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An investigation of gaseous hydrogen storage characterizations of Mg–Y–Ni–Cu alloys synthesized by melt spinning

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Melt spinning was successfully utilized to prepare Mg_{25–x}Y_xNi₉Cu (x = 0, 1, 3, 5, 7) alloys, producing nanocrystalline and amorphous structures with improved hydrogenation and dehydrogenation performances. The influence of spinning rate on hydrogenation and dehydrogenation thermodynamics and kinetics was studied in detail. XRD and TEM were utilized to characterize the alloy structures. Hydrogenation and dehydrogenation performances were investigated by Sievert apparatus, DSC and TGA connected to a H₂ detector. Dehydrogenation activation energies were estimated using both Arrhenius and Kissinger methods. Results show that melt spinning significantly decreases thermodynamic parameters (ΔH and ΔS) and ameliorates desorption kinetics. Dehydrogenation activation energy markedly lowers with increase in spinning rate and is the real driver of amelioration of dehydrogenation kinetics caused by increasing Y content.

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Introduction

Booming consumption of fossil fuels results in acceleration of their exhaustion and seriously damages the environment, especially through progressively serious global warming and air pollution. Reducing consumption of fossil fuels and preventing further damage to our natural environment are two urgent issues that must be addressed. Universally, wide application of clean energy vehicles is a promising strategy for the above issues due to the fact that transport consumes a great deal of energy worldwide¹ and vehicular combustion of fossil fuels produces about 23% of CO₂ globally.² Recently, the Ministry of Environmental Protection of China published an authoritative investigation report and proclaimed that the most responsible factor for atmospheric haze in Beijing is vehicle exhaust. Hydrogen is regarded as the best fuel of all available clean energy, due to its advantages of no CO₂ emission, inexhaustible supply and high energy efficiency.^{3–5} The overarching technical challenge for transportation and vehicles is the development of a practical hydrogen storage system that can reach the conventional requirement of at least a 300 mile driving range.^{6–8} Thus, development of hydrogen storage technology determines whether a hydrogen fuel cell vehicle could be applied commercially and widely in the 21st century.⁹ Among all hydrogen storage systems, one of the best candidates to satisfy

mobile application is metal hydride.^{10,11} Many studies have been performed on different metal hydrides, seeking a suitable hydrogen storage system. Many metal hydrides have been found and are thought to have potential application prospects. In particular, in the USA, Japan and China, rare-earth-based AB₅- and AB₂-type alloys have achieved mass production. Unfortunately, satisfying all requirements of performance for vehicular application, as presented by the U.S. Department of Energy (DOE), remains an unrealized goal.^{12,13} High theoretical gaseous hydrogen storage capacity (3.6 wt%) and electrochemical capacity (1000 mA h g⁻¹) of Mg₂NiH₄ (ref. 14 and 15) give Mg–Ni-based metal hydrides an advantage in hydrogen storage application in hydrogen fuel cell vehicles and negative electrode material in Ni–MH batteries.¹⁶ Nevertheless, Mg-based alloys are still restricted in practical application by intrinsic imperfections, such as poor hydrogenation/dehydrogenation kinetics, relatively high dehydrogenation temperatures and exceedingly weak electrochemical cyclic stability.^{17,18} Two approaches to ameliorate Mg-based alloys' weaknesses in hydrogen storage properties are alloying and microstructure modification.^{19,20} Particularly, in the Mg₂Ni-type alloy, partially substituting Mg with rare earth elements decreases hydride stability and facilitates hydrogen desorption.^{21–24} Moreover, it has been confirmed that the microstructure of Mg-based alloys has a significant impact on their hydrogenation/dehydrogenation kinetics.^{25,26} As reported by Siarhei *et al.*,²⁷ hydrogenation/dehydrogenation properties of Mg-base alloys are ameliorated drastically when their grain sizes reduce far below micrometer scale. Some preparation technologies, particularly mechanical milling²⁸ and melt spinning,²⁹ have been successfully used in the synthesis of

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nanocrystalline and amorphous structures with the elements distributed homogeneously. Spassov *et al.*³⁰ also found that Mm (Mm = Ce, La-rich misch metals) could significantly increase the hydrogenation capacity of as-spun amorphous and nanocrystalline $\text{Mg}_{75}\text{Ni}_{20}\text{Mm}_5$ alloy. Huang *et al.*³¹ prepared an as-spun amorphous and nanocrystalline $(\text{Mg}_{60}\text{Ni}_{25})_{90}\text{Nd}_{10}$ alloy and found that its maximal discharge capacity is 580 mA h g^{-1} . The improved de-/hydrogenation kinetics of ternary LaCuMg_8 alloy could be explained by the influences of MgCu_2 and Mg_2Cu .³² Transition metals can play the role of catalysts to dissociate the H_2 molecule into absorbable monotonic hydrogen atoms.^{33–35}

To investigate the effects of Y content on hydrogen storage and thermodynamic and dynamic characteristics, we chose Y to partially substitute for Mg in Mg–Ni-based alloy and applied melt spinning technology to prepare nanocrystalline and amorphous $\text{Mg}_{25-x}\text{Y}_x\text{Ni}_9\text{Cu}$ ($x = 0–7$) alloys in the present study.

Experimental

$\text{Mg}_{25-x}\text{Y}_x\text{Ni}_9\text{Cu}$ ($x = 0, 1, 3, 5, 7$) alloys were prepared in a vacuum induction furnace filled with 0.04 MPa helium atmosphere to prevent magnesium volatilization. As the alloys have different Y contents, we labeled these alloys as Y_0 , Y_1 , Y_3 , Y_5 and Y_7 for convenience. Cast ingots were fabricated by pouring the molten alloy in a copper cooled mould. An appropriate number of as-cast alloys were re-melted and dealt by melt-spinning with spinning rates set at 15, 20, 25 and 30 m s^{-1} . As the accurate cooling rate of samples was difficult to measure, we treated the linear velocity of copper roller as the approximate spinning rate.

X-ray diffraction (XRD) (D/max/2400) equipped with a $\text{CuK}_{\alpha 1}$ radiation filtered by graphite was used for determining the

phase structure of alloys. Experimental parameters were set at 160 mA, 40 kV and $10^\circ \text{ min}^{-1}$.

A high-resolution transmission electron microscope (HRTEM) (JEM-2100F) was used to observe the as-spun alloys at 200 kV. Electron diffraction (ED) was performed to ascertain their crystalline states.

An automatically controlled Sieverts apparatus (temperature error within $\pm 2 \text{ K}$) was used to measure P – C – T curves and hydrogenation/dehydrogenation curves of the alloys; 300 mg sample was weighed for each measurement. Hydrogenation was conducted at 3 MPa initial hydrogen pressure and different temperatures of 553, 573 and 593 K. Dehydrogenation was measured at initial hydrogen pressure of $1 \times 10^{-4} \text{ MPa}$ and the same temperatures corresponding to those for hydrogenation. Dehydrogenation properties of alloys at different heating rates of 5, 10, 15 and 20 K min^{-1} were also measured by differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) (SDT-Q600).

