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Fast hydrogen sorption from MgH$_2$-VO$_2$(B) composite materials

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ABSTRACT

The hydrogen sorption kinetics of MgH$_2$–VO$_2$(B) composites synthesised by mechanical milling have been studied. The microstructural properties of composites were characterized by means of X-ray diffraction (XRD), Raman spectroscopy, Scanning electron microscopy (SEM), Particle size analysis (PSD), while sorption behaviour was followed by differential scanning calorimetry (DSC) and Sievert measurements. Results have shown that although desorption temperature reduction is moderate; there is a substantial improvement in hydrogen sorption kinetics. The complete desorption of pure MgH$_2$ at elevated temperature takes place in more than 30 minutes while the composite fully desorbs in less than 2 minutes even at lower temperatures. It has been shown that the metastable γ–MgH$_2$ phase and the point defects have a decisive role in desorption process only in the first sorption cycle, while the second and the subsequent sorption cycles are affected by microstructural and morphological characteristics of the composite.

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1. INTRODUCTION

Regarding the fact that world-wide use of fossil fuels brings series of ecological problems as well as economy dependence hydrogen energy emerges as viable and sustainable alternative, in a sense of abundance, availability and price. As one of the main problems in hydrogen economy is hydrogen storage and since nowadays known materials for solid-state storage cannot fulfil the targets for automotive applications, fundamental research is needed in this area. Much effort has been devoted to study metal hydrides as materials for hydrogen storage. Metallic magnesium and magnesium based compounds are considered as the most promising candidates. Magnesium is abundant, cheap and a light metal with high gravimetric hydrogen capacity (7.66 wt.%) [1]. Still, high sorption temperature and slow kinetics are main drawbacks of this material.

It has been proven that high energy ball milling of magnesium hydride with various additives (oxide, non-oxide ceramics, amines etc.) improves the hydrogen sorption by reducing particle size and therefore decreasing the desorption temperature at given operating conditions [2–4]. In contrast to hydrides with large particles, the fine powder with its high specific surface area favours hydrogen diffusion and nucleation of MgH$_2$/Mg [5]. Regarding additives, metal oxides are both catalysts and very efficient milling reagent which, together with milling process, can create desirable changes in magnesium hydride structure such as crystal defects, lattice distortion, increase of specific surface area, increase of active sorption sites density and reduction of the effective diffusion length for sorption (by decrease of the average inter-particle

distance) [6]. A wide range of different metal oxides were used to improve the sorption properties of MgH_2 [7–14]. *Borgschulte et al.* proposed that the oxides present at the interfaces may locally destabilize the magnesium hydride phase [8]. Further, *Oelerich et al.* showed that the highest desorption rates are achieved using V_2O_5 and Fe_3O_4 [9]. Metal oxides are dispersed homogeneously due to their brittleness in a way that can improve desorption properties [2]. In addition, oxide additives reduce apparent activation energy and accelerate the reaction in general [7,9,11–13]. The most favourable effect on desorption temperature was reported by *Aguey-Zinsou et al.* [10]. They used Nb_2O_5, (described as a lubricant, dispersing and cracking reagent) and obtained a shift of DSC maximum from 414°C to 264°C, although amount of additive was relatively high (17 wt.%) and the sample was milled for 200 hours.

Ma et al. [11] investigated dependence of apparent activation energy E_a^{des} for hydrogen desorption on milling time in MgH_2-1mol% Nb_2O_5. They showed that E_a^{des} decreases with the increase of milling time: for 0.02h of milling E_a^{des} =147 kJmol^{-1} H_2 while for 20h of milling E_a^{des} =63 kJmol^{-1} H_2. *Croston et al.* [12] examined the influence of TiO_2 on desorption properties of MgH_2 and report the reduction of temperature onset from 360°C to 257°C and the reduction of the apparent activation energy to E_d^{des} =72±3 kJmol^{-1}. Another remarkable reduction of the apparent activation energy for desorption of hydrogen from MgH_2 was reported by *Li et al.* [13]. They used 7mol% of $MnFe_2O_4$ as a dopant and reported a reduction from 255 kJmol^{-1} to 65 kJmol^{-1}. Similar effect was reported for CeO_2 addition. The authors obtained apparent activation energy for desorption of E_d^{des} =60±10 kJmol^{-1} [7].

