Book Chapter

Development of Metastable Catalysts to Improve the Hydrogen Sorption Characteristics of Nanostructured Magnesium Hydrides for Hydrogen Storage

Gracia Shokano^{1*}, Zahir Dehouche^{1*}, George Fern¹, Basile Galey² and Georgeta Postole²

¹College of Engineering, Design and Physical Sciences, Brunel University London, UK

²Univ Lyon, Université Claude Bernard Lyon 1, CNRS, IRCELYON, France

***Corresponding Author:** Gracia Shokano, College of Engineering, Design and Physical Sciences, Brunel University London, Uxbridge UB8 3PH, UK

Zahir Dehouche, College of Engineering, Design and Physical Sciences, Brunel University London, Uxbridge UB8 3PH, UK

Published **December 28, 2020**

This Book Chapter is a republication of an article published by Gracia Shokano, et al. at Catalysts in July 2020. (Shokano, G.; Dehouche, Z.; Galey, B.; Postole, G. Development of a Novel Method for the Fabrication of Nanostructured $Zr_{(x)}Ni_{(y)}$ Catalyst to Enhance the Desorption Properties of MgH_2 . Catalysts 2020, 10, 849.)

How to cite this book chapter: Gracia Shokano, Zahir Dehouche, George Fern, Basile Galey, Georgeta Postole. Development of Metastable Catalysts to Improve the Hydrogen Sorption Characteristics of Nanostructured Magnesium Hydrides for Hydrogen Storage. In: VSR Rajasekhar Pullabhotla, editor.

Prime Archives in Chemistry: 2nd Edition. Hyderabad, India: Vide Leaf. 2020.

© The Author(s) 2020. This article is distributed under the terms of the Creative Commons Attribution 4.0 International License(<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Author Contributions: Conceptualization, G.S. and Z.D.; methodology, G.S., Z.D. and B.G.; validation, Z.D., B.G. and G.P.; writing—original draft preparation, G.S.; supervision, Z.D. and G.F. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Conflicts of Interest: The authors declare no conflict of interest.

Abstract

The present work involves the development of a novel method for the fabrication of zirconium nickel ($Zr_{(x)}Ni_{(y)}$) alloy used as a nanocatalyst to improve the hydrogen storage properties of the Mg/MgH₂ system. The catalyst was fabricated through the high-pressure reactor and activated under hydrogen prior to being mechanically milled with the MgH₂ for 5 h under argon. The microstructure characterisation of the samples was determined via SEM-EDX (scanning electron microscope analysis–energy dispersive X-ray spectroscopy), XRD (X-ray diffraction) and FE-HRTEM (field emission high resolution transmission electron microscopy), and the desorption characteristic of the nanocomposite (10 wt.% $Zr_{(x)}Ni_{(y)}$ –MgH₂) was determined via TPD (temperature-programmed desorption). The nanostructured MgH₂ powder milled with 10 wt.% of the activated $Zr_{(x)}Ni_{(y)}$ based nanocatalyst resulted in a faster hydrogen release—5.9 H₂-wt.% at onset temperature 210 °C/peak temperature 232 °C. The observed significant improvement in the hydrogen desorption properties was likely to be the result of the impact of the highly dispersed catalyst on the surface of the Mg/MgH₂ system, the

reduction in particle size during the ball milling process and/or the formation of $\text{Mg}_{0.996}\text{Zr}_{0.004}$ phase during the milling process.

Keywords

Zirconium Nickel Nanocatalyst; Magnesium Hydride; Hydrogen Storage

Introduction

The existing problem of environmental pollution leading to global warming and the rapid growth of industrial sectors requiring a high demand for fossil fuels such as coal, oil and natural gas is a global concern. Therefore, as hydrogen plays a significant role in the field of secondary energy production, and due to its potential to be stored securely for transportation applications, hydrogen storage methods were implemented early on by a number of internationally renowned researchers [1,2]. Hydrogen can be stored in three different ways: in a liquid state, in a gas state and in a solid state. The conventional method of storing hydrogen in a compressed gas or in a liquid phase comes with several disadvantages for vehicular applications [3]. Therefore, storing hydrogen in a solid state is often the convenient approach. Magnesium/magnesium hydride (Mg/MgH_2) is one of the metal hydride systems that serves as a solid state hydrogen storage system, captivating the attention of researchers due to its high and reversible volumetric, $110 \text{ g L}^{-1} \text{ H}_2$, and gravimetric, 7.6 wt.% H_2 , capacities [4]. Furthermore, cost-wise, magnesium is affordable as well as abundantly available [5]. Despite these advantages, the practical applications of the Mg/MgH_2 system come with several drawbacks which limit its utilisation in automobiles [6]. (Refer to Table 1 for the U.S. Department of Energy (DOE) target requirements for on-board hydrogen storage systems 2020, 2025 and ultimate in comparison to the Mg/MgH_2 system).

Table 1: U.S. Department of Energy (DOE) target requirements for on-board hydrogen storage systems versus Mg/MgH₂ storage properties, adapted from [1].