Results and discussion

Microstructure characteristics

XRD was used to analyze structure and phase composition of the as-cast and as-spun $\text{Mg}_{25-x}\text{Y}_x\text{Ni}_9\text{Cu}$ ($x = 0–7$) alloys, as shown in Fig. 1. The as-cast and as-spun Y_0 alloys exhibit narrow and sharp diffraction peaks in Fig. 1(a), indicating a typical crystal structure. Moreover, when the spinning rate reaches 25 m s^{-1} , amorphous structure appears in the alloy, as broad and flat diffraction peaks appear in the XRD curves of as-spun Y_3 alloy. Therefore, it can be conjectured that promotion of glass forming in Mg–Ni-based alloys was caused by Y substituting Mg. Fig. 2 shows SEM images and EDS spectra of alloys and demonstrates that Y_0 alloy is consistent with major

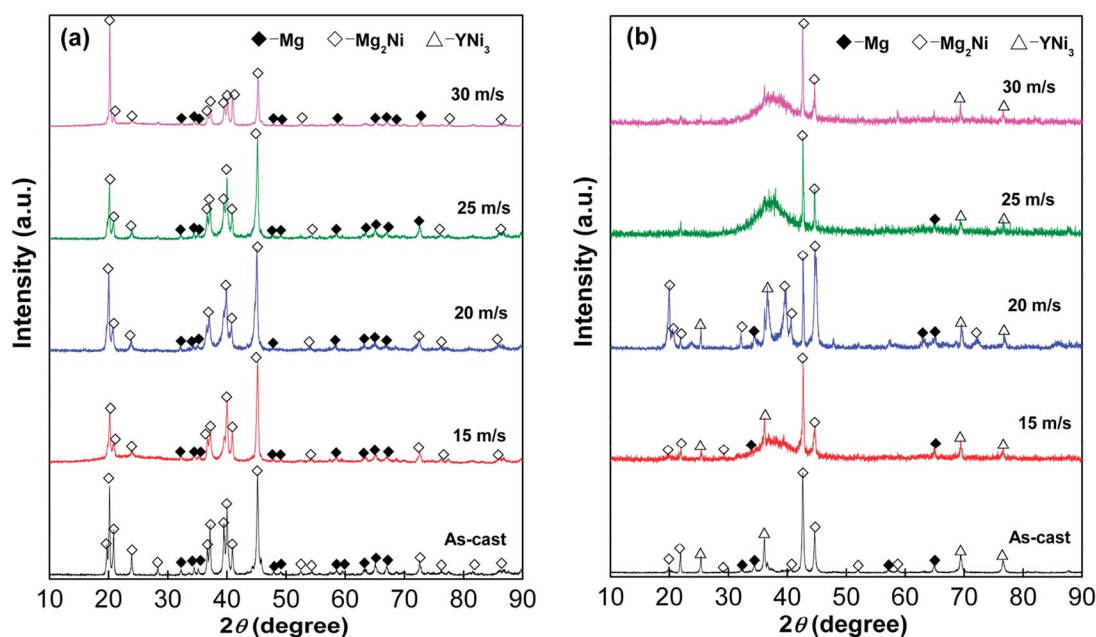


Fig. 1 XRD profiles of as-cast and as-spun Y_0 and Y_3 alloys: (a) Y_0 alloy and (b) Y_3 alloy.

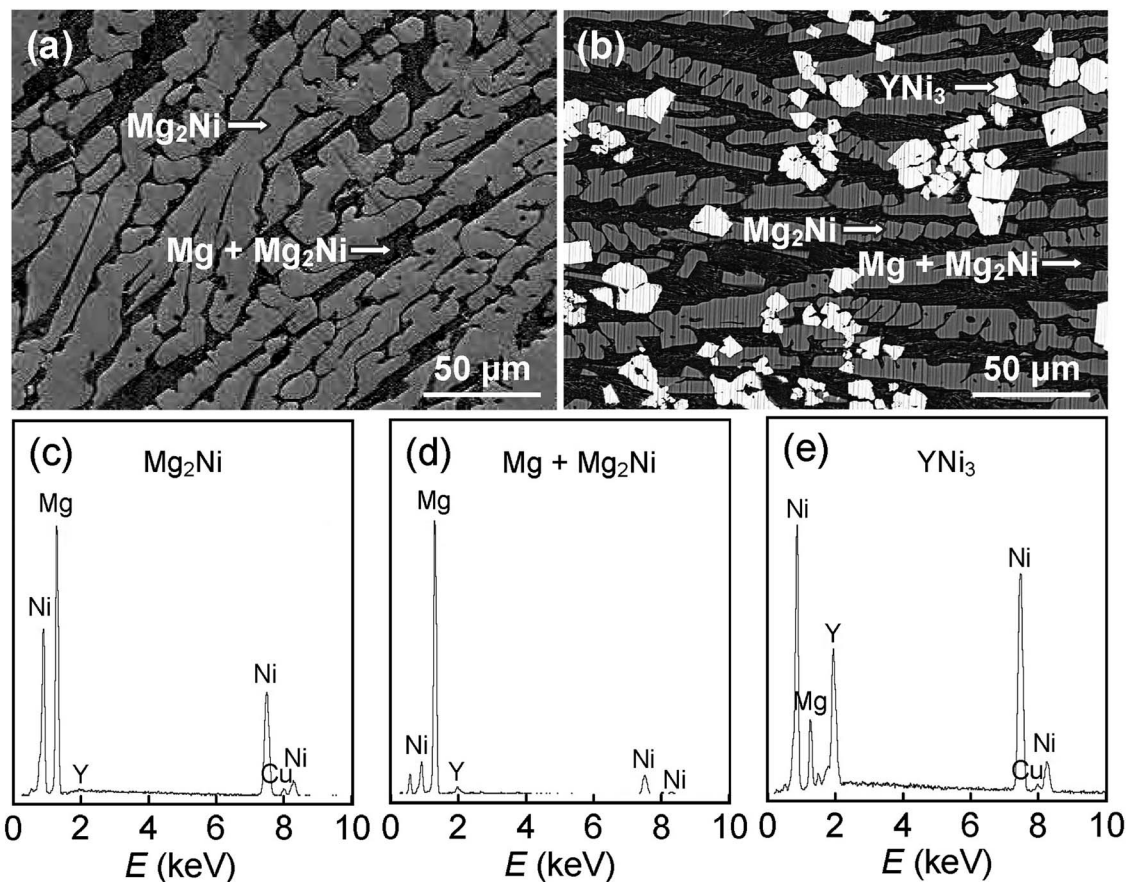


Fig. 2 (a and b) SEM images and (c–e) typical EDS spectra of as-cast alloys: (a) Y_0 alloy and (b) Y_3 alloy.

phase Mg_2Ni and secondary phase Mg. Addition of Y brings on the appearance of secondary phase YNi_3 while keeping the Mg_2Ni and Mg phases unaltered, as can be seen comparing Fig. 1(a) and (b) and by SEM analysis, as shown in Fig. 2.