Having in mind the fact that catalytic activity of transition-metal compounds can be correlated to several factors, starting from high number of structural defects and valence state of

the transition metal to affinity of the transition metal to hydrogen and low stability of the compound, in this paper we have used $VO_2(B)$, characterised by both vacant structure and high valence state to enhance kinetics of MgH_2 [15,16].

2.EXPERIMENTAL DETAILS

Composites were synthesized by high-energy ball milling process, using Spex 8000M mixer mill with different $VO_2(B)$ quantity (5 and 15 wt. %). $VO_2(B)$ was hydrothermally synthesized starting from V_2O_5, (Merck) [17] as a precursor, while MgH_2 was obtained from Alfa Aesar (98% purity). Composites were milled for 2h using 10:1 ball to powder ratio. All samples were kept and handled in the Ar atmosphere in a glove box. Microstructure, phase composition and morphology of the samples were characterized by X-ray Diffraction (XRD, Panalytical X'Celerator θ–θ diffractometer with filtered Cu-K_α radiation), Scanning Electron Microscopy (SEM, JEOL JSM 6460LV) and particle size analysis. To evaluate the average lattice parameters, crystallite size and the relative amount of the different phases before and after cycling, Rietveld analysis of the XRD patterns was carried out using MAUD package [18]. Raman spectra were excited by 514.5nm line of an argon laser and measured at room temperature using a Jobin Yvon model T64000 monochromator, with a CCD detector. Typical resolution was 2cm^{-1}.

Hydrogen sorption properties were examined using Differential Scanning Calorimetry (DSC) and Sievert's measurements. DSC measurements were performed by TA instrument DSC Q10P in hydrogen flowing environment using 10K/min heating rate in temperature range from 80 to 500ºC. The samples were submitted to repeated hydrogen absorption/desorption cycles in a Sievert type apparatus under isothermal conditions in temperature range 330–360ºC and pressure

from 0.4–1 bar for desorption and 8–15 bar for absorption keeping the thermodynamic driving force constant according to the formulation of *Rudman et al.* [19].

3.RESULTS AND DISSCUSION

XRD profiles of both composites obtained by mechanosynthesis with different quantity (5 and 15 wt.%) of $VO_2(B)$ are presented in Figure 1a. The samples are labelled as MV5 and MV15, respectively. The XRD profile both composite after cycling in Sievert's apparatus (entitled as MV15ac) is also presented in Figure 1b. The broad peaks corresponding to β–MgH_2 tetragonal structure are observed in all composites as a result of milling and consequent crystallite size reduction (see Table 1). The XRD profiles show the presence of $VO_2(B)$ as well as reflections attributed to metastable γ-MgH_2 and MgO that were obtained after milling. Formation of γ-MgH_2 and MgO phases is very common after ball-milling [3,6,7,10].

As shown in Table 1, the quantity of MgO in cycled samples is about 46%. This finding suggests that oxidation arises from both, high-temperature reaction with oxygen impurities physisorbed on the samples during handling [6, 7] and from the reactions:

$MgH_2 \rightarrow Mg + H_2$

$2Mg + VO_2 \rightarrow 2MgO + V$
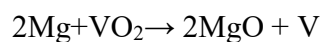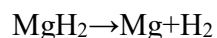
An increase of MgO quantity due to reaction with vanadium oxides has been already reported by several authors [20,21]. The interaction of V with H_2 may lead to formation of an interstitial solid solution, of $VH_{\sim 0.8}$, or of VH_2 depending on the operating temperature and pressures At the conditions of our H-sorption cycles, an interstitial solid solution of H in bcc V is expected according to V-H thermodynamics [5]. However, cooling down to room temperature under H_2 results in the successive formation of $VH_{\sim 0.8}$ and VH_2, when the temperature decreases below the dissociation temperature of these hydrides at the actual H_2 pressure.

