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A metal-oxide catalyst enhanced the desorption properties in complex metal hydrides

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In this study, LiTi2O4 was synthesized as a possible catalyst for complex metal hydrides. LiTi2O4 was stable in the sample after high-energy ball milling and heat treatment. LiTi2O4 exhibited a catalytic effect among the samples of MgH2, LiAlH4 and LiNH2. The desorption kinetics and the purity of the desorbed hydrogen gas have been improved by doping LiTi2O4. Furthermore, the catalytic mechanism of LiTi2O4 was discussed in accordance with the experimental results.

1. Introduction

Hydrogen is a kind of sustainable energy carrier compared to conventional ones such as fossil fuels. It has advantages such as being clean, is environmentally friendly and forms water as a non-polluting product during practical applications. Accordingly, the reduction of our dependence on fossil fuels and emissions of pollutants and greenhouse gases could be realized by using hydrogen as a primary fuel successfully. This would have a great influence on both changing energy structure and improving global warming. Recently hydrogen energy has been actively investigated in terms of production, storage and application (comprising combustion engines in vehicles and power application). Nowadays, compressed gaseous hydrogen is used in prototype and liquid form in advanced tank systems. However, storing hydrogen in solid-state materials has definite advantages on accessible hydrogen content and energy efficiency compared to complex system that use high-pressure gas or cryogenic liquid. A series of storage materials have been proposed, such as metal-hydrides, alanates and amides. While the hydrogen capacity of complex metal hydrides is reasonable (often in the 3–12 wt% range), relatively high thermal stability, slow kinetics of the pure hydrides, undesirable by-product gases (e.g. ammonia) and irreversibility on cycling are still limitations for on-board applications.

Extensive efforts are currently being made for the improvement of hydrogen storage properties by improving the absorption/desorption kinetics, reducing the operating temperatures and improving the purity of the desorbed hydrogen gas. One of the strategies for the improvement of hydrogen storage properties is the addition of catalysts. Ti compounds have been well known to catalyze the dehydrogenation of complex aluminium hydrides in solution. Wieberg et al. in 1951 observed that Ti catalysed the dehydrogenation of LiAlH4 in a diethyl ether suspension. A breakthrough came when Bogdanovic and Schwickardt certified that doping a few mol% Ti in the complex metal hydride NaAlH4 lowered the dehydrogenation temperature, improved the kinetics and allowed rehydrogenation of the decomposition products. This is an indication that Ti worked effectively for solid-state NaAlH4. Interest in using complex metal hydrides as hydrogen storage materials has dramatically reawakened. Meanwhile, the catalytic effect of Ti compounds was also found in a LiH/LiNH2 system which was reversible and had a high hydrogen capacity of 6.5 wt%. The absorption/desorption kinetics and the purity of the desorbed hydrogen gas had been improved. Among the metal hydrides, MgH2 exhibits a high hydrogen capacity up to 7.6 wt%. However, the absorption and desorption reactions of Mg/MgH2 itself are too slow. Some of the researchers reported that Ti compounds had a catalytic effect on the absorption and desorption. Subsequently, Hanada et al. reported the valence state of the Ti compounds that were doped in MgH2. However, the catalyst state of Ti is not the same as the original after ball-milling and/or heat treatment. In this case, the mechanism of the catalytic effect is difficult to understand clearly due to the instability of Ti compounds.

Normally, the reactions that occur between the solid phases for complex metal hydrides are solid-state reactions. Diffusion plays an important role in the solid-state reactions since they require the coming together of reactive species. The rigid structure makes solids different from liquids and gases. The kinetics of solid-state reactions is therefore greatly dependent on the crystal structure and its defects. Based on these results, one proposal for improving the kinetics of the solid-state reaction is increasing the mobility of ions, such as Li+, Na+ and H− in the complex metal hydrides. This process would progress by doping the catalyst, especially with Ti compounds. Our previous
study suggested that LiTi$_2$O$_4$ was discovered by transmission electron microscopy (TEM) in the Li–N–H sample which was doped with TiCl$_3$ as the catalyst. The catalytic Ti compounds were active at the interface between LiH and LiNH$_2$. Herein, we choose LiTi$_2$O$_4$ as the catalyst for hydrogen storage materials in lithium alanate, magnesium hydride and lithium amides. As a result of their investigation, hydrogen can be desorbed via the following reaction:

$$\text{LiAlH}_4 \leftrightarrow \text{LiH} + \text{Al} + 3/2\text{H}_2$$  \hspace{1cm} (1)  

$$\text{MgH}_2 \leftrightarrow \text{Mg} + \text{H}_2$$  \hspace{1cm} (2)  

$$\text{LiH} + \text{LiNH}_2 \leftrightarrow \text{Li}_2\text{NH} + \text{H}_2$$  \hspace{1cm} (3)  