Fig. 3 shows morphologies and crystal states of as-spun alloys, measured by HRTEM and ED, respectively. The crystalline alloy was disordered by melt spinning and forms a nanostructure. Meanwhile, some crystal defects appear, for instance, subgrains and grain boundaries. Through the linear intercept method, we confirm the grain size range to be 20–30 nm on average. Careful observation finds nanocrystalline structure in as-spun Y_0 alloys and sharp multi-haloes in their ED patterns, which confirms nanocrystalline structure of Mg_2Ni and Mg phases in the alloys. Through the Debye–Scherrer rings, the structure of as-spun Y_3 alloy is confirmed to show both clear nanocrystalline and amorphous features.

Hydrogen storage thermodynamics

Fig. 4 shows the P – C – T curves of the as-prepared $Mg_{2.5-x}Y_xNi_9Cu$ ($x = 0$ – 7) alloys at 573 K. It is clear that there are two pressure plateaus in each P – C – T curve, corresponding to the hydriding and dehydriding processes of Mg/ MgH_2 (lower pressure plateau) and Mg_2Ni/Mg_2NiH_4 (higher pressure plateau), respectively. Moreover, the pressure plateaus of hydrogen absorption and desorption show clear gradients and have large hysteresis ($H_f = \ln(P_a/P_d)$). Melt spinning visibly lifts the

absorption and desorption pressure plateaus, which is ascribed to the increase in internal strain introduced by melt spinning. It is worth noting that the increase in spinning rate has a marked influence on hydrogen storage capacity; the correspondence between spinning rate and hydrogen storage capacity is shown in the inset of Fig. 4. Evidently, the hydrogen storage capacity first increases and later decreases with the increase in spinning rate.

The increase in hydrogenation capacity can be attributed to the refined grain size, namely, formation of a nanocrystalline structure.³⁶ As the spinning rate is further increased, more amorphous phase is formed inside the alloy, resulting in a decrease in hydrogenation capacity of the alloy. Maximal capacities of Y_0 and Y_3 alloys are 4.36 wt% and 3.65 wt% respectively, which indicates that substitution of Mg by Y leads to a significant decrease in hydrogen absorption capacity. It is well known that hydrogenation and dehydrogenation thermodynamic parameters of an alloy are the main standards to evaluate thermal stability of the hydride, which can be derived with Van't Hoff equation as shown below:³⁷

$$\ln[P(H_2)/P_0] = \Delta H/(RT) - \Delta S/R \quad (1)$$

where $P(H_2)$ is the equilibrium hydrogen gas pressure, P_0 is 1 atm, R is the gas constant and T is the temperature of the alloy.

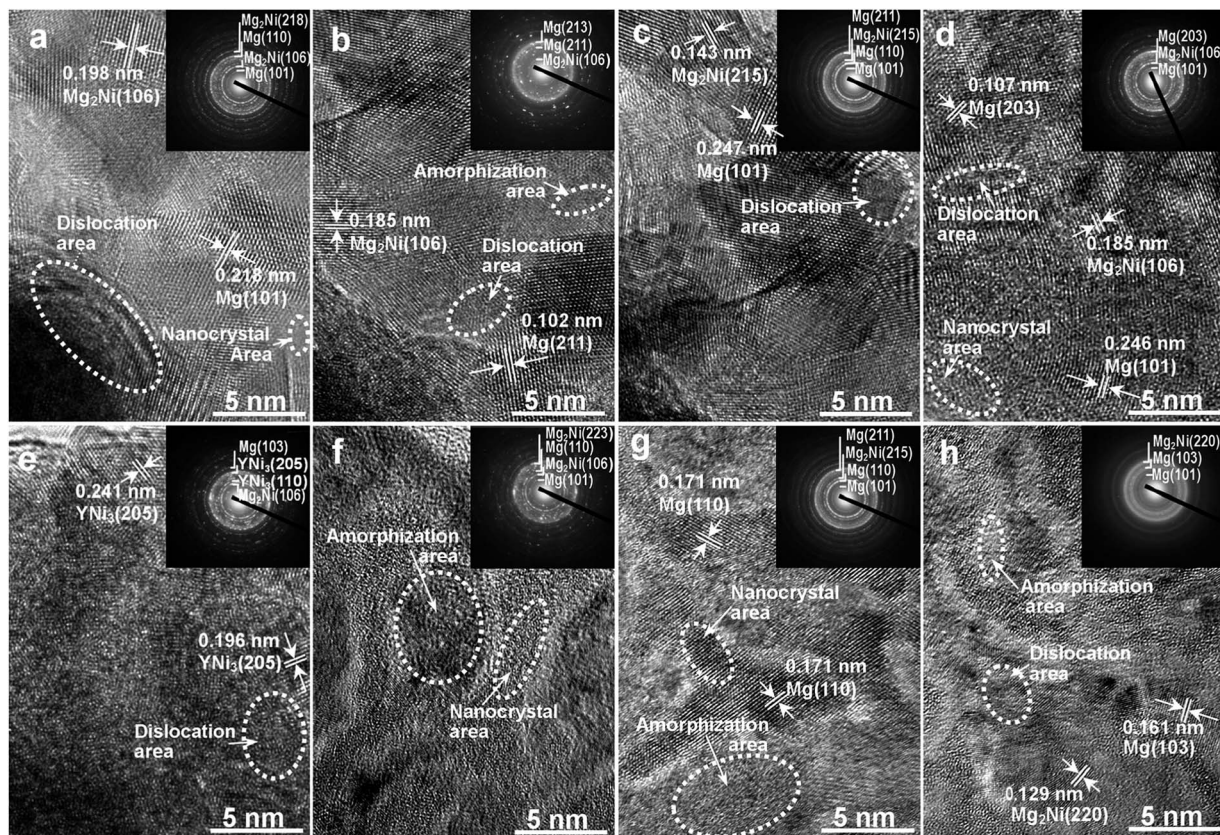


Fig. 3 HRTEM micrographs and ED patterns of as-spun alloys: (a), (b), (c) and (d) Y_0 alloy, spun at 15, 20, 25 and 30 $m s^{-1}$, respectively; (e), (f), (g) and (h) Y_3 alloy spun at 15, 20, 25 and 30 $m s^{-1}$, respectively.

For using Van't Hoff equation to calculate thermodynamic parameters, three temperatures (553, 573 and 593 K) were chosen to test the P - C - T curves of the as-cast and as-spun (30 $m s^{-1}$) Y_3 alloys, as illustrated in Fig. 5. With the logarithmic transformation of eqn (1), Van't Hoff graphs of $\ln[P(H_2)/P_0]$ vs. $1/T$ for the two alloys can be plotted, as shown in the inset of Fig. 5, based on the data of the P - C - T curves. These figures indicate that $\ln[P(H_2)/P_0]$ is linear against $1/T$. Thus, with the help of eqn (1), enthalpy (ΔH) and entropy (ΔS) can be expediently calculated. The obtained ΔH and ΔS values of the as-cast and as-spun Y_3 alloys are listed in Table 1. The visible decrease in thermodynamic parameters indicates that thermal stability of the alloy hydrides is decreased by melt spinning.

