and higher amount of Mn\textsuperscript{3+} are the main factors that contribute to the excellent performance of OMS-2-Ac. This study can improve our understanding of ozone decomposition on OMS-2 catalysts and serve as a guide in using OMS-2 for ozone removal.

1. INTRODUCTION

As we all know, ozone is a double-edged sword. The ozone layer in the stratosphere can prevent ultraviolet light from reaching the earth directly and protect life on Earth. However, tropospheric ozone is a common pollutant that causes health problems to human beings including neurological disease, increased frequency of respiratory symptoms, and reduced immune system function.\textsuperscript{1} Because of its powerful oxidizing ability, ozone is widely used in purification of wastewater and drinking water, oxidation of organic substances, and food preservation. The resulting tail gas still contains some amount of ozone, and its release into the environment must be avoided. The major indoor source of ground level ozone comes from photocopiers, laser printers, and other modern office equipment. Because most people spend a majority of time inside buildings, it is a significant cause of exposure. According to the U.S. Environmental Protection Agency (U.S. EPA) regulations, the allowable exposure to ozone in terms of 8 h average concentration is 0.075 ppm and the Chinese “indoor air quality standards” (GB/T 18883-2002) requires that the indoor ozone concentration should not exceed 0.07 ppm.\textsuperscript{2} Therefore, the decomposition of ozone is an important area of research from the viewpoint of environmental protection and health.

Because it is safe, economical, and efficient, catalytic decomposition is the most effective method for purifying waste gases containing ozone. Metal oxide catalysts are reported to be among the most active catalysts for ozone decomposition, and considerable efforts have been directed toward the design of ozone decomposition catalysts based on noble metals (e.g., Pd, Au, and Ag)\textsuperscript{3−6} or oxides of transition metals such as Mn\textsuperscript{2+−9}, Fe\textsuperscript{3+−12}, Cu\textsuperscript{2+}, Co\textsuperscript{3+}, and Ni\textsuperscript{2+} Dhandapani and Oyama\textsuperscript{16} pointed out that among the

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common metal oxides (with γ-Al₂O₃ and cordierite as carrier), MnO₂ possessed the highest activity in the catalytic decomposition of ozone. Its better activity may be associated with its changeable valence and variety of morphologies. Cryptomelane-type manganese oxide (OMS-2) is a form of manganese dioxide with a one-dimensional tunnel structure composed of 2 × 2 edge-shared MnO₆ octahedral chains, which are corner-connected to form 4.6 × 4.6 Å tunnels. Manganese is present in OMS-2 as Mn²⁺, Mn³⁺, and Mn⁴⁺ located in octahedral sides, and potassium ions reside in the tunnels along with a small amount of water to stabilize the tunnel structure. The mechanism of ozone catalytic decomposition consists mainly of redox steps, thus the presence of mixed-valent manganese ions probably accelerates the ozone decomposition. The redox pair of Mn(IV)/Mn(III) in OMS-2 seems to facilitate the oxidation thus the presence of mixed-valent manganese ions probably accelerates the ozone decomposition. The redox pair of Mn(IV)/Mn(III) in OMS-2 seems to facilitate the oxidation steps.

2. EXPERIMENTAL SECTION

2.1. Catalyst Synthesis. The OMS-2 materials were prepared via reaction between Mn²⁺ and potassium permanganate with a hydrothermal method. Different Mn²⁺ precursors can have varying effects on the properties of OMS-2, such as crystallinity, surface area, and redox properties and consequently affect the catalytic performance. In this work, OMS-2 catalysts were successfully prepared by using different Mn²⁺ precursors. The as-prepared OMS-2 catalysts were characterized by XRD, SEM, N₂ adsorption–desorption, Raman, XAFS, H₂-TPR, and ICP.

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2.2. Catalyst Characterization. The crystalline structure of the catalysts was determined by a powder X-ray diffractometer (XRD; XPert PRO, PANalytical) using Cu Kα (λ = 0.154 06 nm) radiation at 40 kV and 40 mA. The data of 2θ from 5 to 80° were collected with the step size of 0.06°.

