Correlation between kinetics and chemical bonding state of catalyst surface in catalyzed magnesium hydride

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Abstract

We reported on the hydrogen desorption properties, microstructure, kinetics, and chemical bonding state of catalyst surface for composites of MgH2 and 1 mol% Nb2O5 ball-milled for 0.02 h, 0.2 h, 2 h, 20 h under 1 MPa H2 atmosphere, as well as hand-mixed (HM) one. Hydrogen desorption properties were significantly improved by ball-milling with Nb2O5. Then, we estimated by Kissinger Method the activation energy \( E_a \) of hydrogen desorption reaction that decreased with the increase of ball-milling time. Especially, \( E_a \) of the sample ball-milled for 0.2 h was drastically decreased, compared with that of the sample ball-milled for 0.02 h. TEM observations revealed that the distribution of Nb2O5 in MgH2 was gradually improved during ball-milling. On the other hand, we confirmed by XPS that in the sample ball-milled for 0.2 h, Nb2O5-x phase(s) existed at least on the surface. It can be suggested that these deoxidized Nb2O5-x phases eventually decrease \( E_a \) as substantial catalyst rather than Nb2O5 itself.

1. Introduction

Many kinds of hydrogen absorbing materials have been studied in order to store hydrogen, such as sodium alanates [1,2], metal nitrides [3,4], as well as MgH2 [5,6], which is one candidate because of its high hydrogen capacity (7.6 mass%). However, the absorption and desorption reaction of Mg/MgH2 itself is too slow. Recently, scientists found out that the composite of MgH2 and transition metal compounds prepared by ball-milling showed improved kinetics for both hydrogen absorption and desorption [7–11], which can be seen as a good indicator for practical application. Barkhordarian et al. reported that the Nb2O5 had a superior catalytic effect for absorption and desorption, compared to other metal oxide catalysts. The magnesium hydride catalyzed with 0.5 mol% Nb2O5, which was ball-milled for 100 h, finished full desorption within 90 s under vacuum conditions at 300 °C. The desorbed products could also absorb roughly 7 mass% of hydrogen within 60 s under 8.4 bar of hydrogen at 300 °C [10]. Hanada et al. reported that the desorbed composite, MgH2 with 1 mol% Nb2O5 milled for 20 h, was able to absorb ~4.5 mass% of hydrogen under lower pressure than 1.0 MPa within 15 s even at room temperature [11]. However, the mechanism of this significant catalytic effect has not been clarified yet. Hanada et al. observed the ball-milled MgH2–Nb2O5 mixture by the Transmission Electron Microscope (TEM) and found that the additive particles were homogeneously distributed in MgH2 particles within nanometer scale, thus leading to the fast kinetics [12]. Additionally, they performed X-ray absorption spectroscopy for investigating the valence of metal oxide...
additives such as Nb₂O₅ and V₂O₅. The results showed that the metal oxide had been reduced by ball-milling with MgH₂ for 20 h. Therefore, they suggested that the improvement of hydrogen sorption was caused by the catalytic effect of reduced metal oxides [13]. Meanwhile, Porcu et al. confirmed via High Resolution TEM that in the MgH₂–Nb₂O₅ ball-milled composite MgNb₂O₃.67 was present, which may elaborate on the improved kinetics [14]. With respect to the chemical bonding state of niobium, both results indicated that the reduced niobium oxide existed in the sample after milling with MgH₂. However, few works have been done to directly clarify the correlation between the kinetics and the reduced catalyst. The exact mechanism of how the reduced oxides affect the kinetics of reaction should be investigated further in order to develop catalytic effect in hydrogen storage materials, not only in MgH₂. In this work, we have investigated the kinetics of desorption, the morphology and the chemical state on the surface of the catalyst in MgH₂–Nb₂O₅ composites ball-milled for varied time. Especially, we have tried to trace the change of chemical bonding states of catalyst surface by X-ray photoelectron spectroscopy (XPS) under the consideration that the state of catalyst surface is the most important factor affecting the kinetics of desorption.