Temperature programmed desorption curves of the as-cast and as-spun Y_0 and Y_3 alloys, which were in a saturated hydrogen absorption condition at 3 MPa and 573 K, were measured in confined space at 5 $K min^{-1}$ heating rate and shown in Fig. 6. Samples were taken in equal weight to avoid the influence on desorption temperature caused by increased pressure. As seen from the figures, the increase in spinning rate reduces the initial hydrogen desorption temperature in as-cast and as-spun alloys, which suggests that melt spinning reduces stability of the hydride. Increasing the spinning rate of alloy from 0 (as-cast alloy) to 30 $m s^{-1}$ results in the decrease in the initial hydrogen desorption temperatures of Y_0 and Y_3 alloys from 532.8 to 504.2 K and from 511.7 to 494.3 K, respectively. Comparison of Fig. 6(a) and (b) demonstrates that for a fixed

spinning rate, Y_3 alloy exhibits a lower starting hydrogen desorption temperature compared with Y_0 alloy, indicating that replacement of Mg by Y reduces thermal stability of hydride. The decreased starting hydrogen desorption temperature caused by melt spinning can be ascribed to formation of the nanocrystalline and amorphous structures because the refined microstructure plays a dominant role in decreasing the dehydrogenation temperature.^{38,39}

Hydrogen absorption and desorption kinetics

Fig. 7 shows the hydriding curves of Y_0 and Y_3 alloys of different spinning rates at 573 K and 3 MPa. The as-cast and as-spun alloys absorb hydrogen quickly during the inception phase and then, the hydrogen absorption rate decreases sharply. It takes a long time for alloys to achieve nearly full hydrogen saturation. Apart from hydrogenation capacity, hydrogen absorption and desorption rates are also important influences on the application of hydrogen storage materials. Herein, hydrogen absorption saturation ratio ($R^a(t)$) was used to characterize hydrogen absorption kinetics of the alloy. $R^a(t)$ is the ratio of absorbed hydrogen content at time t to the maximum absorbed hydrogen content and is expressed as $R^a(t) = C^a(t)/C^a(100) \times 100\%$, where $C^a(t)$ and $C^a(100)$ respectively represent the absorbed hydrogen content at times t and 100 min. The $C^a(100)$ value, which is found to be higher than 98% of the maximum hydrogen absorption capacity, is seen as

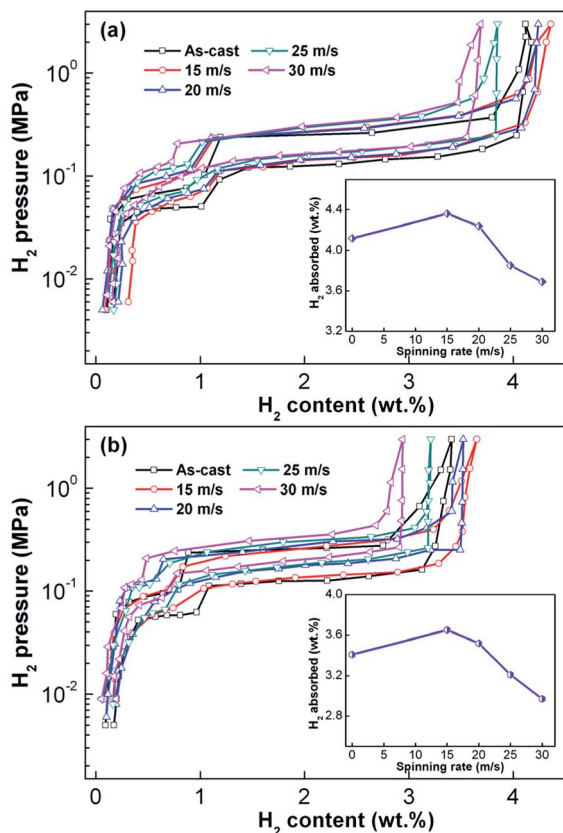


Fig. 4 P - C - T curves of the as-cast and as-spun Y_0 and Y_3 alloys at 573 K: (a) Y_0 alloy and (b) Y_3 alloy.

approximately the maximum absorbed hydrogen content of alloys. We chose 5 minutes as the duration for hydrogen absorption and the changes in $R^a(5)$ ($t = 5$) values with the increase in spinning rate of Y_0 and Y_3 alloys were recorded, as shown in the insets of Fig. 7(a) and (b). The data show that increasing the spinning rate leads to the $R^a(5)$ value of Y_0 alloy to first increase and later decrease, while the value markedly decreases for Y_3 alloy. With the spinning rate increasing from 0 to 30 m s^{-1} , the $R^a(5)$ ($t = 5$) value of Y_3 alloy reduces from 88.8% to 78.6%, but grows from 85.8% (0 m s^{-1}) to 91.5% (15 m s^{-1}) first and then decreases to 83.3% (30 m s^{-1}) for Y_0 alloy.

Similarly, hydrogen desorption ratio ($R^d(t)$) was used to signify the hydrogen desorbing kinetics of the alloy. $R^d(t)$ is the ratio of the desorbed hydrogen content at time t to the maximum absorbed hydrogen content, which can be calculated by $R^d(t) = C^d(t)/C^a(100) \times 100\%$, where $C^d(t)$ represents the desorbed hydrogen content at t min and $C^a(100)$ represents the same as previously mentioned. We chose 10 minutes as the duration of hydrogen desorption, and the changes in $R^d(10)$ ($t = 10$) values with the increase in spinning rate of Y_0 and Y_3 alloys were established, as shown in Fig. 8. It is clear that with the increase in spinning rate, the dehydrogenating rates of Y_0 and Y_3 alloys strictly increase. Specifically, the increase in spinning rate of Y_0 and Y_3 alloys from 0 to 30 m s^{-1} enhances their $R^d(10)$ values from 50.4% to 65.2% and from 61.5% to 71.2%, respectively. This evidences that Y_3 alloy has a higher $R^d(10)$ value

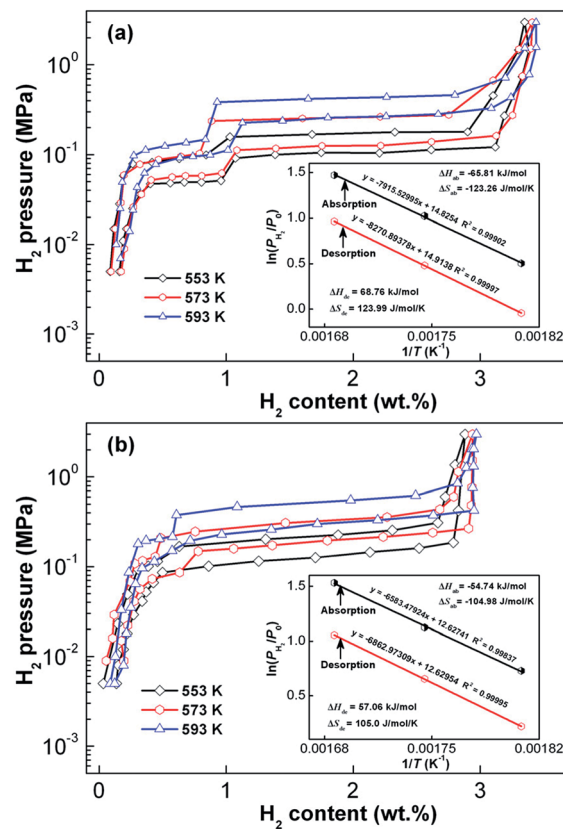


Fig. 5 P - C - T curves of the as-cast and as-spun (30 m s^{-1}) Y_3 alloys in the temperature range of 553–593 K. Van't Hoff plots (insets): (a) as-cast, (b) as-spun (30 m s^{-1}).