	MgH ₂	2020	2025	Ultimate
Gravimetric capacity (H ₂ -wt.%)	7.6	4.5	5.5	6.5
Volumetric capacity kg H ₂ L ⁻¹	1.4	0.03	0.04	0.05
Operating temperature (max/min) °C	>250	-40/85	-40/85	-40/85
Storage system cycle life (cycles)	1500	1500	1500	1500
Charging rate-filling time for 5 kg of H ₂ (min)	-	3–5	3–5	3–5
Storage system cost (\$/kg H ₂)	Meet target	333	300	266

MgH₂ has an ionic phase with an Mg–H bond charge distribution of Mg 1.91+ H 0.26– [7]. The ionic structure contributes to its high thermodynamic stability of $\Delta H = \pm 75 \text{ kJ mol}^{-1} \text{ H}_2$ and entropy value of $130 \text{ J K}^{-1} \text{ mol}^{-1}$. Therefore, the high stability results in an operating temperature above 350 °C [8]. Moreover, the nanostructured Mg/MgH₂ suffers from poor hydrogenation and dehydrogenation kinetics at elevated temperatures [9]. There are numerous factors that contribute to the reduction in the rate of hydrogenation, which prevents the hydrogen from penetrating into the molecules. For example, the formation of magnesium oxide (MgO) and/or magnesium hydroxide (Mg(OH)₂) on the surface of magnesium may take place when exposing magnesium to air [10]. The oxide layer on the surface of the magnesium particles can be cracked or perforated during the activation of the metal or during the ball milling preparation [11]. Activation consists of cycling and cooling the material in a vacuum atmosphere, and, as a result, the metal surface is exposed to hydrogen [10], [12]. The drawback of Mg/MgH₂ is still under investigation by researchers through different approaches, such as the reduction of the particles' sizes during the ball milling process and the catalyst doping of metallic particles with different fractions of catalytic agents [10]. Nickel (Ni) and zirconium (Zr) were reported to improve the dissociation of hydrogen at the magnesium surface [11]. The metal oxides (ZrO₂, TiO₂) and the metastable phases Zr_(x)Ni_(y) have shown remarkable progress in past studies in enhancing the hydrogen storage properties of MgH₂ [12–16]. Pighin et al. [13] reported the study of 10 wt.% Zr₈Ni₂₁ alloy, where the hydrogen

storage capacity of 5.9 wt.% at 250 °C was obtained. The catalytic effects of various $\text{Zr}_{(x)}\text{Ni}_{(y)}$ alloys used as additives to the Mg/MgH_2 system have also been reported by Dehouche et al. [14], where the best desorption rate was observed with 0.2 mol% $\text{Zr}_9\text{Ni}_{11}/\text{MgH}_2$ at 250 °C (5.88 H_2 -wt.%). The preparation of zirconium nickel alloys, such as the ZrNi_5 , ZrNi_3 , $\text{Zr}_8\text{Ni}_{21}$, ZrNi_2 , $\text{Zr}_7\text{Ni}_{10}$, $\text{Zr}_9\text{Ni}_{11}$, ZrNi and Zr_2Ni alloys, have been reported in several studies by arc melting under a flow of argon gas [13–20]. Arc melting is a long process for catalyst fabrication, requiring the melting of the material up to five times in a repeated cycle to ensure the uniform chemical composition of the resulting alloy, crushing the resulted material (spherical balls) into small pieces and milling the produced alloy powder under argon atmosphere for up to 100 h prior to being milled with magnesium [15]. The sol-gel method has been proposed by Parayil et al. [21] in a past study for the synthesis of TiO_2 photocatalytic hydrogen production. The sol-gel method involves the hydrolysis reaction and the gel formation via condensation. The drying process is the most critical step of the sol-gel method—xerogel drying is a simple method that takes place under ambient conditions. The aerogel drying method, on the other hand, is a time-consuming procedure involving multiple stages, such as the washing of the gel—known as supercritical drying of the wet gel—that may take a few weeks. In addition, drying by aerogel results in the formation of materials with low crystallinity, even when the materials are calcinated at a high temperature of 500 °C [21].

The present work proposes a faster alternative method for the fabrication of a zirconium nickel nanostructured catalyst via a modified sol-gel process conducted in a compact high-pressure hydrogen reactor to enhance the hydrogenation and dehydrogenation kinetics of the Mg/MgH_2 system by producing nanostructured catalyst particles.

Results

Characterisation of $\text{Zr}_{(x)}\text{Ni}_{(y)}$ Catalyst

The zirconium nickel nanostructured catalyst was synthesised via a modified sol-gel method in a compact high-pressure hydrogen reactor and activated prior to being used as an additive

to MgH_2 . The catalyst preparation method used in the present work is further detailed in Section 4. To fully characterize the catalyst after synthesis and activation, XRD, SEM-EDX and FE-HRTEM analytical methods were used. The results of the SEM-EDX are presented in Section 2.4. Firstly, XRD was carried out to identify and quantify the crystalline phases present in the synthesised $\text{Zr}_{(x)}\text{Ni}_{(y)}$ based catalyst powder prepared by the high-pressure reactor (Figure 1). It is visible from Figure 1 that the Ni phase was predominant after the 5-h heat treatment at 350 °C. The precursor Ni powder is indicated by the sharp Bragg peaks at around 2θ of 44.5°, 51.8° and 76.2° (PDF file # 01-070-1849). Additional broad peaks were identified at around 2θ of 30° and 60.5°, corresponding to the development of Zr_2O (PDF file #01-071-6425).