2. Experimental procedures

MgH₂ powder was purchased from Alfa Aesar with a purity of 98%. Nb₂O₅ powder with the purity of 99.99% was also purchased from the Kojundo chemical Laboratory, with a size of ~1 μm. Ball-milling was performed using a Fritsch P7 device. The samples, mixtures of 300 mg MgH₂ and 1 mol% Nb₂O₅, were ball-milled with 20 steel balls (7 mm in diameter) for 0.02 h, 0.2 h, 2 h and 20 h. The ball-to-powder ratio was 100:1. The hydrogen pressure inside the milling spot was 1 MPa and the rotating speed was 400 rpm during milling. Also, hand-mixed (HM) sample with the same composition was prepared by agate mortar. All the samples were treated without exposure to air, because of the instability of MgH₂. The hydrogen desorption properties were examined by thermal desorption spectroscopy (TDS) with several rates from 1 to 20 °C/min under a highly pure helium flow. X-ray diffraction (XRD) was performed by using JEOL JDX-3500 powder diffractometer with Cu Kα radiation. The samples were observed by 200 kV TEM (JEOL JEM-2010). XPS was carried out using a JPS-90MX photoelectron spectrometer with Mg Kα radiation. The energy scale of the spectrometer is calibrated by setting oxygen (O1s) peak to 532 eV. The spectra were smoothed and subtracted the background according to the Shirley method. Peak separation was performed using Gaussian-Lorentz routines with the ratio of 0.8. The area ratio was fixed to 2/3 due to spin-orbit coupling and the distance 2.7 eV between Nb3d₃/₂ and Nb3d₅/₂ peaks.

3. Results and discussion

Fig. 1 shows the profiles of TDS of hydrogen for the composites with a heating rate of 5 °C/min. It can be seen that as the ball-milling time increased, the peak temperatures of hydrogen...
desorption are decreasing, meaning the catalytic effect was gradually activated and improved during ball-milling. By measuring TDS at different heating rate, the activation energy ($E_a$) of desorption was calculated by Kissinger Method [15], according to the following equation:

$$\ln\left(\frac{\beta}{T_p^2}\right) = -\frac{E_a}{R} + \ln\left(\frac{k_0 R}{E_a}\right),$$

where $\beta$ is the heating rate, $T_p$ indicates the peak temperatures of desorption, $R$ is the gas constant and $k_0$ is the frequency factor. If plotting $\ln\left(\frac{\beta}{T_p^2}\right)$ as the function of the inverse of $T_p$, straight lines can be obtained, namely the Kissinger plot. From the slope of the straight lines $E_a$ for the HM sample and those ball-milled for 0.02 h, 0.2 h, 2 h and 20 h was estimated to be 147 kJ/molH$_2$, 138 kJ/molH$_2$, 82 kJ/molH$_2$, 70 kJ/molH$_2$ and 63 kJ/molH$_2$, respectively. The correlation between $E_a$ and ball-milling time was shown in Fig. 2. It can be seen that $E_a$ decreases with the increase of ball-milling time. Especially between the samples ball-milled for 0.02 h and 0.2 h, a large reduction of $E_a$ (~56 kJ/molH$_2$) has been reached, which is worth discussion. This indicates that some changes were brought by ball-milling and hence improved the catalytic effect of Nb$_2$O$_5$. To understand the mechanism of the catalytic effect, these changes should be investigated.

XRD patterns of the samples were obtained to evaluate the crystalline information, as shown in Fig. 3. The width of the peaks widened and the intensity drastically decreased when the samples were milled for a long time. It indicates that the grain size of both MgH$_2$ and Nb$_2$O$_5$ becomes smaller during ball-milling. This refinement of grains may suggest the decrease of size for both MgH$_2$ and Nb$_2$O$_5$ particles, leading to