Table 1 ΔH and ΔS values of as-spun (20 m s^{-1}) Y_0 and Y_3 alloys

States	Absorption		Desorption	
	ΔH (kJ mol^{-1})	ΔS ($\text{J mol}^{-1} \text{ K}^{-1}$)	ΔH (kJ mol^{-1})	ΔS ($\text{J mol}^{-1} \text{ K}^{-1}$)
As-cast	-65.81	-123.26	68.76	123.99
As-spun	-54.74	-104.98	57.06	105.0

compared with that of Y_0 alloy at a fixed spinning rate, indicating that the substitution of Y for Mg has an ameliorative influence on the hydrogen desorption kinetics of alloys.

It is generally acknowledged that hydrogen absorption rate is decided by the following factors: the decomposition rate of hydrogen molecules on surface, the penetration rate of hydrogen atoms on surface into the bulk metal, and the formation rate of hydride in metal. Among these factors, the slowest step primarily dominates the hydrogen absorption rate. The experimental results reveal that hydrogen storage capacity and kinetics are significantly affected by the variation in spinning rate. Regarding the active role of melt spinning in hydrogen storage capacity and kinetics, the change in alloy structure resulting from melt spinning cannot be ignored. Structure identification shows that melt spinning disorders the

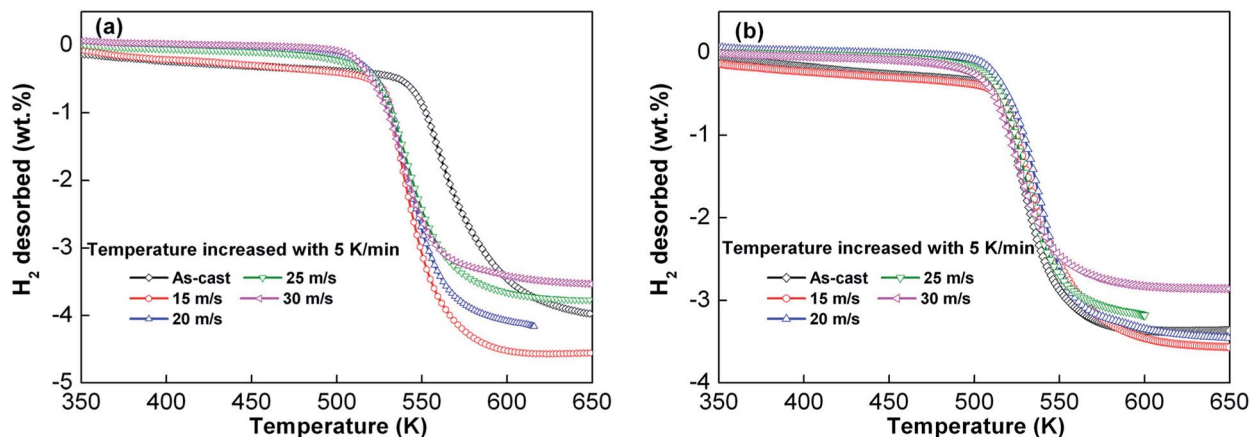


Fig. 6 Temperature programmed desorption curve of as-cast and as-spun Y_0 and Y_3 alloys after hydrogen absorption at a heating rate of 5 K min^{-1} : (a) Y_0 alloy and (b) Y_3 alloy.

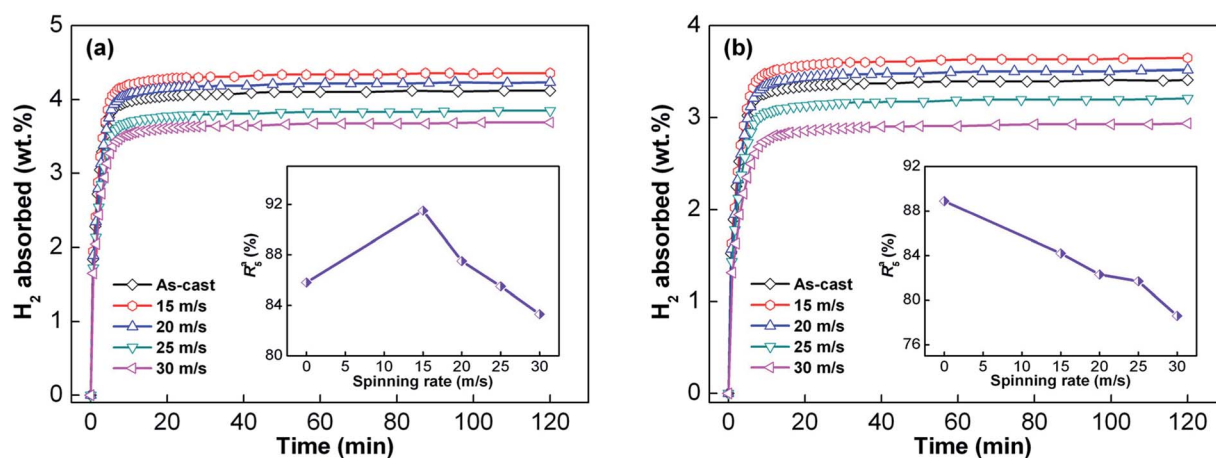


Fig. 7 Hydrogen absorption kinetic curves of the as-cast and as-spun Y_0 and Y_3 alloys at 573 K : (a) Y_0 alloy and (b) Y_3 alloy.

crystalline structure and turns it, or at least part of it, into nanocrystalline. As grain boundary contains a much higher hydrogen concentration than grain interior and amorphous

region,⁴⁰ the hydrogen storage capacity of this alloy is enhanced through melt spinning due to the numerous crystallites and grain boundaries created. Also, a grain boundary facilitates diffusion of hydrogen inside the alloy and improves its hydrogen absorption kinetics by providing numerous sites that require low diffusion activation energy.⁴¹ Regarding the negative influence of melt spinning on hydrogen storage property, the glass phase facilitated by Y replacing Mg and the increase in spinning rate are mainly responsible owing to the fact that it is much easier for hydrogen atoms to diffuse in a nanocrystalline phase than in an amorphous phase. Moreover, the negative influence on hydrogen storage capacity and kinetics resulting from the stored strain energy created during melt spinning cannot be ignored. The promotion in hydrogen desorption kinetics caused by melt spinning is thought to be connected with the decreased grain size and enhanced internal strain because of the drastic improvement in ultrafine powder size (far below micrometer scale) on dehydrogenation performance of Mg-based alloy.⁴² Moreover, Northwood *et al.*⁴³ considered that the increase in spinning rate leads to the growth of stored internal strain, which consequently causes a significant decline

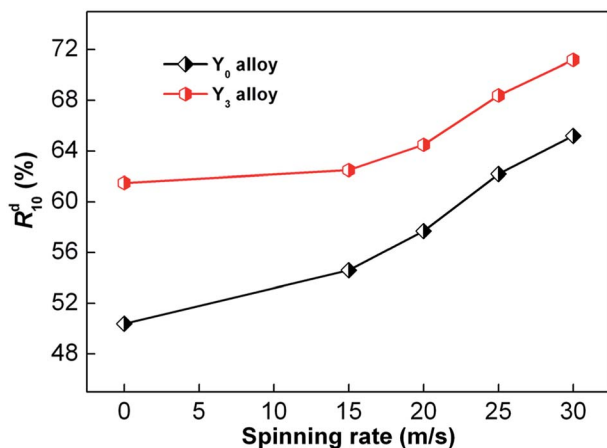


Fig. 8 Evolution of R_{10}^d values of as-cast and as-spun Y_0 and Y_3 alloys with spinning rate.