Figure 1: The XRD patterns of $\text{Zr}_{(x)}\text{Ni}_{(y)}$ based nanocatalyst.

The crystallite size (d), was calculated using Scherrer's equation. Refer to Equation (1).

$$D = \frac{k \cdot \lambda}{FWHM \cdot \cos \theta} \quad (1)$$

where D is the crystallite size (nm), k is the Scherrer's constant (0.9), λ is the wavelength of the X-ray source (0.15406 nm), $FWHM$ is the full width at half maximum, and θ is the peak

position. The crystallite size D (nm) of the catalyst particles was calculated from the XRD results (refer to Figure 1), where the crystallite size of zirconium(II) oxide determined from $2\theta = 30^\circ$ was 2.8 nm and nickel from the $2\theta = 44.5^\circ$ was 4.2 nm. The specific surface area of the catalyst was measured using the BET (Brunauer–Emmett–Teller) adsorption method. The catalyst $Zr_{(x)}Ni_{(y)}$ revealed a specific surface area of $88.5 \text{ m}^2 \text{ g}^{-1}$.

Lastly, FE-HRTEM analysis was performed to determine the surface morphologies of the catalyst $Zr_{(x)}Ni_{(y)}$ and the nanocrystal structures (Figure 2). The powder consisted of nanocrystal structures corresponding to zirconium and nickel, represented in zones I, II and III. The Mire fringes with dissimilarity in the interplanar structure were identified. The d spacing was determined in nm, and the identities of each characterised element were confirmed by the XRD phases. Zone I corresponds to the d spacing of 0.14 nm, determined as the zirconium crystal (211) phase; zone II corresponds to the d spacing of 0.13 nm, determined as the nickel crystal (220) phase, as shown in Figure 2c; zone III correlates to the d spacing of the 0.15 nm/(311) phase, corresponding to the Zr_2O particle, as shown in Figure 2d.

Figure 2: Field emission high resolution transmission electron microscopy (FE-HRTEM). (a) Zr_(x)Ni_(y) nanocatalyst, (b) the lattice image of zone I; (c) the lattice image of zone II; (d) the lattice image of zone III.

Characterisation of MgH₂/10 wt.% Zr_(x)Ni_(y) Nanocomposite

After being activated under hydrogen, the Zr_(x)Ni_(y) catalyst was used as an additive in the Mg/MgH₂ system. The commercial MgH₂ was thus milled with 10 wt.% of Zr_(x)Ni_(y) for 5 h, as detailed in Section 4. The dehydrogenation properties of the formed MgH₂/10 wt.% Zr_(x)Ni_(y) nanocomposite were analysed via temperature-programmed desorption (TPD).

Dehydrogenation Properties of the Nanocomposite $\text{MgH}_2/10 \text{ wt.}\% \text{Zr}_{(x)}\text{Ni}_{(y)}$

To understand how the $\text{Zr}_{(x)}\text{Ni}_{(y)}$ catalyst impacts the dehydrogenation properties of MgH_2 , the hydrogen released properties of the nanocomposite $\text{MgH}_2/10 \text{ wt.}\% \text{Zr}_{(x)}\text{Ni}_{(y)}$ were compared with those of the as-received (un-milled MgH_2) and the milled MgH_2 (in similar conditions as used for nanocomposite preparation). Figure 3 represents the temperature profiles of the MgH_2 decomposition, whilst Figure 4 indicates the hydrogen capacity. The milled MgH_2 released 6.74 wt.% of hydrogen in a wide range of desorption temperatures between 300 °C and 400 °C, which may be a result of a large particle size distribution. It is worth noticing that the milled powder presents an important decrease in the decomposition temperature compared with the as-received powder, which is directly related to the reduction in the particle sizes of MgH_2 (the mean size of crystallites was approximately 34 nm for the milled MgH_2 versus approximately 213 nm for the as-received MgH_2 sample, as deduced from XRD analysis by Rietveld method). The obtained result correlated with the findings reported in previous studies [22,23]. The addition of 10 wt.% $\text{Zr}_{(x)}\text{Ni}_{(y)}$ nanocatalyst to MgH_2 induced the shift of the onset hydrogen desorption temperature from 300 °C to a starting temperature below 210 °C. As observed for the nanocomposite $\text{MgH}_2/10 \text{ wt.}\% \text{Zr}_{(x)}\text{Ni}_{(y)}$, only 5.9 wt.% of hydrogen was released in a wide range of temperatures between 200 °C and 400 °C. This result was expected due to the high amount of additive used (10 wt.% of $\text{Zr}_{(x)}\text{Ni}_{(y)}$), which reduced the overall storage capacity of the system. To evaluate the effect of temperature on the discharge characteristics of the sorption materials, the desorption kinetic rates of MgH_2 as-received (un-milled), MgH_2 (milled 5 h) and $\text{MgH}_2/10 \text{ wt.}\% \text{Zr}_{(x)}\text{Ni}_{(y)}$ (milled 5 h) were calculated from the TPD curves of Figure 3, at peak desorption temperatures of 418 °C, 328 °C and 232 °C, respectively; these are shown in Table 2.