The morphologies of the OMS-2 materials were investigated using a SU8000 field emission scanning electron microscope (FESEM), and the accelerating voltage was 3000 V. The samples for FESEM measurements were prepared by depositing the powder on a conductive tape using N₂ vertical purging.

The surface area and pore structure (volume and size) of the catalysts were determined with a physisorption analyzer (Autosorb-1C-TCD, Quantachrome) by N₂ adsorption–desorption at 77 K. Prior to the N₂ physisorption, all samples were degassed at 90 °C for 12 h to evaporate the physisorbed moisture before measurement. The surface area (S BET) was determined by applying the Brunauer–Emmett–Teller (BET) method to the adsorption isotherm in the partial pressure range of 0.05–0.35. The pore volume (V BH) and the pore diameter (D BH) were determined by the Barrett–Joyner–Halenda (BJH) equation from the desorption isotherm.

Raman spectra of the catalysts were recorded at room temperature on a homemade UV resonance Raman spectrometer (UVRLDPC-DL-03), which consisted of three optional excitation lasers (244, 325, and 352 nm), a three-stage grating spectrograph, and a CCD detector cooled by liquid nitrogen. The instrument was calibrated against the Stokes Raman signal of Teflon at 1378 cm⁻¹. A continuous diode-pumped solid state (DPSS) laser beam (532 nm) was used as the exciting radiation, and the power output was about 48 mW. The diameter of the laser spot on the sample surface was focused at 25 μm. The spectral resolution was 2.0 cm⁻¹. All Raman spectra used in the paper were original and unsmoothed.

The X-ray near edge structure function (XANES) and extended X-ray absorption fine structure (EXAFS) of the Mn−K edge in the OMS-2 catalysts were measured in transmission mode at room temperature on the 1W1B beamline, Beijing Synchrotron Radiation Facility and BL14W1 beamline, Shanghai Synchrotron Radiation Facility. Data processing was performed by the software IFEFFIT 2.11. XANES data were normalized with edge height and then the first-order derivatives were taken to compare the variation of absorption edge energies. EXAFS oscillation χ(κ) was extracted using spline smoothing with a Cook-Sayers criterion, and the filtered k³-weighted χ(κ) was Fourier transformed into R space in the k range of 3–12 Å⁻¹. In the curve-fitting step, the possible backscattering amplitude and phase shift were calculated using FEFF8.4 code.

The H₂ temperature-programmed reduction (H₂-TPR) measurements were carried out using a Micromeritics AutoChem 2920 chemisorption analyzer equipped with a TCD detector. About 50 mg of sample was loaded in a U-shaped quartz reactor and pretreated at 100 °C in a flow of air (50 mL/min) for 1 h. After cooling the reactor to room
temperature, the catalyst was then heated to 1000 °C in a 10 vol% H₂/Ar gas flow of 50 mL/min at a heating rate of 10 °C/min.

The mass ratio K/Mn of the catalysts was analyzed using an inductively coupled plasma instrument (OPTIMA 8300) with a radial view of the plasma. All samples were dissolved using strong acid solution before tested. The calibration solution was prepared using pure materials. The average of three atomic emission lines was used to determine the mass ratio K/Mn in the catalysts.

2.3. Catalyst Activity for Ozone Decomposition. The catalytic performance for ozone decomposition was tested in a fixed bed continuous flow quartz reactor (4 mm i.d.) at a temperature of 30 °C and a space velocity of 600000 h⁻¹. The total gas flow was 1000 mL/min with 800 mL/min N₂ and 200 mL/min O₂. Ozone was generated by low-pressure ultraviolet lamps (185 nm, Beijing Lighting Research Institute) and the inlet ozone concentration was 40 ± 2 ppm. Inlet and outlet ozone concentrations were analyzed by an ozone monitor (model 202, 2B Technologies). The temperature and pressure drop were also recorded at the same time to verify uniformity throughout the experiments. Ozone conversion was calculated on the basis of the following equation:

\[ \text{O}_3 \text{ conversion} = \frac{C_{\text{in}} - C_{\text{out}}}{C_{\text{in}}} \times 100\% \]