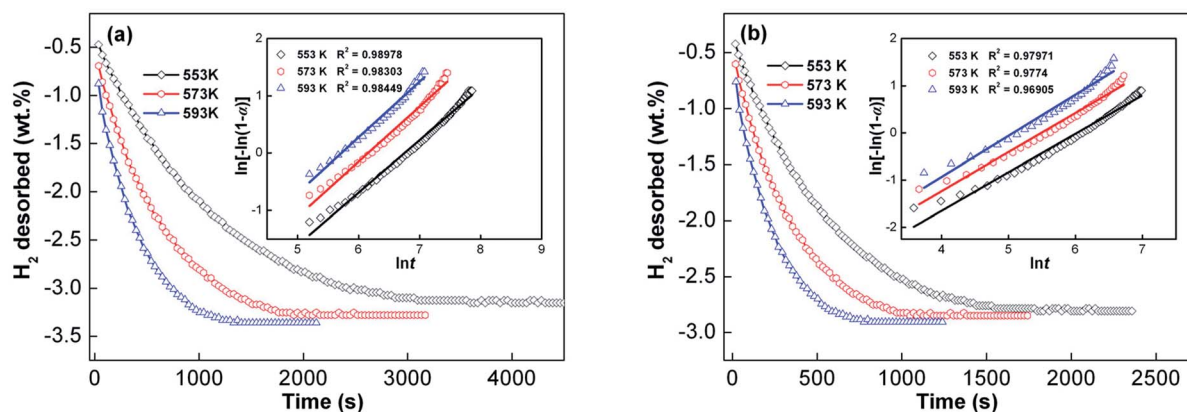


Fig. 9 Hydrogen desorption kinetic curves of the as-cast and as-spun (30 m s^{-1}) Y_3 alloy at 553, 573 and 593 K and Avrami plots of $\ln[-\ln(1 - \alpha)]$ vs. $\ln t$: (a) as-cast and (b) as-spun (30 m s^{-1}).

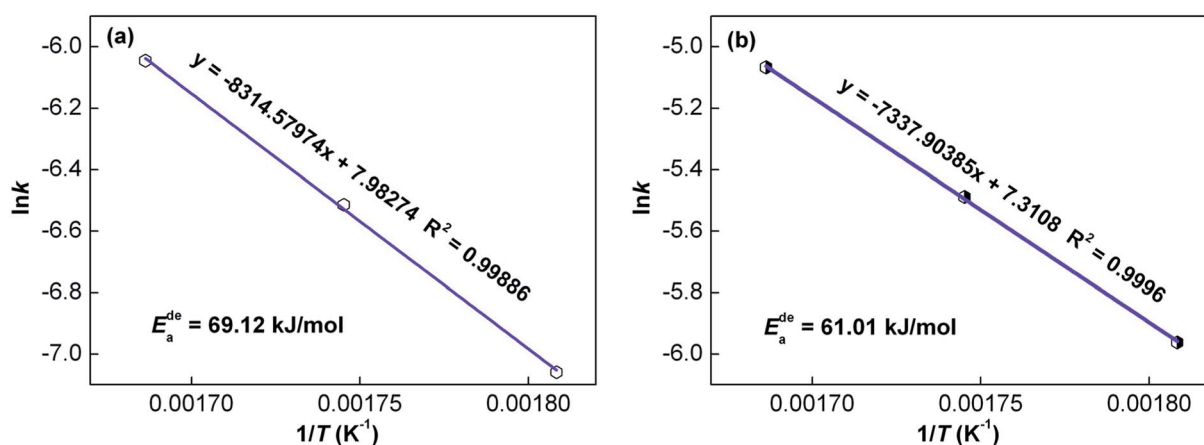


Fig. 10 Arrhenius plots of the as-cast and as-spun (30 m s^{-1}) Y_3 alloy: (a) as-cast and (b) as-spun (30 m s^{-1}).

in the stability of metal hydrides, thus facilitating hydrogen desorption.

Dehydrogenation activation energy

Generally, the total energy barrier that a reaction needs to surmount is known as the activation energy, which is an important factor that determines the kinetics of a gas–solid reaction. With respect to hydrogen desorption reaction, activation energy is the total energy barrier during dehydrogenation processes. Hence, the driving force for the dehydrogenation process can be measured through activation energy. In this study, the Arrhenius and Kissinger methods, two common methods for estimating hydrogen desorption activation energy, were used to reveal the influence of melt spinning on dehydrogenation kinetics. To determine the kinetics of alloys by the Arrhenius method, experimental alloys were measured in the temperature range of 553 to 593 K. The as-cast and as-spun Y_3 alloys were chosen as representatives, and their dehydrogenation kinetic curves are shown in Fig. 9. Usually, the nucleation and growth processes of dehydrogenation can be simulated by the Johnson–Mehl–Avrami (JMA) model with the following JMA equation:⁴⁴

$$\ln[-\ln(1 - \alpha)] = \eta \ln k + \eta \ln t \quad (2)$$

where α represents the phase fraction transformed at time t , k is the rate constant, and η represents the Avrami exponent. With logarithmic transformation of eqn (2), JMA graphs of $\ln[-\ln(1 - \alpha)]$ vs. $\ln t$ can be constructed, as shown in the inset of Fig. 9. The JMA plots are found to be nearly linear, implying that the dehydrogenation reaction of alloys abides by instantaneous nucleation and interface controlled three-dimensional growth process.⁴⁵ Thus, the η and $\eta \ln k$ values at different temperatures can be derived from the slope and intercept of JMA plots and the value of k can be easily calculated. The activation energy (E_a) of dehydrogenation can be derived from the Arrhenius equation as follows:⁴⁶

$$k = A \exp[-E_a/(RT)] \quad (3)$$

where the rate constant k has been defined previously, A is a temperature independent coefficient, and R and T represent the universal gas constant ($8.3145 \text{ J mol}^{-1} \text{ K}^{-1}$) and the absolute temperature of alloys, respectively. Fig. 10 shows the Arrhenius plots of $\ln k$ vs. $1/T$, from the slopes of which the activation

