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# Hydrogen storage in Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> nano particles synthesized from coarse-grained Mg and nano sized Ni(Fe) precursor

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In this work, Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> was synthesized using precursors of nano Ni(Fe) composite powder prepared through arc plasma method and coarse-grained Mg powder. The microstructure, composition, phase components and the hydrogen storage properties of the Mg–Ni(Fe) composite were carefully investigated. It is observed that the Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> particles formed from the Mg–Ni(Fe) composite have a diameter of 100–240 nm and a portion of Fe in the Ni(Fe) nano particles transformed into α-Fe nano particles with the diameter of 40–120 nm. DSC measurements showed that the peak desorption temperature of the Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> was reduced to 501 K and the apparent activation energy for hydrogen desorption of the Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> was 97.2 kJ mol<sup>-1</sup> H<sub>2</sub>. The formation enthalpy of Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> was measured to be –53.1 kJ mol<sup>-1</sup> H<sub>2</sub>. The improvements in hydrogen sorption kinetics and thermodynamics can be attributed to the catalytic effect from α-Fe nano particles and the destabilization of Mg<sub>2</sub>NiH<sub>4</sub> caused by the partial substitution of Ni by Fe, respectively.

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## 1 Introduction

Magnesium hydride has a relatively high hydrogen storage capacity (7.6 wt%), an environmentally friendly nature and low cost, which meets some basic requirements for onboard and stationary applications set by the US DOE. However, the high thermodynamic stability of MgH<sub>2</sub> ( $\Delta H = -75$  kJ mol<sup>-1</sup> H<sub>2</sub>) is a big hurdle for lowering the hydrogen desorption temperature of Mg-based hydrides.<sup>1,2</sup> It has been established that when Mg alloys contain a non-hydride forming element, the value of hydrogenation enthalpy can be decreased. Therefore, alloying is a traditional and effective strategy for altering the thermodynamics of Mg-based alloys for hydrogen storage.

One of the typical examples is Mg<sub>2</sub>Ni, which can react with H<sub>2</sub> to form Mg<sub>2</sub>NiH<sub>4</sub>. It should be noted that the formation enthalpy of Mg<sub>2</sub>NiH<sub>4</sub> ( $\Delta H = -64.5$  kJ mol<sup>-1</sup> H<sub>2</sub>) is lower than that of MgH<sub>2</sub> which allows the former to desorb at lower temperatures than MgH<sub>2</sub>.<sup>3</sup> Morinaga *et al.* found that hydrogen interacted more strongly with Ni atoms rather than Mg atoms in

Mg<sub>2</sub>NiH<sub>4</sub>.<sup>4</sup> This existence of Ni in Mg<sub>2</sub>NiH<sub>4</sub> weakened the Mg–H bond as compared to MgH<sub>2</sub>, and leads to a lower formation enthalpy of Mg<sub>2</sub>NiH<sub>4</sub>. The crystallographic and hydrogen storage properties of Mg<sub>2</sub>NiH<sub>4</sub> were firstly reported by Reilly and Wiswall.<sup>3</sup> Since single phase Mg<sub>2</sub>Ni cannot be simply obtained by casting as phase separation occurs during solidification,<sup>5,6</sup> reactive mechanical alloying (RMA), mechanical milling (MM) and high-energy ball milling (HEBM) have been employed to synthesize Mg<sub>2</sub>Ni or Mg<sub>2</sub>NiH<sub>4</sub>.<sup>7–11</sup> However, it usually takes a long time to prepare Mg<sub>2</sub>Ni or Mg<sub>2</sub>NiH<sub>4</sub> *via* these methods. Therefore, the size of the raw material was reduced to nano-scale to promote the formation of Mg<sub>2</sub>Ni or Mg<sub>2</sub>NiH<sub>4</sub>.<sup>12,13</sup> Shao *et al.* prepared Mg<sub>2</sub>Ni from Mg and Ni nanoparticles produced by hydrogen plasma-metal reaction.<sup>13</sup> It was established that at 553 K and 3 MPa hydrogen pressure, Mg<sub>2</sub>NiH<sub>4</sub> could be generated.