In fact, XRD patterns after cycling confirmed the formation of VH_2 phase in both samples. One can notice from Figure 1b that MgH_2, MgO, and VH_2 phases (with some Mg traces in MV5 sample) are present. Some of the Bragg reflections of VH_2 and MgO overlap (see Figure 1b) as the two cubic phases have similar lattice parameters. The intensity of the broad reflection centred at 2θ ~36.8 ° is too high, compared to the MgO (200) reflection centred at 2θ ~42.7 °, to be assigned exclusively to MgO (111) and it originates also from the most intense (111) reflection of VH_2. The lattice parameter of VH_2 was determined as 4.205(5) Å. Accurate Rietveld refinement permitted to determine the contribution of the two phases to the XRD patterns and to derive the relative phase fractions, which are reported in Table 1.

The XRD profile of MV15ac indicates that after sorption cycling there is a conversion of metastable γ-MgH_2 to stable β–MgH_2 phase, in accordance with literature data [3, 6]. It is well known that due to the presence of γ-MgH_2 in the first desorption cycle there is an appearance of a low temperature DSC peak [3]. In tested samples on the other hand, XRD pattern after cycling does not reveal presence of this metastable phase, so one has to disregard γ-MgH_2 phase in practical applications.

Further, since the lattice parameters and c/a ratio of commercial MgH_2 [4] and of composites before and after cycling are very similar [4], no incorporation of V-atom into MgH_2 lattice is expected. Quite interesting is the fact that in the case of MV5 sample, the most of the $VO_2(B)$ is transformed into VH_2. On the other hand in MV15 sample, $VO_2(B)$ has been partially transformed to VH_2 (see Table 1). Nevertheless, the presence of $VO_2(B)$ after cycling is not detected by XRD analysis, meaning that the quantity of this phase is low and/or the crystallite size is very small.

To gain further information on the V-containing phases after cycling, we employ Raman spectroscopy, which can provide details of short range ordering since it is sensitive to the degree of crystallinity [22].

Raman spectra of hydrothermally synthesized $VO_2(B)$, commercial MgH_2 [23] and MV15ac are shown in Figure 2. The Raman phonon vibrations of $VO_2(B)$ obtained in this study are quite similar to those reported by *Ni et al.* [24] (see Table 2). However, the spectrum in Figure 2 shows only five of eighteen possible Raman bands for low temperature $VO_2(B)$ polymorph. Full spectrum, on the other hand, can be obtained for thin films or for a single crystal [25–28]. The Raman shift of VO_2 bands at 160 cm^{-1} and 272 cm^{-1} can be assigned to V–O–V bending modes and external mode (bending/wagging), respectively [24]. The bands at 523 cm^{-1} can be attributed to V–O–V stretching mode while the band at 705 cm^{-1} is due to coordination of vanadium atoms with three oxygen atoms. The band at 1015 cm^{-1} is attributed to V=O stretching of distorted octahedral and distorted square-pyramids [24,29,30].

Regarding composite material MV15ac, one can notice that the whole spectrum exhibits broader features i.e. the bands are wider and the intensity is lower. The sharp band at 311 cm^{-1} related to MgH_2 Raman active-phonons B_{1g} [23,31] disappears after ball milling and cycling treatment, while a band at 374 cm^{-1} appears. Several authors reported the VO_2 Raman scattering mode around 390 cm^{-1}. If one takes into account that mechanical milling causes a change in crystal lattice parameters, and therefore a slight shift in position and significant broadening of bands [31–33], the band at 374 cm^{-1} can be assigned to A_g Raman active-phonon of VO_2.