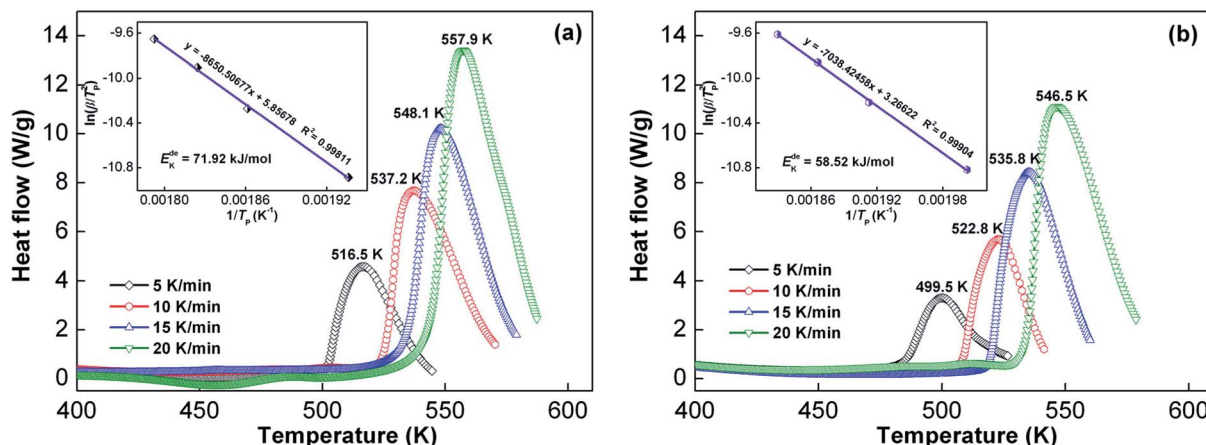


Fig. 11 DSC curves of the as-cast and as-spun (30 m s^{-1}) Y_3 alloy at various heating rates and Kissinger plots: (a) as-cast, (b) as-spun (30 m s^{-1}).

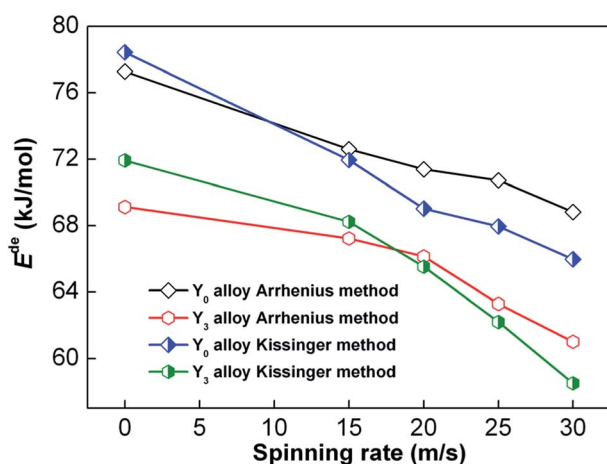


Fig. 12 Evolutions of hydrogen desorption activation energies of Y_0 and Y_3 alloys calculated by Arrhenius and Kissinger methods with increase in spinning rate.

energy E_a can be derived. E_a values of the as-cast and as-spun (30 m s^{-1}) Y_3 alloys are 69.12 and $61.01 \text{ kJ mol}^{-1}$, respectively, indicating that melt spinning can decrease the dehydrogenation activation energy.

To verify the result achieved from the Arrhenius method, the Kissinger method was used for calculating the dehydrogenation activation energy using the Kissinger equation given below:⁴⁷

$$d\{\ln[\beta/(T_p)^2]\}/d(1/T_p) = -E_k/R \quad (4)$$

where β represents the heating rate, T_p represents the absolute temperature corresponding to the maximal desorbing rate of DSC curves, E_k represents the activation energy, and R represents the universal gas constant. DSC curves of the dehydrogenation reactions of the as-cast and as-spun $\text{Mg}_{25-x}\text{Y}_x\text{Ni}_9\text{Cu}$ ($x = 0-7$) alloys, saturated at 3 MPa and 573 K , were recorded at $5, 10, 15$ and 20 K min^{-1} , respectively. We choose the as-cast and as-spun (30 m s^{-1}) Y_3 alloys as representatives and illustrated their dehydrogenation DSC curves in Fig. 11. It is clear that the

shapes of all alloys are similar, with each having an endothermic peak corresponding to the dehydrogenation process, indicating that the reaction processes of all alloys are similar. Furthermore, all of the endothermic peaks of as-spun alloys drift to low temperature at each heating rate, which implies the positive effect of melt spinning on dehydrogenation reaction. On the basis of Fig. 11, Kissinger plots of $\ln[\beta/(T_p)^2]$ vs. $1/T_p$ can be created by utilizing the logarithmic transform of eqn (4), as shown in the inset of Fig. 11. The Kissinger plots are nearly linear and their corresponding slopes can be used to calculate activation energy E_k . The E_k values of as-cast and as-spun (30 m s^{-1}) Y_3 alloys are 71.92 and $58.52 \text{ kJ mol}^{-1}$, respectively.

Arrhenius and Kissinger methods were both used to calculate the activation energy E_{de} ($E_{de} = E_a$ for Arrhenius method and $E_{de} = E_k$ for Kissinger method). Fig. 12 shows the relationship between E_{de} values and spinning rate, which reveals that the E_{de} values markedly decline with the increase in spinning rate. Moreover, it is also noted that Y_3 alloy has a lower E_{de} than Y_0 alloy for any fixed spinning rate. We draw a conclusion from the abovementioned findings that the decrease in dehydrogenation activation energy, which is the result of Y partly substituting Mg and melt spinning, is the real driver for the improvement of dehydrogenation kinetics.

Conclusions

Hydrogen storage thermodynamics and kinetics of as-cast and as-spun $\text{Mg}_{25-x}\text{Y}_x\text{Ni}_9\text{Cu}$ ($x = 0-7$) alloys were systematically studied and the major conclusions are listed below.

(1) Melt spinning results in a significant decrease in thermodynamic parameters (ΔH and ΔS) and a dramatic improvement in dehydrogenation kinetics due to the declining dehydrogenation activation energy incurred by melt spinning.

(2) Melt spinning facilitates reduction of hydride stability. Increasing spinning rate from 0 to 30 m s^{-1} decreases starting dehydrogenation temperature from 532.8 to 504.2 K for Y_0 alloy and 511.7 to 494.3 K for Y_3 alloy.

(3) Dehydrogenation activation energies of experimental alloys were evaluated by both Arrhenius and Kissinger methods.