Figure 3: The evolution of the temperature-programmed desorption (TPD) profile of as-received MgH₂ after milling without/with 10 wt.% Zr_(x)Ni_(y). Experimental conditions: from ambient temperature to 500 °C under an argon flow of 20 mL min⁻¹ and at a heating rate of 2 °C min⁻¹.

Figure 4: Thermally programmed H₂ desorption capacity curves of studied powders. Same experimental conditions as given in Figure 3.

Table 2: Desorption kinetic properties of pure MgH₂ as-received (un-milled), MgH₂ (ball-milled 5 h) and MgH₂/10 wt.% Zr_(x)Ni_(y). The samples were analysed via TPD from room temperature to 500 °C under an argon flow of 20 mL min⁻¹ and at a heating rate of 2 °C min⁻¹.

Samples	Peak Desorption Temperature (°C)	Discharge Rate (H-wt.% min ⁻¹)	Discharge Capacity (H-wt.%)
MgH ₂ as-received (un-milled)	418	1.51	7
MgH ₂ (milled 5 h)	328	0.17	6.7
MgH ₂ /10 wt.% ZrNi	232	0.07	5.9

Characterisation of Crystal Structures

XRD was carried out to quantify the compositions of the crystalline phases of the composite after the milling process with MgH₂ powder. The patterns of the composite MgH₂/10 wt.% Zr_(x)Ni_(y) were superimposed on the reference MgH₂ (un-milled) powder to reveal broadening in the diffracted line of the nanocomposite sample, which resulted from the grain refining of the powders during the planetary ball milling process (refer to Figure 5a). The composite powder obtained after 5 h milling in an inert atmosphere consisted of the Bragg peaks β -MgH₂ (PDF file #: 00-012-0697) at 2θ of 27.95°, 35.74°, 39.86° and 54.62°; Ni (PDF file #: 00-001-1258) at 2θ of 44.37° and 51.60°; Mg_{0.996}Zr_{0.004} (PDF file #: 01—71-9629) at 2θ of 32.23°, 34.42°, 36.67°, 47.88°, 57.480°, 63.13°, 67.45° and 68.75°. There was a small fraction of fcc-MgO observed at 2θ of 43.04°. The powder was mainly oxidised as a result of the sample preparation for the X-ray analysis handled outside the glove box, as shown in Figure 5b, although a low amount of MgO was also revealed in the as-received MgH₂.

Figure 5: The XRD patterns of (a) 10 wt.% $\text{Zr}_{(x)}\text{Ni}_{(y)}$ nanocomposite in red and the un-milled MgH_2 used as reference in black and (b) $\text{MgH}_2/10\text{ wt.\% } \text{Zr}_{(x)}\text{Ni}_{(y)}$ nanocomposite to identify the crystal phase of each element.

The SEM was used to determine the distribution of the catalyst on the surface of magnesium and to determine the size of the particles. There is a homogenous distribution of the catalyst particles, zirconium and nickel, which can be well distinguished from the SEM image, as seen in Figure 6a. The SEM image of the milled MgH_2 powder used as a reference is shown in Figure 6b. The SEM image of the nanocomposite $\text{MgH}_2/10\text{ wt.\% } \text{Zr}_{(x)}\text{Ni}_{(y)}$ in Figure 6c shows the distribution of the catalyst on the surface and edge of the MgH_2 matrix. The zirconium particles are the white spots, with an average particle size of 10 nm in diameter, as shown in Figure 7a. This correlates with the study

reported by El-Eskandarany et al. [15] where the milled Zr_2Ni alloy powder agglomerated to form particle sizes of approximately 8 nm in diameter. The average particle size of nickel is 100 nm in diameter, as shown in Figure 7b. The wide range of MgH_2 particle size distribution of the composite $\text{MgH}_2/10 \text{ wt.}\% \text{ Zr}_{(x)}\text{Ni}_{(y)}$ is below 1000 nm, as shown in Figure 7c. Pighin et al. [13] reported a study of a $\text{MgH}_2/10 \text{ wt.}\% \text{ Zr}_8\text{Ni}_{21}$ alloy. They established that the white spots were the result of the elastic collisions of the backscattered electrons with the heavy nuclei of zirconium, considering smaller particle sizes under 1000 nm, whereas the grey region, as shown in Figure 6c, was reported to occur due to the light nuclei of magnesium hydride [13]. Figure 6d shows the zoomed image of the $\text{MgH}_2/10 \text{ wt.}\% \text{ Zr}_{(x)}\text{Ni}_{(y)}$ nanocomposite, which represents the catalyst aggregation on the surface MgH_2 matrix. The effect of powder aggregation was reported in previous study to be the result of the high concentration of magnesium deposited at or near the edges of the catalyst [6]. Figure 6e represents the smooth surface of the matrix corresponding to MgO that formed during the handling of the powder outside the glove box.