Due to milling and cycling, the strong and sharp E_g Raman active-phonon at 953 cm^{-1} of MgH_2 sample is shifted toward lower values (922cm^{-1}, see composite sample). This band is wide and with low intensity. It is interesting to note that the band at 1270 cm^{-1}, which exists in

commercial MgH$_2$ [23, 34], completely disappears after cycling and milling. Since bands at higher frequencies are related to H-H interactions, one can conclude that this interaction is rather weak after cycling. On the other hand, the low-frequency mode at 120 cm^{-1} [34, 35] related to Mg vibration still exists in the composite sample. Further, the Raman band that corresponds to V–O–V stretching mode (see VO$_2$ signal, 523 cm^{-1}) is shifted to 542 cm^{-1} in composite material, due to sample milling and interaction with hydrogen during cycling. The VO$_2$ (B) Raman active mode at 271 cm^{-1} is shifted to 254 cm^{-1}.

Based on the above arguments, Raman analysis suggests that VO$_2$(B) is still present in sample MV15 after cycling, i.e. that only partial reduction of VO$_2$(B) occurs. In sample MV5, Rietveld analysis implies that VO$_2$ (B) reduction may be more relevant, but quantitative analysis is difficult due to the low fraction of V-containing phases in this sample.

The composites with different VO$_2$(B) amount exhibit a similar morphology, as demonstrated by the SEM micrographs in Figure 3. In both samples, very small particles (less than 2μm in size) as well as larger agglomerates (more than 50μm in size) are observed, typical for ball-milled powders [5,36]. Particle size distribution (PSD) (see insets in Figure 3) demonstrates a polymodal distribution for both composites in the range of 0.6-100μm. 30% of particles have a size smaller than 10μm, while 70% of particles have size ranging from 10 to 100μm in sample MV5. Regarding sample MV15, 27% of particles have size ranging from 0.6-8μm, 70% of particles from 8 to 80μm and 3% of particles between 80-100μm. The changed morphology and reduced particle size have positive effect on sorption properties, as it will be shown later.

Thermal analysis performed by DSC is shown in Figure 4. The endothermic signals of composites, associated with hydrogen desorption, clearly develop in a lower temperature range

with respect to commercial MgH_2. DSC trace of MV5 exhibits two broad peaks at ~395°C and ~420°C, while MV15 is characterized by a maximum at ~400°C with strong asymmetry towards higher temperatures. These features suggest that hydrogen desorption occurs as two steps process. One can argue that the bimodal particle size distribution is one of the reasons for such DSC behaviour. On the other hand, as shown by XRD, the presence of γ-MgH_2 in composites after milling cannot be excluded as a possible reason of two DSC maxima. Similar temperatures for hydrogen desorption during DSC scans are reported in the literature for different MgH_2-based composites [36,37]. The area under the endothermic desorption peak in the DSC traces decreases from $1380 Jg^{-1}$ for commercial MgH_2 to about 1200 Jg^{-1} for the composites. Such reduction reflects a decrease of the overall gravimetric hydrogen storage capacities of the composites, due to a presence of vanadium based compounds and formation of MgO.

The Sievert's apparatus was used to record hydrogen sorption kinetics under isothermal conditions. In the commercial MgH_2, at temperature of 380°C and pressure of 1bar, the hydrogen desorption capacity saturates at 6.9 wt.% in about 30 min (Figure 5a). At lower temperature (350°C/0.7bar), hydrogen desorption process is significantly slower, reaching only 2.9 wt.% in 5000 min (Figure 5a).

As it can be seen in Figure 5a, composite MV5 shows significant acceleration of hydrogen desorption: at 350°C/1bar capacity of 4.9 wt.% is reached after ~120 seconds, while composite MV15 reaches the capacity of 4.3wt.% after 85 seconds at 350°C/0.7bar. Similar improvement was reported by *Grigorova et al.* [38] for composites with vanadium in valence state 5 (capacity of 5.4% after 25 minutes at 350°C) and *Khrussanova et al.* [39] from 90%Mg+10%V_2O_5 (~4.3% after 30 minutes and the activation energy of 89.7 $kJmol^{-1}$). It is worth noticing that all these observations are valid for the first desorption cycle.