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**Fig. 4** – TEM micrographs of HM and ball-milled samples: bright field images of HM (a), 0.02 h (b), 0.2 h (c), 2 h (d), 20 h (e) and the selected area diffraction from area A (f) and B (g).
a better distribution of catalyst in the samples. TEM observation provides evidence for the inference above. In the bright field images shown in Fig. 4a–e (all of them were in the same scale), two kinds of particles could be distinguished by different contrast. The selected area diffraction was applied for both of them (see area A and B in Fig. 4b). In Fig. 4g the spots with streak (obtained from area B) are recognized as Nb2O5, while in Fig. 4f the Debye rings (obtained from area A) are confirmed as Mg and MgO. Therefore, in the bright field images the particles with very dark contrast should be Nb2O5 while the others are Mg related phases. MgH2 was not found because of the fast decomposition under the electron beam, as discussed in other papers [12,14]. When comparing the images, it can be seen that the Nb2O5 particles were gradually attenuated by ball-milling and a homogeneous distribution was reached in the samples milled for long time. In Fig. 4e, e.g., the Nb2O5 particles imbed homogeneously in the Mg related phases. The size of the Nb2O5 particles was estimated to be less than 100 nm. This gradually improved distribution of the catalyst with the refined size should be one factor responsible for the decreased Ea and improved kinetics.

However, we noticed that Ea decreased not gradually but in fact drastically when the sample was milled for 0.2 h. Thus there should be other factors that affect the kinetics more essentially. Under this consideration, XPS measurements were carried out for further investigation. The niobium (Nb3d) spectra were drawn in Fig. 5 after calibration. A decrease of the Nb3d signal with ball-milling can be seen, reasserting the results from elsewhere [16,17]. The decrease could be explained as the milled MgH2 particles covered the surface of Nb2O5, preventing the detector from getting enough signal from inside. It can be seen that the position of the Nb3d peaks detected almost remain the same, even after the sample was milled for a substantial period. However, the obviously widened peaks were obtained in the sample milled for 0.2 h. Further separation of those peaks shown in the inserted graph resulted in a coexistence of 4 peaks. There are two peaks at 210.2 eV and 207.5 eV, which are corresponding to Nb3d5/2 and Nb3d3/2 (210.1 eV and 207.4 eV respective in reference to [18]), and two new peaks appearing to the right. Though the signal is not strong enough to identify the exact phase for the new peaks, the shift to lower energy infers that the chemical state of Nb has changed at least on the surface of the sample. A certain or a complication of Nb2O5-x phase(s) appeared in the sample milled for 0.2 h. The deoxidized Nb2O5 on the surface with the valence of Nb less than +5 may act as a more effective catalyst that decreases Ea and also improve the kinetics of desorption. Here it is also noticed that there was a small shoulder at higher energy side (~211 eV). We considered it as noise because fully oxidized niobium (Nb2O5) should have the Nb3d peak with the highest binding energy.

4. Conclusion

In conclusion, we were able to perform a study on the HM MgH2–Nb2O5 composite as well as those ball-milled for 0.02 h, 0.2 h, 2 h and 20 h under 1 MPa H2 atmosphere. An improved desorption kinetic was reached by ball-milling with Nb2O5. Ea of HM, 0.02 h, 0.2 h, 2 h and 20 h samples was estimated to be 147 kJ/molH2, 138 kJ/molH2, 82 kJ/molH2, 70 kJ/molH2 and 63 kJ/molH2, respectively, thus decreasing in accordance with the improved kinetics. XRD and TEM results showed a decrease of particle size for both MgH2 and Nb2O5 occurred with ball-milling. In the sample milled for 20 h, the Nb2O5 particles were found imbedding homogeneously in Mg related phases, with a size of less than 100 nm. The better distribution and the refined size of the Nb2O5 may be responsible for the decreased Ea and improved kinetics for desorption. The chemical state of Nb was found changed at least on the surface, as evident by XPS results. The peak separation was performed on the XPS spectrum of the sample milled for 0.2 h, indicating that a certain or a complication of deoxidized Nb2O5-x phase(s) appeared, which may play a more important role in decreasing Ea and improving kinetics for desorption.

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Fig. 5 – XPS spectra of HM sample and those ball-milled for 0.02 h, 0.2 h, 2 h, and 20 h. The insert graph shows the Nb3d peak separation of the sample milled for 0.2 h.
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