For the case of LiAlH$_4$, eqn (1) can be recast as a two step reaction as follows:

$$\text{LiAlH}_4 \rightarrow 1/3\text{Li}_3\text{AlH}_6 + 2/3\text{Al} + \text{H}_2$$  \hspace{1cm} (4)  

$$1/3\text{Li}_3\text{AlH}_6 \rightarrow \text{LiH} + 1/3\text{Al} + 1/2\text{H}_2$$  \hspace{1cm} (5)  

Consequently, the catalytic effect of LiTi$_2$O$_4$ will be discovered for samples with the aim of enhancing hydrogen desorption properties in this study.

2. Experimental

2.1. Sample preparation

Powders of LiAlH$_4$ (95%), Li$_2$CO$_3$ (99.977%), TiO$_2$ (99%), Ti$_2$O$_3$ (99.9%), LiH (95%) and LiNH$_2$ (95%) were purchased from Sigma-Aldrich. MgH$_2$ (98%) powder was purchased from Alfa Aesar. The single phase of LiTi$_2$O$_4$ was synthesized by a two-step solid-state reaction. Li$_2$Ti$_2$O$_5$ forms as an intermediate compound. The reaction equations used here are:

$$\text{Li}_2\text{CO}_3 + 2\text{TiO}_2 \rightarrow \text{Li}_2\text{Ti}_2\text{O}_5 + \text{CO}_2$$  \hspace{1cm} (6)  

$$\text{Li}_2\text{Ti}_2\text{O}_5 + \text{Ti}_2\text{O}_3 \rightarrow 2\text{LiTi}_2\text{O}_4$$  \hspace{1cm} (7)  

The synthesis method of LiTi$_2$O$_4$ according to Xu et al. has been used. Considering the volatility of Li, it was hard to get the exact composition of Li. Here an excess molar fraction 0.15 of Li$_2$CO$_3$ is added in eqn (6) for LiTi$_2$O$_4$. An additional factor is the sensitivity of LiTi$_2$O$_4$ to air. This issue was mentioned by several groups, regardless of the preparation method used. The aging reaction is the topotactic oxidation of LiTi$_2$O$_4$. LiTi$_2$O$_4$ reacts with oxygen or H$_2$O in air and is progressively transformed into Li$_{1-x}$Ti$_2$O$_4$ (0 ≤ x ≤ 0.8). The polycrystalline sample is stored in Ar atmosphere in order to avoid the aging reaction with oxygen and moisture.

Typically, in order to ensure a homogeneous mixing between the starting materials and the additive, a ball-milling equipment (Fritsch P7) is used. Samples of LiAlH$_4$ and 0.5 mol% LiTi$_2$O$_4$ were milled for 2 h. The samples, mixtures of MgH$_2$ and 1 mol% LiTi$_2$O$_4$, were milled for 20 h. LiNH$_2$ and LiH powders with 1 : 1.2 molar ratio and 0.5 mol% LiTi$_2$O$_4$ were milled for 2 h. The total weight for each ball-milled sample was 300 mg. In the high energy ball-milling process, powder and 20 steel balls (SUJ-2) with a diameter of 7 mm were brought into a Cr steel pot (SKD-11) and milled at 400 rpm under a hydrogen gas (99.9999% purity) at a pressure of 1 MPa at room temperature. The ball-to-powder ratio was 100 : 1. The milling was interrupted every 1 h for 30 min in order to prevent frictional heat during the milling process.

2.2. Characterization measurements

Structure properties were characterized by X-ray diffraction (XRD) measurement. A sample was firstly put on a glass plate in a glove box. Then the glass plate was covered with a plastic film to protect the sample from contact with air. The plastic film does not have any obvious influence on the obtained XRD patterns. The samples were measured with a diffraction 2θ angle from 10° to 90° under a speed of 0.04° per step. Furthermore, after the milling process, the mixtures were examined by thermogravimetry and differential thermal analysis (TG-DTA) equipment (HITACHI BRUKER TAPS3000S) combined with thermal gas desorption mass spectrometry (TDMS). TG-DTA equipment was installed in another glove box to avoid exposing the samples to air in their measurements.