Result reveal that dehydrogenation activation energy declines with the increase in spinning rate, which is regarded to be the real driver for improvement of dehydrogenation kinetics caused by melt spinning.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 D. Mori and K. Hirose, *Int. J. Hydrogen Energy*, 2009, **34**, 4569–4574.
- 2 R. Lan, J. T. S. Irvine and S. Tao, *Int. J. Hydrogen Energy*, 2012, **37**, 1482–1494.
- 3 I. P. Jain, *Int. J. Hydrogen Energy*, 2009, **34**, 7368–7378.
- 4 Y. Bai, C. Wu, F. Wu, J. H. Yang, L. L. Zhao, F. Long and B. L. Yi, *Int. J. Hydrogen Energy*, 2012, **37**, 12973–12979.
- 5 V. S. Arunachalam and E. L. Fleischer, *MRS Bull.*, 2008, **33**, 264–275.
- 6 L. Schlapbach and A. Züttel, *Nature*, 2001, **414**, 353–358.
- 7 V. Bhat, A. Rougier, L. Aymard, G. A. Nazri and J. M. Tarascon, *Int. J. Hydrogen Energy*, 2007, **32**, 4900–4906.
- 8 Y. Zhang, M. Ji, Z. Yuan, W. Bu, Y. Qi and S. Guo, *RSC Adv.*, 2017, **7**, 37689–37698.
- 9 W. Hui, S. Takenaka and K. Otsuka, *Int. J. Hydrogen Energy*, 2006, **31**, 1732–1746.
- 10 Z. M. Yuan, W. Zhang, P. L. Zhang, Y. H. Zhang, W. G. Bu, S. H. Guo and D. L. Zhao, *RSC Adv.*, 2017, **7**, 56365–56374.
- 11 W. J. Song, J. S. Li, T. B. Zhang, X. J. Hou and H. C. Kou, *Mater. Lett.*, 2015, **145**, 193–196.
- 12 L. E. Klebanoff and J. O. Keller, *Int. J. Hydrogen Energy*, 2013, **38**, 4533–4576.
- 13 L. Shaw, J. Pratt, L. Klebanoff, T. Johnson, M. Arienti and M. Moreno, *Int. J. Hydrogen Energy*, 2013, **38**, 2810–2823.
- 14 D. Chandra, A. Sharma, R. Chellappa, W. N. Cathey, F. E. Lynch, R. C. Bowman Jr., J. R. Wermer and S. N. Paglieri, *J. Alloys Compd.*, 2008, **452**, 312–324.
- 15 N. Juahir, N. S. Mustafa, A. M. Sinin and M. Ismail, *RSC Adv.*, 2015, **5**, 60983–60989.
- 16 A. Ebrahimi-Purkani and S. F. Kashani-Bozorg, *J. Alloys Compd.*, 2008, **456**, 211–215.
- 17 I. P. Jain, C. Lal and A. Jain, *Int. J. Hydrogen Energy*, 2010, **35**, 5133–5144.
- 18 Y. B. Pan, H. Y. Leng, J. Wei and Q. Li, *Int. J. Hydrogen Energy*, 2013, **38**, 5133–5144.
- 19 Y. F. Liu, H. G. Pan, M. X. Gao and Q. D. Wang, *J. Mater. Chem.*, 2011, **21**, 4743–4755.
- 20 A. L. Eric, *Int. J. Hydrogen Energy*, 2011, **36**, 10787–10796.
- 21 T. Spassov and U. Köster, *J. Alloys Compd.*, 1999, **287**, 243–250.
- 22 B. Darriet, M. Pezat, A. Hbika and P. Hagemuller, *Int. J. Hydrogen Energy*, 1980, **5**, 173–178.
- 23 E. A. Lass, *Int. J. Hydrogen Energy*, 2011, **36**, 10787–10796.
- 24 D. H. Xie, P. Li, C. X. Zeng, J. W. Sun and X. H. Qu, *J. Alloys Compd.*, 2009, **478**, 96–102.
- 25 A. Teresiak, A. Gebert, M. Savyak, M. Uhlemann, C. Mickel and N. Mattern, *J. Alloys Compd.*, 2005, **398**, 156–164.
- 26 A. Montone, J. G. Novakovic, M. V. Antisari, A. Bassettib, E. Bonetti, A. L. Fiorini, L. Pasquini, L. Mirengic and P. Rotolo, *Int. J. Hydrogen Energy*, 2007, **32**, 926–2934.
- 27 K. Sjarhei, R. Lars, R. Thomas, G. Thomas, W. Thomas and K. Bernd, *Int. J. Hydrogen Energy*, 2011, **36**, 1592–1600.
- 28 M. Y. Song, Y. J. Kwak, H. S. Shin, S. H. Lee and B. G. Kim, *Int. J. Hydrogen Energy*, 2013, **38**, 1910–1917.
- 29 T. Spassov, L. Lyubenova, U. Koster and M. D. Baro, *Mater. Sci. Eng., A*, 2004, **375–377**, 794–799.
- 30 T. Spassov and U. Köster, *J. Alloys Compd.*, 1999, **287**, 243–250.
- 31 L. J. Huang, G. Y. Liang, Z. B. Su and D. C. Wu, *J. Power Sources*, 2006, **160**, 684–687.
- 32 S. Couillaud, E. Gaudin, J. Andrieux, S. Gorsse, M. Gayot and J. L. Bobet, *Int. J. Hydrogen Energy*, 2012, **37**, 11824–11834.
- 33 J. Cui, J. Liu, H. Wang, L. Ouyang, D. Sun, M. Zhu and X. D. Yao, *J. Mater. Chem. A*, 2014, **2**, 9645–9655.
- 34 M. Pozzo and D. Alfe, *Int. J. Hydrogen Energy*, 2009, **34**, 1922–1930.
- 35 F. Tang, T. Parker, H. F. Li, G. C. Wang and T. M. Lu, *Nanotechnology*, 2008, **19**, 465706.
- 36 W. J. Song, J. S. Li, T. B. Zhang, H. C. Kou and X. Y. Xue, *J. Power Sources*, 2014, **245**, 808–815.
- 37 H. Falahati and D. P. J. Barz, *Int. J. Hydrogen Energy*, 2013, **38**, 8838–8851.
- 38 W. J. Song, J. S. Li, T. B. Zhang, X. J. Hou and H. C. Kou, *RSC Adv.*, 2015, **5**, 54258–54265.
- 39 T. B. Zhang, W. J. Song, H. C. Kou and J. S. Li, *Appl. Surf. Sci.*, 2016, **371**, 35–43.
- 40 T. Spassov, V. Rangelova and N. Neykov, *J. Alloys Compd.*, 2002, **334**, 219–223.
- 41 L. Hima Kumar, B. Viswanathan and S. S. Murthy, *J. Alloys Compd.*, 2008, **461**, 72–76.
- 42 T. Sadhasivam, M. Sterlin Leo Hudso, S. K. Pandey, A. Bhatnagar, M. K. Singh, K. Gurunathan and O. N. Srivastava, *Int. J. Hydrogen Energy*, 2013, **38**, 353–7362.
- 43 H. Niu and D. O. Northwood, *Int. J. Hydrogen Energy*, 2002, **27**, 69–77.
- 44 M. Pourabdoli, S. Raygan, H. Abdizadeh and D. Uner, *Int. J. Hydrogen Energy*, 2013, **38**, 11910–11919.
- 45 T. Czujko, R. A. Varin, Ch. Chiu and Z. Wronski, *J. Alloys Compd.*, 2006, **414**, 240–247.
- 46 T. Kimura, H. Miyaoka, T. Ichikawa and Y. Kojima, *Int. J. Hydrogen Energy*, 2013, **38**, 13728–13733.
- 47 H. E. Kissinger, *Anal. Chem.*, 1957, **29**, 1702–1706.