Desorption curves in Figure 5a and 5b show the difference between the first and all subsequent desorption cycles. One can argue that the second and subsequent cycles are affected by chemical composition (type of matrix and catalyst), microstructural and morphological characteristics of composite (particle size of matrix and the catalyst, distribution of the catalyst). Induced vacancies and introduced defects in crystal lattice affect only the first desorption cycle [40]. After subsequent sorption, the material relaxes, so the vacancies and defects disappear and their influence on desorption is negligible. Therefore the activation process has been followed in details. Figure 6 shows sorption cycling of MV15 composite at 320°C before ''activation''[2] and after activation at 380°C. Comparing the storage capacities, one can notice that the capacity without activation process is lower (4.3 wt.%, 4.8 wt.%, 5.0 wt.% in 1st, 2nd and 3rd desorption, respectively) and gradually increases with cycling. After activation, one can see that the hydrogen desorption capacity attains a higher value of 5.4 wt.%, which remains constant during subsequent cycles. Korablov et al. spotted similar behaviour after cycling of MgH_2-V_2O_5 system and correlate it to the *in situ* formation of VH_2/V in the sample [20]. In fact, in the system MgH_2-VO_2 ,Raman and XRD analysis shows formation of VH_2 upon cycling and therefore the existence of VO_2/VH_2 (V^{4+}/V^{2+}) system is probably responsible for strongly improved (de)sorption kinetics. Regarding the mechanism of kinetic reaction one can argue that the existence of multivalent V acts as an intermediate between Mg^{2+} and H^-. Indeed, based on XPS analysis Cui et al. [41] proposed similar mechanism for addition of Ti based compounds to MgH_2. They have found that the interfaces between high valence and low valence Ti compounds are actually responsible for improved kinetics by favoring electron transfer between Mg^{2+} and H^- at the interface and proposed following three-step mechanism: 1) at the hydride/metal interface, H^-

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[2] Activation process: uptake and release of hydrogen at elevated temperature

donates an electron to high valence V, which transforms into low valence V; 2) e⁻ transfer weakens Mg–H bonds and dissociated H is formed, leading to dehydrogenation; 3) finally, two H atoms recombine to form one H_2 molecule, while the metallic Mg phase nucleates and grows.

Recently Paskaš Mamula et al [42] showed using DFT calculations that the substitution of Mg with 3d transition metals results in weaker Mg-H bonds and shorter and stronger TM-H bonds. Further, they have revealed that Bader charge analysis [42] (number, distribution and charge of bonding critical points and attraction basin integrated charges) goes in favor of more directional bonding with larger share of covalence. One must have in mind that the effect is quite localized and constrained to several coordination shells. They have shown that V-H bonds are slightly shorter (1.845 Å) compared to 1.959 Å in pure MgH_2 and are almost the same as in VH_2 (1.85 Å [43]), despite the lower coordination number. The stability of MgH_2-V system is lowered as a direct consequence of dopant concentration (6.25 at%), meaning that localized changes in host lattice caused by V doping are compensated by its concentration.

Hydrogen sorption kinetic curves were fitted according to the Johnson-Mehl-Avrami (JMA) nucleation and growth model: $f(t) = 1 - \exp\left[-\left(kt\right)^n\right]$ where f is the transformed fraction as a function of time t, k is the rate constant and n is Avrami exponent (see Figure 7). The kinetic parameters obtained from the best fit are listed in Table 3. The mechanism of phase transformation was determined using the n value and the apparent activation energies [44, 45]. It can be noticed that n values fluctuate around 1.5 for MV5 and 1.9 for MV15 for composites indicating that there is change in the mechanism of desorption regarding dimensionality of nuclei and the nucleation rate [35]. As it can be seen from Table 4 Avrami exponent can adopt different values depending on the parameters. Avrami exponent is given by $n = a + b * c$ where a is determined by the nucleation mechanism and $a = 0$ for an instant nucleation, $0 < a < 1$ for

decreasing nucleation rate, $a = 1$ for constant nucleation rate and $a>1$ for increasing nucleation rate. On the other hand, b is related to the growth dimensionality ($b = 1$, 2 and 3) and c indicates diffusion-controlled ($c = 0.5$) or interface-controlled ($c = 1$) growth [44, 45]. Therefore, it is difficult to identify the exact mechanism of desorption, but it is obvious that the probability of 2D growth increases with VO_2 (B) addition (see Table 3).