3. Results and discussion

LiTi$_2$O$_4$ was synthesized by sintering a mixture of Li$_2$Ti$_2$O$_5$ and Ti$_2$O$_3$ at 880 °C under a pressure less than 10$^{-4}$ Pa. Fig. 1 shows the powder X-ray diffraction (XRD) profile. There were no apparent impurities in the product of the single phase LiTi$_2$O$_4$. The color of this single phase was black, which is consistent with previous reports. The stability of LiTi$_2$O$_4$ is reflected by the XRD patterns in Fig. 2. Three samples after high-energy ball milling and dehydrogenation, 0.5 mol% LiTi$_2$O$_4$ + 99.5 mol% LiAlH$_4$ mixture, 1 mol% LiTi$_2$O$_4$ + 99 mol% MgH$_2$ mixture and the composite of LiH and LiNH$_2$ with 0.5 mol% LiTi$_2$O$_4$ were prepared for XRD measurement. The results indicated that the
structure of LiTi$_2$O$_4$ was the same as the single phase of the raw LiTi$_2$O$_4$ after high-energy ball milling and dehydrogenation.

Fig. 3 shows the TG-DTA results of the as received LiAlH$_4$ and the 0.5 mol% LiTi$_2$O$_4$ + 99.5 mol% LiAlH$_4$ mixture. It could be confirmed that during the examined temperature range, the dehydrogenation process showed rather different behavior from these two samples. In the case of the as received sample, two conspicuous peaks appeared around 165–180 °C. The endothermic peak (170 °C) resulted from the melting of LiAlH$_4$. After the endothermic peak, there was an exothermic peak (179.5 °C) that indicated the decomposition of liquid LiAlH$_4$. The reaction started at 160 °C and ended at 240 °C. The total weight loss was 6.4 wt%. In the case of the mixed sample, the hydrogen desorption started around 100 °C, and the whole decomposition of LiAlH$_4$ was complete at around 220 °C. Two endothermic peaks could be seen from the DTA curve. The peak temperature of each peak was 160 °C and 210 °C. Accordingly, some authors have calculated in their reports that the enthalpy of the first dehydrogenation of LiAlH$_4$ with an additive is in agreement with the enthalpy difference between the melting transition and the exothermic decomposition immediately following it for as received LiAlH$_4$.

Consequently, the composite's endothermic-exothermic peak at 170–180 °C in the sample without the additive is in agreement with the absorption of latent heat of melting by LiAlH$_4$ with fast decomposition of Li$_3$AlH$_6$ and desorption of H$_2$, and immediate releasing the latent heat of fusion of Li$_3$AlH$_6$ with desorption of H$_2$. Hence, the first dehydrogenation of LiAlH$_4$ is intrinsically endothermic. Doping LiAlH$_4$ with LiTi$_2$O$_4$ improved its thermal decomposition, bringing this dehydrogenation below the melting point of LiAlH$_4$ and separating it from the complex melting-desorption-solidification event that appears at 170 °C for the undoped sample. This result indicated that LiTi$_2$O$_4$ has some catalytic effect on the lithium alanate.

The TG-DTA results of the milled MgH$_2$ and the 1 mol% LiTi$_2$O$_4$ + 99 mol% MgH$_2$ mixture are shown in Fig. 4. There were two endothermic peaks for the milled MgH$_2$ sample during the measurement. Dehydrogenation started around

![Differential thermal analysis and corresponding weight loss for the dehydrogenation of the as received LiAlH$_4$ and the 0.5 mol% LiTi$_2$O$_4$ + 99.5 mol% LiAlH$_4$ mixture.](image1)

![Differential thermal analysis and corresponding weight loss for the dehydrogenation of the milled MgH$_2$ and the 1 mol% LiTi$_2$O$_4$ + 99 mol% MgH$_2$ mixture.](image2)
260 °C and ended at 380 °C. The peak temperature for each peak was 290 °C and 340 °C. The weight loss of 5.1 wt% was much less than the theoretical value of 7.6 wt%, which indicated an incomplete decomposition of the milled sample under the conditions of heating up to 400 °C at a rate of 5 °C min⁻¹. For the sample with an additive the DTA curve showed that hydrogen desorption started at 220 °C and finished at 330 °C. There was only one main endothermic peak for the mixture sample during the measurements. The peak temperature was at 260 °C. A weight loss of 6.6 wt% was achieved from the TG measurement compared with the theoretical value of 7.0 wt%. The difference between the experimental value and the theoretical value was caused by impurities in the original sample, such as MgO. The much better hydrogen desorption kinetics of the mixture sample was attributed to the catalytic effect of LiTi₂O₄. Recently, Nb₂O₅ doped MgH₂ has been actively studied. LiTi₂O₄ was introduced to this system, the kinetics of MgH₂ has been accelerated at the same level compared with previous work.²⁷

The sluggish sorption kinetics of MgH₂ can be improved dramatically to release 6 wt% of reversible hydrogen at LiTi₂O₄. Recently, Nb₂O₅ doped MgH₂ has been actively studied. The sluggish sorption kinetics of MgH₂ can be improved dramatically to release 6 wt% of reversible hydrogen at LiTi₂O₄. Recently, Nb₂O₅ doped MgH₂ has been actively studied. LiTi₂O₄ was introduced to this system, the kinetics of MgH₂ has been accelerated at the same level compared with previous work.²⁷