Figure 6: SEM micrograph of (a) $Zr_{(x)}Ni_{(y)}$ based nanocatalyst after 5-h heat treatment under hydrogen; (b) MgH_2 after 5 h milling time under argon, here used for reference; (c–e) $MgH_2/10$ wt.% $Zr_{(x)}Ni_{(y)}$ nanocomposite. The homogeneous distributions of the $Zr_{(x)}Ni_{(y)}$ on the MgH_2 surface are shown in (c). The enlarged image of the $MgH_2/10$ wt.% $Zr_{(x)}Ni_{(y)}$ showing the aggregation of $Zr_{(x)}Ni_{(y)}$ is represented in (d) and the presence of MgO is represented in (e).

Figure 7: Histograms of particle size distribution measured by the SEM/ImageJ analysis. (a) Zr; (b) Ni size distribution within the catalyst sample; (c) milled MgH₂.

The EDX was used to confirm the presence of the analysed elements and impurities. Figure 8a shows the activated Zr_(x)Ni_(y) based nanocatalyst. The presence of impurities such as chlorine (Cl) came from the starting precursor material ZrCl₄ powder. Figure 8b shows the presence of magnesium in the milled MgH₂. Figure 8c shows the presence of magnesium, zirconium and nickel particles in the MgH₂/10 wt.% Zr_(x)Ni_(y) nanocomposite. The additional peaks are gold (Au), from the coating during the sample preparation of the SEM analysis; carbon (C) from the carbon tape; oxygen (O) from preparing the sample outside the gloves box, and iron (Fe) from the reactor during the milling process.

(a)

(b)

(c)

Figure 8: The energy dispersive X-ray spectroscopy (EDX) of (a) $\text{Zr}_{(x)}\text{Ni}_{(y)}$ nanocatalyst; (b) milled- MgH_2 as reference; (c) $\text{MgH}_2/10 \text{ wt.}\% \text{Zr}_{(x)}\text{Ni}_{(y)}$ nanocomposite.

The TEM analysis was performed to determine the surface morphologies of the composite $\text{MgH}_2/10 \text{ wt.}\% \text{Zr}_{(x)}\text{Ni}_{(y)}$ on a nanometre scale and to understand the catalytic effect of

$Zr_{(x)}Ni_{(y)}$ on the dehydrogenation properties of MgH_2 powder. The TEM image of the composite can be seen in Figure 9a, where the atomic array with a long-range ordered crystal structure is represented in the three zones. Zone I corresponds to the d spacing of 1.63 nm in Figure 9b and correlates with the $(Mg_{0.996}Zr_{0.004})$ (110) phase. Zone II corresponds to the d spacing of 2.58 nm, as shown in Figure 9c, which correlates with the $(Mg_{0.996}Zr_{0.004})$ (002) phase. Zone III is confirmed by the d spacing of 2.52 nm, as shown in Figure 9d, which correlates with the nanocrystalline β - MgH_2 grain and is well-matched with the tetragonal- MgH_2 (101) phase. In Figure 10, the darker particles show the agglomeration of the metallic powder $Zr_{(x)}Ni_{(y)}$ based nanocatalyst on the surface of the MgH_2 matrix after milling with the catalyst. Nickel particles are the dark grey elements covering the surface of the magnesium matrix, and the small, darkest spots are the zirconium particles. The agglomeration of the $Zr_{(x)}Ni_{(y)}$ was also observed in a previous study, where a heterogeneous distribution catalyst on the surface of the MgH_2 powder was reported [15].

Figure 9: Field emission high resolution transmission electron microscopy (FE-HRTEM) of (a) $\text{MgH}_2/10 \text{ wt.}\% \text{ Zr}_{(x)}\text{Ni}_{(y)}$ nanocomposite; (b) the lattice image of zone I; (c) the lattice image of zone II; (d) the lattice image of zone III.

Figure 10: TEM of the bright field image of MgH_2 powders milled with 10 wt.% $\text{Zr}_{(x)}\text{Ni}_{(y)}$, demonstrating the agglomeration of catalyst on the surface of the MgH_2 matrix after 5 h milling time.

Discussion and Conclusions

With the aim of improving the kinetic and thermodynamic properties of the Mg/MgH_2 hydrogen storage system, $\text{Zr}_{(x)}\text{Ni}_{(y)}$ catalyst was synthesised via a modified sol-gel method, in a compact high-pressure hydrogen reactor, and activated prior to being used as an additive to MgH_2 . The TPD results confirmed that the addition of $\text{Zr}_{(x)}\text{Ni}_{(y)}$ to MgH_2 induced an important shift towards the lower hydrogen desorption temperature. This positive impact may be due to several factors. First of all, the activation of the catalyst prior to the milling process with the MgH_2 metamorphosed the alloys into fine powders; as a result, this generated a more homogeneous catalyst distribution [14].

The grain size reduction of the current catalyst was confirmed by the SEM image analysis. The reported size of the zirconium particles in the present study agreed with a number of literature reports [15] and was up to ten times smaller in comparison to the report by Pighin et al. [13]. Although the SEM and TEM analyses confirmed that the catalyst is heterogeneously distributed, it aggregates with a high concentration at the surface and near the edges of the MgH_2 power. The TEM micrograph in Figure 9a shows that the catalyst covered the surface of the MgH_2 during the milling process. The efficiency of the ball milling procedure in dispersing the $\text{Zr}_{(x)}\text{Ni}_{(y)}$ nanocatalyst on the MgH_2 was also reported by M. S. El-Eskandarany et al. [6]. Additionally, because of the ductility differences between $\text{Zr}_{(x)}\text{Ni}_{(y)}$ and MgH_2 particles, the efficiency of the MgH_2 ball milling preparation increases in the presence of the catalyst [24,25].