3. RESULTS AND DISCUSSION

Figure 1 shows the ozone decomposition activity over the OMS-2 catalysts and amorphous MnO₂ material when relative humidity (RH) was 45%. The activities of OMS-2-Ac and OMS-2-NO₃ were nearly 100% even after 6 h. The activity of OMS-2-Cl decreased with time, to only 85% after 6 h. However, the amorphous MnO₂ material was by far the least active, with a conversion percentage of only 35% after 6 h. The OMS-2 catalysts showed high activity for ozone decomposition compared with nonporous manganese oxides, which may be due to the special structure of the OMS-2 material.

Water vapor has a severe influence on catalytic performance in ozone decomposition. H₂O molecules compete with ozone for adsorption, leading to a decrease in catalytic activity. When the RH was 90%, ozone conversion over these catalysts decreased to greater or lesser extents (Figure 2). Compared with the other catalysts, OMS-2-Ac maintained better ozone conversion performance (∼75%), which indicates that the catalysts prepared from the different precursors were well-crystallized with the cryptomelane structure, characteristic of the octahedral molecular sieve material OMS-2 (JCPDS 29-1020). The variations in peak broadening among the synthesized materials revealed differences in crystal size. The sizes of OMS-2 crystals prepared with different precursors were calculated from the line broadening of XRD peaks using Scherrer’s formula \(d = \frac{0.9λ}{β \cos θ}\). The as-calculated crystal sizes of OMS-2-Ac, OMS-2-NO₃, and OMS-2-Cl were 9.0, 11.4, and 12.1 nm, respectively.

Figure 4 shows the FESEM images of the samples obtained from different precursors. All of the samples displayed fibrous morphologies typical of cryptomelane, regardless of the precursors used. On the basis of measurements of 100 particles for each catalyst material, the average diameters of OMS-2-Ac, OMS-2-NO₃, and OMS-2-Cl were 8.9 ± 2.0, 11.9 ± 2.1, and 14.1 ± 2.4 nm, respectively. This was consistent with the XRD results.
The OMS-2-Ac catalyst showed the smallest particle diameter, consistent with the results calculated from the XRD peaks. Combining the results obtained from XRD and FESEM, it can be speculated that the MnAc2 precursor resulted in the formation of a smaller particle diameter for the OMS-2-Ac catalyst, which is probably related to the presence of acetate groups in the solution. When using MnAc2 as the precursor, the concentration of acetate groups in solution was 0.5 mol/L, much higher than that in the solutions for the other two precursors (0.006 mol/L). Carbonyl groups from the acetate in the solution and the surface OH groups on the manganese oxide materials may have formed bonds, which could cause the isolation of manganese oxide particles from each other by the surrounding acetate groups and prevent further aggregation.42

Smaller particles can introduce more defects and enhance the surface area. The N2 adsorption–desorption isotherms of the three OMS-2 catalysts are shown in Figure 5. All of the samples show similar N2 adsorption/desorption isotherms typical of mesoporous materials. The isotherms show a characteristic type II isotherm pattern for the three catalysts, with microporous filling at low P/P0 and capillary condensation at high P/P0. The surface areas of OMS-2-Ac, OMS-2-NO3, and OMS-2-Cl were 137, 83, and 78 m2/g, respectively. Large surface area can provide more sites for the adsorption of ozone and thus improve the catalytic performance for ozone decomposition on OMS-2-Ac.

Compared with XRD, Raman spectroscopy is more surface-sensitive for the characterization of the catalysts. Figure 6 shows the Raman spectra of the OMS-2 catalysts. All the synthesized OMS-2 materials showed bands at 180, 386, 506, 576, and 630 cm\(^{-1}\). The first three bands correspond to the deformation modes of the metal–oxygen chain of Mn–O–Mn, and the distinct bands at 576 and 630 cm\(^{-1}\) correspond to the stretching mode of the Mn–O lattice.27,43–45 The OMS-2-Ac catalyst showed Raman spectra characteristics similar to both OMS-2-NO3 and OMS-2-Cl but gave much broader and weaker peaks. This is a size-dependent phenomenon commonly observed with nanoparticles and can be explained by the inhomogeneous strain broadening associated with dispersion in particle size and by phonon confinement.46 This observation is consistent with the XRD and SEM measurements, in which the particle diameter of the OMS-2 samples decreases in the sequence of OMS-2-Cl > OMS-2-NO3 > OMS-2-Ac.