More recently a number of research groups focused on various synthesis methods for doping with additives to further improve the kinetic properties of Mg<sub>2</sub>Ni.<sup>14–17</sup> Xie *et al.* synthesized nanostructured Mg<sub>2</sub>Ni<sub>1-x</sub>Co<sub>x</sub> ( $x = 0.05, 0.1$ ) alloys through hydrogen plasma-metal reaction.<sup>14</sup> The alloys could absorb 2 wt% H<sub>2</sub> at 473 K but desorption still required about 573 K. Hara *et al.* synthesized Mg<sub>2</sub>Ni alloys with the addition of Y and depleted Mg by casting and vacuum heating.<sup>15</sup> The alloys could absorb ~2–3 wt% H<sub>2</sub> at 473 K. Lu *et al.* prepared the alloy Mg<sub>90</sub>In<sub>5</sub>Ni<sub>5</sub> containing the Mg<sub>14</sub>In<sub>3</sub>Ni<sub>3</sub> phase by sintering followed by mechanical milling.<sup>16</sup> The alloy has higher reversible hydrogen storage capacity of 3.3 wt% and the minimum dehydrogenation temperature is reduced to 493 K.

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Iron is considered as a good catalytic/alloying element to improve kinetics/thermodynamics of Mg-based hydrogen storage materials.<sup>18,19</sup> In addition, the hydride of Fe is highly unstable, which may be beneficial for the structural reversibility of Mg<sub>2</sub>Ni in dehydriding.<sup>18</sup> In the present work, the Ni(Fe) nano particles were prepared through an arc plasma method described in previous works.<sup>13,20</sup> The formation of the Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> by using Ni(Fe) precursor and its hydrogen sorption kinetic and thermodynamic properties were carefully investigated. Based on the experimental investigations, the mechanisms of the formation and hydrogen sorption behaviors of the Mg<sub>2</sub>-Ni(Fe)H<sub>4</sub> were proposed.

## 2 Experimental

### 2.1 Sample preparation

In this work, the Ni(Fe) composite nano powder was prepared by an arc plasma evaporation apparatus.<sup>20</sup> The purities of commercially available pure Ni and Fe powders are over 99.5% and the particle size ranges from 50 to 150 μm. The preparation of the powder involves several steps. First, pure Ni and Fe powders were mixed homogeneously with a Ni to Fe weight ratio of 3 : 1. Then the mixture was compressed to cylinders with a diameter of 10 mm and height of 7 mm under a uniaxial pressure of 15 MPa at room temperature. These cylinders were put into the reaction chamber of arc plasma evaporation apparatus filled with mixed gases of 0.01 MPa H<sub>2</sub> and 0.05 MPa Ar. The Ni(Fe) composite nano powder was produced using arc evaporation with the current set at 150 A. After evaporation and condensation, the powder was passivated in mixed argon and air (7 : 1) for 12 h.

About 2.0 g mixture of the commercially available Mg powder and Ni(Fe) composite nano powder in a 2 : 1 molar ratio was ball milled for 4 h in a planetary ball miller. The Mg powder has a purity of 99.5% and its particle size ranges from 50 to 150 μm. The ball milling was carried out in a 50 ml stainless steel vessel under 0.1 MPa Ar atmosphere at 200 rpm with a ball-to-powder weight ratio of 30 : 1. Then these samples were further heated up to 623 K and 673 K in hydrogen under a pressure of 3.7 MPa. After hydrogenation, samples were cooled down to room temperature and the sample container was evacuated to 10<sup>-3</sup> Pa.

### 2.2 Characterization

The compositions of the Ni(Fe) and Mg–Ni(Fe) powders were analyzed by using inductive coupled plasma emission spectrometer (ICP). The phase identifications of as-milled and hydrogenated composite powders were carried out by X-ray diffraction (XRD) using a D/max 2550VL/PCX apparatus equipped with a Cu Kα radiation source. The morphology and microstructure of the powders were characterized by a JEM-2100F transmission electron microscope (TEM) equipped with an energy dispersive spectrometer (EDS) micro-analysis and a backscattered electron detector. The hydrogen sorption isotherms at various temperatures were obtained using a Sievert type pressure–composition–temperature (PCT) volumetric apparatus provided by Shanghai institute of microsystem and information technology. The dehydriding behaviors of

hydrogenated samples were investigated by using differential scanning calorimetry (DSC, Netzsch STA449F3 Jupiter) under 0.1 MPa Ar atmosphere at heating rates of 3, 5 and 10 K min<sup>-1</sup> from room temperature to 773 K.