The combinative impact of MgH_2 crystal size reduction and $\text{Zr}_{(x)}\text{Ni}_{(y)}$ nanocatalyst dispersion at the surface of MgH_2 particles dramatically enhances the dehydrogenation properties in comparison with pure-milled MgH_2 , as shown in Figure 3, and the results from the literature using different $\text{Zr}_{(x)}\text{Ni}_{(y)}$ catalysts are listed in Table 3. This is likely to be due to the presence of zirconium oxide in the starting $\text{Zr}_{(x)}\text{Ni}_{(y)}$ catalyst, as confirmed in the XRD analyses of Figure 1. Indeed, ZrO_2 has been recognised in the past as a transition metal oxide catalyst with the advantage of reducing the activation energy and the crystalline size during the ball milling process [25,26]. In addition, I.Yu. Zavaliy et al. [27] compared the hydrogenation of $\text{Zr}_3\text{V}_3\text{O}_x$ to the $\text{ZrV}_2\text{H}_{4.9}$ catalysts and reported that the zirconium based oxygen catalyst increased the hydrogen storage capacity. However, the presence of ZrO_2 in the $\text{MgH}_2/\text{Zr}_{(x)}\text{Ni}_{(y)}$ composite does not explain the differences to results from literature on its own.

The XRD results presented in Figure 5 show the presence of a new $\text{Mg}_{0.996}\text{Zr}_{0.004}$ phase formed during the milling process; this phase may impact the thermodynamic properties of the Mg/MgH_2 system. In a previous study reported by Galey et al. [9], the presence of a $\text{Mg}_{0.992}\text{Ni}_{0.008}$ phase in a Mg/MgH_2 system doped with a nickel complex reduced the overall hydrogenation

and dehydrogenation enthalpies of the formed composite from ± 75 to ± 65 kJ mol⁻¹ H₂. The presence of the new phase is particularly remarkable as the destabilisation of MgH₂ is the main obstacle to overcome in order to allow the use of Mg/MgH₂ system at moderate temperatures. Despite the important number of studies devoted to the activation of MgH₂, to date, very few researchers have succeeded in modifying its thermodynamic properties.

Table 3: Dehydrogenation properties of various ball milled MgH₂ nanocomposites in comparison to the results obtained in the present study.

Elements	Peak Desorption Temperature °C	H ₂ . wt. %	References
10 wt.% Zr _(x) Ni _(y)	232	5.9	Present study
10 wt.% Zr _{0.67} Ni _{0.33}	325	5.0	Dong et al. [26]
10 wt.% Zr ₈ Ni ₂₁	300	5.9	Pighin et al. [13]
10 wt.% ZrNi ₅	275	5.3	El-Eskandarany et al. [6]
10 wt.% Zr ₉ Ni ₁₁	250	5.9	Dehouche et al. [14]

Although not studied here, the Zr_(x)Ni_(y) alloy is likewise expected to have a positive effect on the hydrogenation properties of the Mg/MgH₂ system. Undoubtedly, the presence of the nanocatalyst sites well dispersed on the Mg particles may allow for the splitting of the hydrogen molecules into hydrogen atoms as well as their adsorption, followed by the subsequent migration of hydrogen atoms onto the adjacent Mg surface via spillover and further surface diffusion [15].

In summary, a novel approach was proposed for the fabrication of the catalyst Zr_(x)Ni_(y) by a high-pressure reactor. The approach demonstrates that mixing such nanocatalysts with MgH₂ via the ball milling process offers promising perspectives. The developed method is a faster and promising approach for the fabrication of catalysts, as it is beneficial in terms of producing nanoparticle materials for improving the hydrogenation/dehydrogenation processes in future work. Therefore, there is work to be done to ameliorate the fabrication

of the catalyst, such as conducting a number of experiments to find out if avoiding the presence of oxygen formation may ameliorate the work of the catalyst and further improve the kinetic and thermodynamic properties of hydrogen storage.

Materials and Methods

Catalyst Synthesis

The catalyst was firstly prepared in the high-pressure reactor PARR 5500 Series Compact Reactor, from a mixture of the starting material of ZrCl_4 fine white powder with a 99.5+ metal basis and $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ (98% green salt) provided by Alpha-Aesar and Sigma-Aldridge, respectively. The solution, 0.99 wt/v.% ZrCl_4 and 1.52 wt/v.% $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, dissolved at room temperature in 150 mL of methanol solvent, was transferred into the reactor vessel (Type 316/austenitic chromium-nickel stainless steel containing molybdenum). The reactor was assembled and tightened vigorously to avoid the pressure drop of the hydrogen. The reactor vessel was purged with hydrogen for several minutes to flush out the air inside the reactor and refill it with hydrogen. The mixture was heated for 2 h at 200 °C under hydrogen, at a starting pressure of 20 bar, whilst stirring at 60 resolutions per minute. The sample was cooled overnight; the supernatant was discarded, whilst the formed gel was dried overnight at room temperature. The sample was fully dried when heat-treated at 350 °C for 5 h and further cooled for 2 h under the flow of hydrogen at 0.6 bar (Figure 11). The prepared catalyst was transferred to the glove box, purged with Argon, then transferred into the reactor vessel for activation via a particulate system HPVA at 30 °C/10 bar.