X-ray absorption fine structure (XAFS), including X-ray absorption near edge structure (XANES) and extended X-ray absorption fine structure (EXAFS), was used to determine the electronic and structural properties of the catalysts. The EXAFS study of MnO2 is fairly straightforward since it has only one kind of Mn site. The XAFS of the Mn–K edge, including XANES and EXAFS, were measured using Mn foil, MnO, Mn2O3, and MnO2 as reference samples. The full XAFS spectra of all the catalysts are listed in Figure S2. As shown in Figure 7A, the XANES spectra of MnO2 and Mn2O3 were similar, and both the pre-edge peaks and the postedge regions of Mn–K XANES in OMS-2 catalysts were similar to those spectra. Yet, no obvious differences between the catalysts prepared by different precursors were observed. Afterward, to further determine the average valence of Mn species in our catalysts, the first-order derivatives of Mn–K XANES in the series of MnO\(_x\) catalysts were taken and compared with those in the reference samples.47 As shown in Figure 7B, the Mn K-edge absorption energy (E\(_0\)) of the reference materials was determined to be 6548.4 and 6552.8 eV for Mn2O3 and MnO2. The absorption edge energy of Mn species in MnO\(_x\) catalysts was 6551.8 eV, which was slightly less than that in MnO2, most probably due to the presence of a small amount of Mn\(^{3+}\). The exact amount of Mn\(^{3+}\) in OMS-2 catalysts calculated by the Rex2000 analysis package was 14.4%, 3.12%, and 2.12% for OMS-2-Ac, OMS-2-NO3, and OMS-2-Cl, respectively. Linear combination fitting of the XANES spectra of the
OMS-2 catalysts with Mn foil, MnO, Mn$_2$O$_3$, and MnO$_2$ reference materials showed that the percentage of Mn$_2$O$_3$ in the catalysts was 7.2, 1.5, and 1%, respectively. These results indicate that the electronic and local structure of our MnO$_x$ catalysts is similar to that of MnO$_2$, which is in good accordance with the XRD results.

Figure 8A shows the filtered $k^3$-weighted EXAFS oscillations Fourier transformed into R space of the Mn−K edge in the series of OMS-2 catalysts, and Figure 8B shows the corresponding filtered $k^3\chi(k)$ in the $k$ range of 3−12 Å$^{-1}$. It should be noted that the EXAFS of the OMS-2 catalysts was modeled based on the structure of MnO$_2$ and no contribution from Mn$_2$O$_3$ was considered in the fitting of these catalysts. Because the highest proportion of MnO$_2$ in the form of Mn$_2$O$_3$ in our three OMS-2 materials is only 7.2%, this fitting is reasonable.

The single and multiple scattering paths from the tetragonal cryptomelane structure (space group I$4/m$) of α-MnO$_2$ were used as the reference model. The crystal structure of the reference material is listed in Table S1. As also shown in Figure 8A, due to the residual potassium species in the catalysts, the coordination shells of Mn−O and Mn−Mn in our catalysts were different to a certain extent when compared with those in the MnO$_2$ standard samples (e.g., the relative peak intensity of Mn−Mn$_1$ and Mn−Mn$_2$). These potassium ions occupied the interstitial sites formed by the stacking of Mn−O octahedrons. From Figure 8B, we can see that good curve fitting degrees could be obtained between experimental data and theoretical results, and the curve fitted data are presented in Table 2.