## 3 Results and discussions

### 3.1 Formation of Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> from Mg and Ni(Fe) powders

Fig. 1 shows the XRD pattern of the as prepared Ni(Fe) powder. These peaks correspond to Ni phase but shift to low angles, suggesting the increased lattice parameter. Based on the XRD patterns, the lattice constant of this phase is determined to be:  $a = 0.3548$  nm, which is higher than that of Ni ( $a = 0.3535$  nm). Such a lattice expansion of Ni can be only attributed to the partial substitution of Ni by Fe. Based on the Scherrer's equation,<sup>21</sup> the average grain size of Ni(Fe) is determined to be 80 nm.

ICP analysis revealed that Ni content in the as prepared Fe(Ni) composite powder is 61.2 wt%, lower than that in the original mixture (75 wt%). According to the Ohno's model, the vapor generation rate of a metal through hydrogen plasma reaction method is proportional to its reaction parameter ( $R_p$ ), which can be expressed by the following equation:<sup>22</sup>

$$R_p = \left( -\frac{\Delta H_r}{L_s} \right) \left[ \frac{N_{H_2}(T)}{N_{H_2}(273)} \right] \quad (1)$$

where  $\Delta H_r$  is the reaction enthalpy between hydrogen and metal,  $L_s$  is the vaporization heat of the metal at temperature  $T$ ,  $N_{H_2}(T)$  and  $N_{H_2}(273)$  refer to the densities of molecular hydrogen in metal at temperature  $T$  and 273 K, respectively. According to eqn (1), the vapor generation rate mainly depends on the hydrogen affinity to metals ( $\Delta H_r$ ) and vaporization heat ( $L_s$ ). The reaction enthalpy between H and Fe is quite similar to the one between H and Ni ( $\sim 20$  kJ mol<sup>-1</sup> H<sub>2</sub>) while the heat of vaporization for Ni (370.4 kJ mol<sup>-1</sup>) is higher than that for Fe (349.6 kJ mol<sup>-1</sup>). Therefore, the evaporation rate of Ni is lower than that of Fe. This is consistent with the ICP result for which the Ni content in the Ni(Fe) powder is lower than that in the mixed powder before arc evaporation.

TEM observations have been done on the as prepared Ni(Fe) composite powder. As seen from Fig. 2a, the shapes of the particles in the Ni(Fe) composite powder are mostly spherical

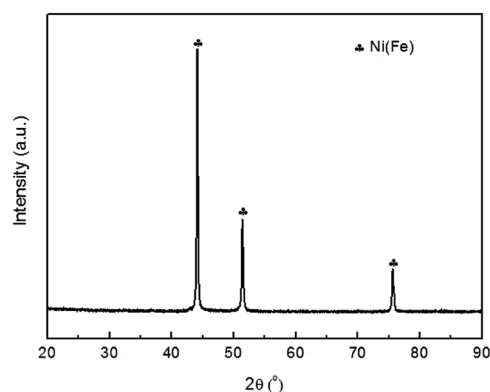


Fig. 1 The X-ray diffraction (XRD) pattern of the Ni(Fe) composite powder prepared using arc plasma method.



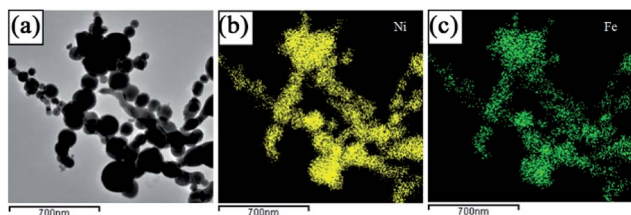


Fig. 2 A bright field TEM micrograph of as prepared Ni(Fe) composite powder (a) and the corresponding EDS elemental distribution maps of Ni (b) and Fe (c).

and the particle size ranges from 50 nm to 260 nm with an average size of about 80 nm. Such morphology and size distribution were also observed in pure Ni powder produced by arc plasma method.<sup>23,24</sup> The corresponding elemental distribution maps of Ni and Fe (Fig. 2b and c) confirm the fairly homogeneous distributions of Ni and Fe in the Ni(Fe) powder.

Furthermore, EDS analyses also reveal that the Ni to Fe weight ratio is 59.2 : 40.8, which is in good agreement with the result obtained by the ICP measurement.