Figure 11: The steps of synthesis of the catalyst preparation.

Preparation of MgH₂-Based Nanocomposite Powders

The MgH₂/Zr_(x)Ni_(y) composite was prepared under an inert argon atmosphere via the planetary ball miller using a PM100 apparatus from Retsch. A total of 10 wt.% of the hydrogenated Zr_(x)Ni_(y) based nanocatalyst was added to MgH₂ in a reactor vessel with zirconium dioxide (ZrO₂) milling balls of 1-mm diameter. The ball to powder mass ratio was 100:1. The reactor was hermetically sealed inside the glove box to prevent any oxidation of the powders. The sample was milled for 5 h at a frequency of 300 revolutions per minute. During the milling process, the rotation of the ball in the mill was paused every 2 min and inverted every 5 min to prevent the increase of temperature in the milling cell. For comparison, MgH₂ was milled under the same conditions but without the addition of the catalyst.

Sample Characterisations

The properties of the synthesised catalyst and the formed composites were studied through X-ray diffraction (XRD). The crystal structures of all samples were investigated by the XRPD Bruker D8 Advance X-Ray Powder Diffractometer (XRPD) with the CuK α (λ = 0.154 nm) radiation over the 2θ ranging from 20° to 80°. The instrument was calibrated with the LaB6 NIST standard (660a). The specific surface area of the synthesised catalyst was measured with nitrogen gas (NOVAtouch LX1, USA) at a temperature of 77 Kelvin. The surface area was determined according to the Brunauer–Emmett–Teller (BET) equation.

The surface morphologies were obtained with a field emission scanning electron microscope (FESEM) (SUPRA 35VP, Zeiss, Germany). Energy dispersive X-ray spectroscopy (EDS) was used for elemental analysis of the deposited layers. For the SEM analysis, the powder placed on a carbon tape was firstly coated with the addition of a thin layer of gold, added through vacuum evaporation to prevent the sample from developing high conductivity and some charging effects during the analysis.

The TEM model JE M-2100F microscope was used for the structural features. The powder was suspended in EtOH and drop-cast onto holey carbon support film, Cu 200 mesh. The surface morphologies of the analysed powders were determined via the field emission scanning electron microscope (FESEM) (SUPRA 35VP, Zeiss, Germany). The images were collected using Gatan software.

The Dehydrogenation Behaviour

The dehydrogenation properties of the composite $\text{MgH}_2/10 \text{ wt.}\%$ $\text{Zr}_{(x)}\text{Ni}_{(y)}$ were studied after the $\text{Zr}_{(x)}\text{Ni}_{(y)}$ catalyst activation under 10 bars of hydrogen at 30 °C for 100 min. Temperature-programmed desorption (TPD) analyses were performed with a TPD/R/O-1100 apparatus from Thermo. To prevent oxidation, all the handlings were performed in a glove box under a protective argon atmosphere. The experiments were performed from ambient temperature to 500 °C at 2 °C min⁻¹ under an argon flow of 20 mL min⁻¹. To determine the amount of hydrogen desorbed by the samples, the instrument was calibrated prior to the analyses by the reduction of different masses of copper (II) oxide.

References

1. Aguey-Zinsou KF, Ares-Fernández JR. Hydrogen in magnesium: New perspectives toward functional stores. *Energy Environ. Sci.* 2010; 3: 526–543.
2. Sadhasivam T, Kim HT, Jung S, Roh SH, Park JH, et al. Dimensional effects of nanostructured Mg/MgH₂ for hydrogen storage applications: A review. *Renew. Sustain. Energy Rev.* 2017; 72: 523–534.
3. Zhang XX, Liu Y, Zhang XX, Hu J, Gao M, et al. Empowering hydrogen storage performance of MgH₂ by nanoengineering and nanocatalysis. *Mater. Today Nano.* 2019; 9: 100064.
4. Matar SF. Intermetallic hydrides: A review with ab initio aspects. *Prog. Solid State Chem.* 2010; 38: 1–37.