Table 2. Curve-Fitting Results of Mn K-Edge EXAFS in OMS-2 Catalysts Prepared Using Different Precursors

| sample            | Mn−K reference | shell | CN$^a$ | R$^b$ (Å) | DW$^c$ (Å) | R factor (%) |
|-------------------|----------------|-------|--------|-----------|------------|-------------|
| MnO$_2$           | α-MnO$_2$      | Mn−O  | 5.9    | 1.91      | 0.063      | 0.3         |
|                   |                | Mn−Mn$_1$ | 4.0    | 2.86      | 0.082      |             |
|                   |                | Mn−Mn$_2$ | 4.0    | 3.44      | 0.032      |             |
| OMS-2-Ac          | α-MnO$_2$      | Mn−O  | 5.6    | 1.92      | 0.06       | 2           |
|                   |                | Mn−Mn$_1$ | 4.0    | 2.87      | 0.057      |             |
|                   |                | Mn−Mn$_2$ | 2.6    | 3.4       | 0.05       |             |
| OMS-2-NO$_3$      | α-MnO$_2$      | Mn−O  | 6.2    | 1.92      | 0.057      | 0.8         |
|                   |                | Mn−Mn$_1$ | 4.2    | 2.86      | 0.06       |             |
|                   |                | Mn−Mn$_2$ | 3.5    | 3.43      | 0.06       |             |
| OMS-2-Cl          | α-MnO$_2$      | Mn−O  | 6.0    | 1.92      | 0.041      | 0.8         |
|                   |                | Mn−Mn$_1$ | 4.2    | 2.86      | 0.058      |             |
|                   |                | Mn−Mn$_2$ | 3.5    | 3.41      | 0.066      |             |

$^a$CN: coordination number. $^b$R: bond distance. $^c$DW: Debye–Waller factor.

After curve fitting, the reference material (α-MnO$_2$) was confirmed to possess Mn−O, Mn−Mn$_1$, and Mn−Mn$_2$ coordination shells at ca. 1.91, 2.86, and 3.44 Å with the coordination numbers being 5.9, 4.0, and 4.0, respectively. Moreover, it is clear that not only the first coordination shell of Mn−O but also the second coordination shells of Mn−Mn$_1$ plus Mn−Mn$_2$ in OMS-2 samples prepared from manganese nitrate and manganese chloride are almost the same as those in the MnO$_2$ reference sample.

However, the OMS-2-Ac catalyst prepared from manganese acetate showed a relatively small coordination number for the Mn−Mn$_2$ coordination shell and had the best ozone decomposition performance, suggesting that this catalyst possessed the most abundant surface defects, which was
beneficial to the adsorption and activation of ozone. The Mn−
Mn₂ species, with longer bond length than that of Mn−Mn₁
species, might be the real active center in our reaction.

Temperature-programmed reduction is an important char-
acterization technique to study the redox properties of metal
oxide catalysts. The H₂-TPR profiles obtained for the OMS-2
catalysts are presented in Figure 9. It could be observed that

![Figure 9. TPR profiles of the OMS-2 catalysts.](image)

OMS-2 prepared from different precursors all showed two
overlapped reduction peaks, which corresponded to a two-step
reduction process. The reduction characteristics of materials
with the cryptomelane structure have been described elsewhere.
The lower temperature reduction process could be assigned to the
MnO₂ to Mn₃O₄ reduction and that at high temperature to the
further reduction to MnO. Nevertheless, the number and intensity of the peaks on the TPR profiles can also depend on the different local environments of such species. In accordance with the literature, the mechanism of ozone decomposition consists mainly of redox steps: adsorption of ozone on the catalysts and desorption of the adsorbed intermediates. Consequently, the faster the catalyst undergoes reduction, the faster will be the rate of the decomposition reaction. As shown in Figure 9, the OMS-2-Ac catalyst, with the best ozone decomposition activity, had the lowest reduction temperature, which is in good accordance with the results obtained by Oyama et al.