Fig. 3a is the TEM image of the mixture of coarse-grained commercial Mg powder and Ni(Fe) nano powder. It can be seen that there are many smaller particles surrounding those larger ones in this Mg–Ni(Fe) powder. The corresponding SAED pattern (Fig. 3b), with diffraction rings of Mg and Ni(Fe) included, confirms the presence of these phases. The corresponding EDS maps of Mg, Ni and Fe (Fig. 3d–f) confirm the segregations of Ni and Fe elements in the smaller spherical particles and the uniform distribution of Mg element in the bigger irregular ones. Therefore, there are many Ni(Fe) nano particles surrounding coarse-grained commercial Mg particles in the Mg–Ni(Fe) mixture.

Fig. 4 shows the XRD patterns of the Mg–Ni(Fe) samples before and after hydrogenation. From Fig. 4a, the main phases present in the as milled Mg–Ni(Fe) composite powder are Mg and Ni(Fe). The result is consistent with the SAED pattern of the composite powder (Fig. 3b). For the synthesis of  $Mg_2NiH_4$ , the Mg–Fe(Ni) samples were hydrogenated at 623–673 K under 3.7 MPa  $H_2$  for 48 h. The phase components of the Mg–Ni(Fe)–H powders under different conditions were examined using XRD and the results were shown in Fig. 4b and c. Based on the RIR analysis of those XRD patterns, the phase components in the hydrogenated powders were calculated and the results were given in Table 1. After hydrogenation at 623 K under 3.7 MPa  $H_2$  for 48 h, peaks from  $Mg_2NiH_4$  and  $\alpha$ -Fe appear on the XRD pattern of the Mg–Ni(Fe)–H powder (Fig. 4b). Meanwhile, Ni(Fe) content reduces from 57.8 wt% in as milled Mg–Ni(Fe) powder to 2.0 wt% in the hydrogenated one. Therefore, with the presence of Ni(Fe) in the Mg–Ni(Fe) powder,  $Mg_2NiH_4$  is able to be synthesized and then most of Ni(Fe) have been transformed into  $\alpha$ -Fe. As the temperature increases from 623 K to 673 K, the diffraction peaks of  $\beta$ - $MgH_2$  become weaker and Ni(Fe) could hardly be detected on the XRD pattern of Mg–Fe(Ni)–H powder. Indeed, higher temperature favors the diffusion of H, Mg and Ni atoms, thus, after hydrogenation at 673 K under 3.7 MPa  $H_2$  for 48 h, the yield of  $Mg_2NiH_4$  increases to 61.2 wt% in the Mg–Ni(Fe)–H powder.

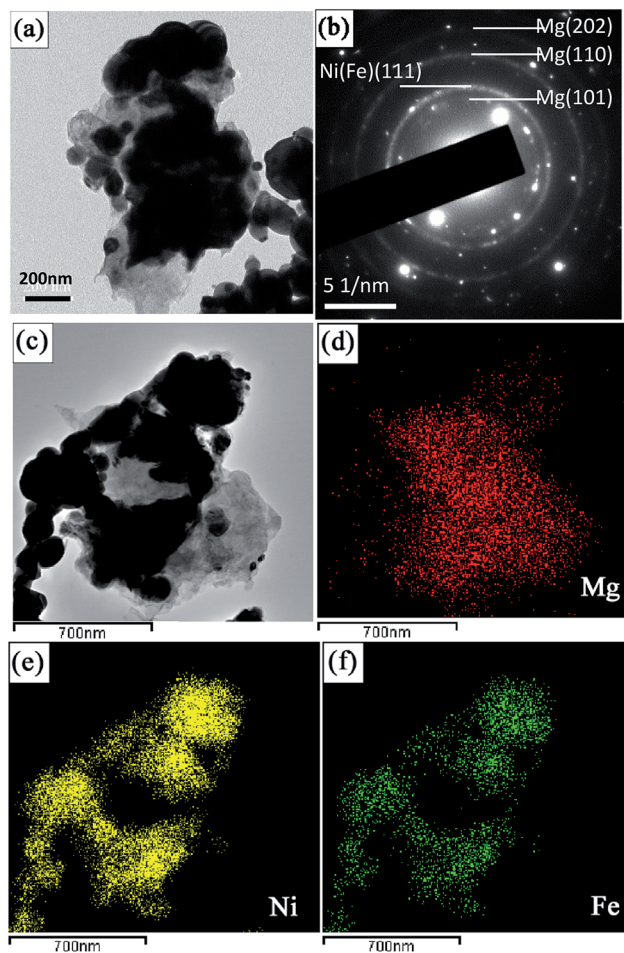


Fig. 3 The bright field TEM micrographs of the Mg–Ni(Fe) mixture (a) and (c), the corresponding selected area electron diffraction (SAED) pattern (b), corresponding EDS maps showing the distributions of Mg (d), Ni (e) and Fe (f).