5. El-Eskandarany MS. Metallic glassy Zr₇₀Ni₂₀Pd₁₀ powders for improving the hydrogenation/dehydrogenation behavior of MgH₂. *Sci. Rep.* 2016; 6: 1–13.
6. El-Eskandarany MS, Shaban E, Al-Matrouk H, Behbehani M, Alkandary A, et al. Structure, morphology and hydrogen storage kinetics of nanocomposite MgH₂/10 wt% ZrNi₅ powders. *Mater. Today Energy.* 2017; 3: 60–71.
7. Wang Y, Wang Y. Recent advances in additive-enhanced magnesium hydride for hydrogen storage. *Prog. Nat. Sci. Mater. Int.* 2017; 27: 41–49.
8. El Khatabi M, Naji S, Bhihi M, Benyoussef A, El Kenz A, et al. Effects of double substitution on MgH₂ hydrogen storage properties: An Ab initio study. *J. Alloys Compd.* 2018; 743: 666–671.
9. Galey B, Auroux A, Sabo-Etienne S, Dhaher S, Grellier M, et al. Improved hydrogen storage properties of Mg/MgH₂ thanks to the addition of nickel hydride complex precursors. *Int. J. Hydrogen Energy.* 2019; 44: 28848–28862.
10. Zaluska A, Zaluski L, Ström-Olsen JO. Nanocrystalline magnesium for hydrogen storage. *J. Alloys Compd.* 1999; 288: 217–225.
11. Zaluski L, Zaluska A, Ström-Olsen JO. Hydrogen absorption in nanocrystalline Mg₂Ni formed by mechanical alloying. *J. Alloys Compd.* 1995; 217: 245–249.
12. Jain IP, Lal C, Jain A. Hydrogen storage in Mg: A most promising material. *Int. J. Hydrogen Energy.* 2010; 35: 5133–5144.
13. Pighin SA, Capurso G, Lo Russo S, Peretti HA. Hydrogen sorption kinetics of magnesium hydride enhanced by the addition of Zr₈Ni₂₁ alloy. *J. Alloys Compd.* 2012; 530: 111–115.
14. Dehouche Z, Peretti HA, Hamoudi S, Yoo Y, Belkacemi K. Effect of activated alloys on hydrogen discharge kinetics of MgH₂ nanocrystals. *J. Alloys Compd.* 2008; 455: 432–439.
15. El-Eskandarany MS, Al-ajmi F, Banyan M. Mechanically-Induced Catalyzation of MgH₂ Powders with Zr₂Ni-Ball Milling Media. *Catalysts.* 2019; 9: 382.
16. Nei J, Young K, Regmi R, Lawes G, Salley SO, et al. Gaseous phase hydrogen storage and electrochemical properties of Zr₈Ni₂₁, Zr₇Ni₁₀, Zr₉Ni₁₁, and ZrNi metal

- hydride alloys. *Int. J. Hydrogen Energy*. 2012; 37: 16042–16055.
17. El-Eskandarany MS, Al-Matrouk H, Behbehani M, Shaban E, Alkandary A, et al. Amorphous-versus big cube-Zr₂Ni for improving the kinetics of hydrogenation/dehydrogenation behaviors for MgH₂ powders. *Mater. Chem. Phys.* 2018; 203: 17–26.
18. Molinas B, Ghilarducci AA, Melnichuk M, Corso HL, Peretti HA, et al. Scaled-up production of a promising Mg-based hydride for hydrogen storage. *Int. J. Hydrogen Energy*. 2009; 34: 4597–4601
19. Ruiz FC, Castro EB, Peretti HA, Visintin A. Study of the different Zr_xNi_y phases of Zr- based AB₂ materials. *Int. J. Hydrogen Energy*. 2010; 35: 9879–9887.
20. Nei J, Young K, Salley SO, Ng KYS. Effects of annealing on Zr₈Ni₁₉X₂ (X = Ni, Mg, Al, Sc, V, Mn, Co, Sn, La, and Hf): Hydrogen storage and electrochemical properties. *Int. J. Hydrogen Energy*. 2012; 37: 8418–8427.
21. Parayil SK, Psota RJ, Koodali RT. Modulating the textural properties and photocatalytic hydrogen production activity of TiO₂ by high temperature supercritical drying. *Int. J. Hydrogen Energy*. 2013; 38: 10215–10225.
22. Galey B, Auroux A, Sabo-Etienne S, Grellier M, Postole G. Enhancing hydrogen storage properties of the Mg/MgH₂ system by the addition of bis(tricyclohexylphosphine)nickel(II) dichloride. *Int. J. Hydrogen Energy*. 2019; 44: 11939–11952.
23. Zhang Q, Zang L, Huang Y, Gao P, Jiao L, et al. Improved hydrogen storage properties of MgH₂ with Ni-based compounds. *Int. J. Hydrogen Energy*. 2017; 42: 24247–24255.
24. Galey B, Auroux A, Sabo-Etienne S, Grellier M, Dhaher S, et al. Impact of the addition of poly-dihydrogen ruthenium precursor complexes on the hydrogen storage properties of the Mg/MgH₂ system. *Sustain. Energy Fuels*. 2018; 2: 2335–2344.
25. Chen BH, Chuang YS, Chen CK. Improving the hydrogenation properties of MgH₂ at room temperature by doping with nano-size ZrO₂ catalyst. *J. Alloys Compd.* 2016; 655: 21–27.

26. Dong J, Panda S, Zhu W, Zou J, Ding W. Enhanced hydrogen sorption properties of MgH₂ when doped with mechanically alloyed amorphous Zr_{0.67}Ni_{0.33} particles. Int. J. Hydrogen Energy. 2020.
27. Zavaliy IY. Effect of oxygen content on hydrogen storage capacity of Zr-based h-phases. J. Alloys Compd. 1999; 291: 102–109.