By assuming Mn²⁺ as the final oxidation state, the average oxidation state (AOS) was calculated for Mn in the catalysts (according to its hydrogen consumption) and listed in Table 3. A lower AOS implies a higher fraction of Mn³⁺ in the catalyst. The fraction of Mn³⁺ was calculated to be 16%, 3%, and 2% for OMS-2-Ac, OMS-2-NO₃, and OMS-2-Cl, respectively, which was consistent with the values calculated by XANES (Table 3). The K/Mn mass ratio of the OMS-2 catalysts determined by ICP was also given in Table 3. K⁺ ions are situated into the channels of the OMS-2 structure in order to stabilize it by providing charge balance, so a higher K/Mn ratio means a lower manganese oxidation state, which is consistent with the results of XANES and H₂-TPR. DFT calculations have confirmed that the presence of K⁺ located in the tunnels of OMS-2 can improve its lattice oxygen activity, and the higher the concentration of K⁺, the lower the removal energy of lattice oxygen. In this study, the concentration of K⁺ is higher in OMS-2-AC, thus the removal energy of lattice oxygen is lower, which is beneficial for the ozone decomposition.

The mixed valency in OMS-2 catalysts was important for electron transport because the efficiencies of catalysts, especially those serving as catalysts for redox reactions, are usually governed by their ability and tendency to cycle between different valence states of the relevant cationic ions. Ozone decomposition on the surface of OMS-2 could be presented by the following reaction scheme:

\[
\text{O}_3 + [\text{Mn}^{3+}] \rightarrow \text{O}_2 + \text{O}_{\text{ads}}[\text{Mn}^{4+}]
\]

\[
\text{O}_3 + \text{O}_{\text{ads}}[\text{Mn}^{4+}] \rightarrow 2\text{O}_2 + [\text{Mn}^{3+}]
\]

Abundant Mn³⁺ present on the surface of the catalyst could favor the ozone decomposition. A linear correlation between ozone decomposition and Mn³⁺ content (wt %) was found by Liu et al. on AgMn/HZSM-5. Therefore, the higher content of Mn³⁺ also contributed to the high decomposition of ozone on OMS-2-AC.

4. CONCLUSIONS

The effects of different Mn²⁺ precursors on the structure and ozone decomposition activity of OMS-2 catalysts were investigated under high-humidity conditions. The OMS-2-Ac material prepared using MnAc₂ as the Mn²⁺ precursor displayed higher ozone conversion and better hydrophobicity than catalysts prepared using Mn(NO₃)₂ or MnCl₂. Under gas hour space velocity (GHSV) of 60000 h⁻¹ and relative humidity of 90%, the OMS-2-Ac catalyst maintained almost 80% conversion after 6 h, which is promising for the purification of waste gases containing ozone under high-humidity conditions. Compared with the other two Mn²⁺ precursors, acetate groups could prevent the aggregation of manganese oxide and form smaller particles as well as abundant Mn³⁺. The OMS-2-Ac catalyst tended to expose more defects channels of the OMS-2 structure in order to stabilize it by providing charge balance, so a higher K/Mn ratio means a lower manganese oxidation state, which is consistent with the results of XANES and H₂-TPR. DFT calculations have confirmed that the presence of K⁺ located in the tunnels of OMS-2 can improve its lattice oxygen activity, and the higher the concentration of K⁺, the lower the removal energy of lattice oxygen.

**Table 3. ICP, XANES, and H₂-TPR Values of the OMS-2 Catalysts**

| Catalysts | K/Mn (%) | Mn³⁺ (%) | H₂ Consumption (mmol/g) | AOS (%) | Mn³⁺ (%) |
|-----------|----------|----------|--------------------------|---------|----------|
| OMS-2-Ac  | 8.274    | 14.4     | 11.01                    | 3.84    | 16       |
| OMS-2-NO₃| 7.406    | 3.12     | 11.76                    | 3.97    | 3        |
| OMS-2-Cl  | 7.476    | 2.12     | 11.75                    | 3.98    | 2        |

**ASSOCIATED CONTENT**

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XRD patterns of the amorphous MnO₂ material, XAFS spectra of the OMS-2 catalysts, and crystal structure of the reference material (PDF)
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AUTHOR INFORMATION

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