Fig. 5a presents the typical bright field TEM micrograph of  $Mg_2NiH_4$  in the Mg–Ni(Fe)–H powder prepared at 673 K under a 3.7 MPa hydrogen pressure for 48 h. The size of the  $Mg_2NiH_4$  nano particles is in the range from 100 to 400 nm. The corresponding SAED pattern is given in Fig. 5b, superimposed with

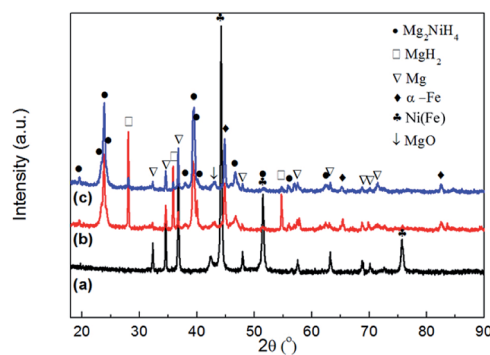


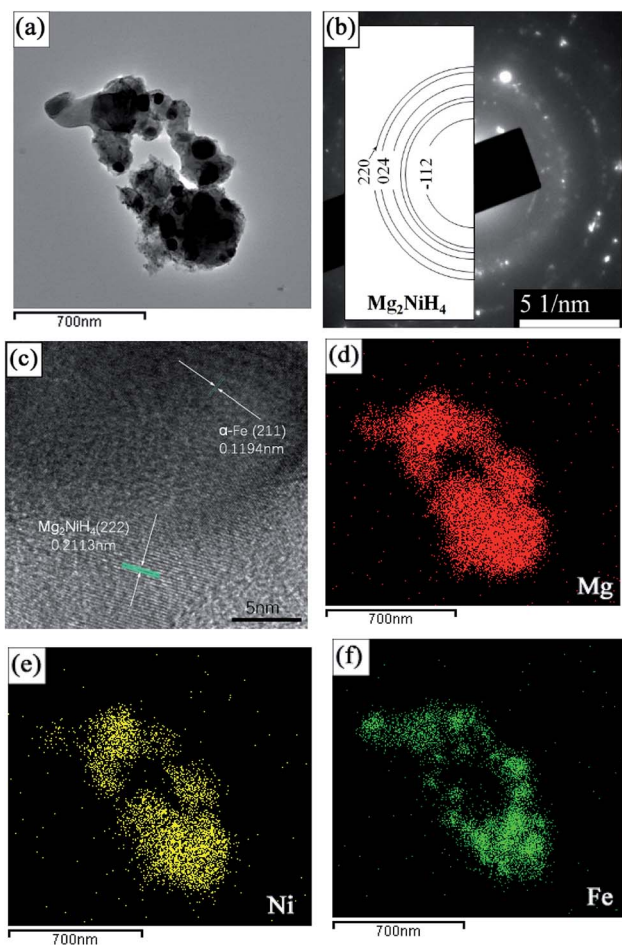
Fig. 4 XRD patterns of as milled Mg–Ni(Fe) powder (a), the Mg–Ni(Fe)–H powder hydrogenated at 623 K under 3.7 MPa  $H_2$  for 48 h (b), the Mg–Ni(Fe)–H powder hydrogenated at 673 K under 3.7 MPa  $H_2$  for 48 h (c).