The dehydrogenation process of a composite of LiH and LiNH₂ with 0.5 mol% LiTi₂O₄ additive was investigated. Fig. 5 presents the TDMS results. The sample of LiH and LiNH₂ with LiTi₂O₄ showed a sharp H₂ peak. The peak temperature was at 227 °C. No ammonia was detected during dehydrogenation. To the best of our knowledge, addition of LiTi₂O₄ to the system as a catalyst resulted in the lowest desorption temperature observed so far, compared with well known catalysts (BN, Si, TiCl₃).²⁸ However, hydrogen and ammonia were released from the sample without LiTi₂O₄. The broad hydrogen desorption curves and obvious ammonia emission can be seen in the temperature range from 200 °C to 400 °C. The results demonstrated clearly that the purity of the desorbed hydrogen gas was improved after doping with LiTi₂O₄. During heating, the decomposition of the LiNH₂ + LiH mixture was observed to start at 200 °C and complete at 400 °C. By contrast, the addition of LiTi₂O₄ led to full desorption within the temperature range of 150–310 °C. The total weight loss for LiNH₂ + LiH + 0.5 mol% LiTi₂O₄ was 5.7 wt% instead of the 7.5 wt% for LiNH₂ + LiH. Accordingly, a lower dehydrogenation temperature indicated that desorption properties can be improved greatly by adding LiTi₂O₄ as the catalyst.

On the basis of these experimental results, it can be figured out that LiTi₂O₄ worked catalytically for the reactions of lithium alanate, magnesium hydride and lithium amides on hydrogen desorption. As far as the catalytic mechanism of a Ti compound is concerned, some investigations have been performed. For the situations of lithium alanate and magnesium hydride, the explanation of how Ti compounds might work as a catalyst is that it destabilizes the metal–H bonds of the complex metal hydrides, such as the bonding between Mg–H or Al–H. Sandrock et al. suggested that the Ti compound aided the “breaking and re-forming of covalent metal–H bonds”²⁹ Subsequently, starting from 2005, a number of researchers have investigated similar ideas.³⁰⁻³⁵ Additionally, Seong Mu Jo certificated that LiTi₂O₄ nanofibers could absorb/desorb hydrogen at ambient temperature.³⁶ That means LiTi₂O₄ could probably have a similar effect on helping to weaken the metal–H bond, and suggests the increasing lability of the hydride. The catalytic effect is generated by nanocontacts between the sample and LiTi₂O₄ in the mechanical milling process.

For the case of a lithium–nitrogen–hydrogen system, David et al. proposed in their report that the reaction mechanism was based on Li⁺ migration across reactive interfaces between LiH and LiNH₂.³⁷ Additionally, our previous results revealed that the catalytic effect is generated by nanocontacts between the sample and the Ti compound in the interface.³² In the present studies, LiTi₂O₄ has been experimentally proven to have a catalytic effect on the Li–N–H system. The catalytic mechanism might be related to the crystal structure of LiTi₂O₄. LiTi₂O₄ has a spinel-structure, where Li⁺ ions could go through into/out of LiTi₂O₄ without high energy barriers at ambient temperature.³⁸ Because of these characteristics of the crystal structure, LiTi₂O₄ has a high diffusion rate of Li⁺ which is reported to be 10⁻⁶ cm² s⁻¹.³⁹ All of these could help increase Li⁺ ion mobility in the Li–N–H system. Accordingly, the catalytic effect of LiTi₂O₄ probably results from the increase of the Li⁺ ion mobility to improve the reaction kinetics. Meanwhile, the improvement of desorption

![Figure 5](image-url)
kinetics could lead to the consumption of LiNH₂ and therefore the ammonia emission at high temperature could be restricted.

4. Conclusions

In conclusion, we performed a study on the catalytic effect of LiTi₂O₄ in complex metal hydrides, such as magnesium hydride, lithium alanate and lithium amide. The purity and stability of LiTi₂O₄ were confirmed by XRD results. LiTi₂O₄ was stable after high energy ball milling and heat treatment. After adding LiTi₂O₄, the dehydrogenation temperature was decreased effectively among the three kinds of materials. The desorption kinetics and the purity of the desorbed hydrogen gas have been improved by doping with LiTi₂O₄. The catalytic mechanism for complex metal hydrides was also discussed in this study. This is the first time these results have appeared in the literature. It is hoped that they are helpful for hydrogen storage materials as well as fuel cell industries.

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