Significant decrease of the apparent activation energies with the addition of catalyst confirms the effectiveness of VO_2(B), which probably acts as nucleation site for the growth of the transformed phase. The obtained E_A^{des} (using JMA), together with literature data, is listed in Table 5. It can be noticed that the apparent activation energies vary depending on the nature of the catalyst but also on the quantity of added phase. Comparing to the apparent activation energy for desorption of pure MgH_2 (161 kJmol^{-1}) obtained by *Fernandez et al*. [46], it can be noticed that E_A^{des} significantly decreases with the addition of VO_2(B). Further, our study confirms the findings of *Barkhordarian et al.* [47], who demonstrated that the apparent activation energy varies with the quantity of added phase.

The results obtained by *Kurko et al*. suggest that lower hydrogen coverage of the surface makes diffusion toward the surface easier [48]. *Borgschulte et al.* suggested that due to formation of vacancies the rate-limiting step is not the dissociation by the active sites, but the number as well as the transport to active sites, i.e. the diffusion on/in oxides [49]. On the other hand, DFT calculation performed by *Du et al.* demonstrates that the diffusion of H-vacancy from an in-plane site to a bridge site on the surface has the smallest activation barrier of 0.15 eV and should be fast at room temperature [50]. The calculated activation barriers for H-vacancy diffusion from the surface into sublayers are all less than 0.70 eV, which is much lower than the activation energy for hydrogen desorption at the MgH_2 (110) surface (1.78-2.80 eV/H$_2$) [50]. This suggests

that surface desorption is the rate determining step rather than vacancy diffusion, so finding an effective catalyst is very important for improving the overall dehydrogenation performances of hydrides. Obviously, obtained apparent activation energy for desorption of about 60kJmol^{-1} (0.63 eV) suggests that $VO_2(B)$ can be used as an excellent catalysts for sorption processes even though the catalytic reaction goes through formation of multivalent system VO_2/VH_2.

4. CONCLUSIONS

Composites MgH_2–$VO_2(B)$ were synthesised by mechanical milling and their hydrogen sorption properties were studied. The broadening of typical peaks corresponding to β–MgH_2 tetragonal structure is observed in all composites as a result of milling and consequent crystallite size reduction. The XRD profiles show the presence of $VO_2(B)$, as well as reflections attributed to metastable γ-MgH_2 and MgO. Upon cycling, the metastable phase γ-MgH_2 converts into a stable β–MgH_2 phase, and the crystalline VH_2 phase appears. Raman spectra confirm the presence of $VO_2(B)$ phase after cycling in the sample with higher quantity of $VO_2(B)$, i.e. MV15ac. XRD analysis indicates that part of $VO_2(B)$ is reduced to V metal at high temperature, with concomitant oxidation of Mg into MgO. Particle size distribution demonstrates polymodal distribution for both composites. There is a shift of desorption temperature toward lower values (about 60°C) and a decrease of gravimetric storage capacity, mainly due to presence of MgO. On the other hand, hydrogen sorption kinetics is substantially improved (from 2.9 wt.% in 5000 min for commercial MgH_2 to 4.9 wt.% in 120 seconds at 350°C/1bar for MV5 sample) due to presence of multivalent V i.e.VO_2/VH_2 system. A considerable difference between first and subsequent sorption cycles is caused by structure relaxation, structure defects disappearance and the presence of VO_2/VH_2. The defect structure and number of vacancies seem to play a more

important role than valence state in the first desorption cycle, while the second and subsequent cycles are more likely improved due to presence of VO_2/VH_2.

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