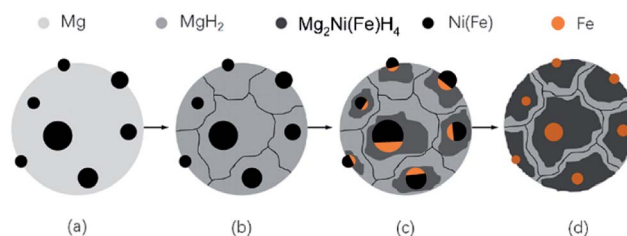
**Table 1** Phase composition (wt%) in as-synthesized samples determined from RIR analysis of XRD data

Sample	Mg–Ni(Fe)	Mg–Ni(Fe)–H (623 K, 3.7 MPa H <sub>2</sub> )	Mg–Ni(Fe)–H (673 K, 3.7 MPa H <sub>2</sub> )
β-MgH <sub>2</sub>	—	27.2	10.2
MgO	—	3.0	3.0
Mg	42.2	10.7	6.8
α-Fe	—	21.3	19.8
Ni(Fe)	57.8	2.0	—
Mg <sub>2</sub> NiH <sub>4</sub>	—	35.8	61.2

the simulated ring patterns of Mg<sub>2</sub>NiH<sub>4</sub>. The majority of the diffraction pattern can be indexed with Mg<sub>2</sub>NiH<sub>4</sub> phase. In addition, the HRTEM observation was further carried out to study the microstructure of nano particles in more detail, as shown in Fig. 5c. It is clearly shown that the bigger bright particles are Mg<sub>2</sub>NiH<sub>4</sub> and the smaller dark particles are α-Fe, according to the measurements of inter-planar spacing. The corresponding EDS maps of Mg, Ni and Fe (Fig. 6d–f) confirm



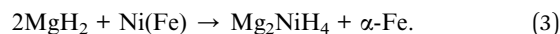
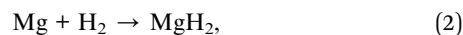
**Fig. 5** A bright field TEM micrograph of the Mg–Ni(Fe)–H powder prepared at 673 K under a 3.7 MPa hydrogen atmosphere for 48 h (a), the corresponding SAED pattern (b), the corresponding high-resolution TEM (HRTEM) image (c) of the selected zone in (a), and corresponding EDS maps showing the distribution of Mg (d), Ni (e) and Fe (f).



**Fig. 6** Schematic illustrations showing the formation procedures of the Mg<sub>2</sub>Fe(Ni)H<sub>6</sub> from coarse-grained Mg powder and γ-Fe(Ni) nano particles.

the segregation of Fe element in the smaller spherical particles and the uniform distributions of Ni and Mg elements in the bigger irregular ones. Therefore, after hydrogenation at 673 K under a 3.7 MPa hydrogen pressure for 48 h, Ni atoms diffused from Ni(Fe) nano particles to the coarse-grained Mg particles to form Mg<sub>2</sub>NiH<sub>4</sub> and then a portion of Fe atoms in the Ni(Fe) nano particles transformed into α-Fe nano particles with smaller particle size. The shapes of the α-Fe nano particles are mostly spherical and the particle size ranges from 40 nm to 120 nm with an average size of about 60 nm. In addition, a portion of Fe atoms have also diffused with Ni atoms to form Mg<sub>2</sub>Ni(Fe)H<sub>4</sub>, as confined by the EDS analyses.

During hydrogenation, two reactions occur in the Mg–Ni(Fe) powder:



The formation of Mg<sub>2</sub>NiH<sub>4</sub> requires a long-distance diffusion of not only small H atoms but also the large metallic species Mg and Ni. Owing to the low diffusion rates of Mg and Ni, the formation of the Mg<sub>2</sub>NiH<sub>4</sub> generally suffers from poor kinetics. In systems based on Mg and Ni that are tailored mainly toward fast sorption kinetics, MgH<sub>2</sub> would rather form prior to Mg<sub>2</sub>NiH<sub>4</sub>.<sup>13,25</sup> Fig. 6 shows a sketch illustrating the formation of Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> nano particles. Firstly, the mixed Mg powders in micron scale and Ni(Fe) nano particles are heated in hydrogen, as shown in Fig. 6a. At the beginning of the absorption, because of the catalytic effect from nickel and iron on the dissociation of gaseous hydrogen,<sup>26</sup> MgH<sub>2</sub> will form through hydrogen spill over mechanism while Ni(Fe) particles still exists, as seen in Fig. 6b. Then Mg<sub>2</sub>NiH<sub>4</sub> will nucleate at the phase boundaries between MgH<sub>2</sub> and Ni(Fe) (Fig. 6c). Due to the partial substitution of Ni by Fe in Ni(Fe) lattice, a portion of Fe atoms have dissolved into the Mg<sub>2</sub>NiH<sub>4</sub> to form Mg<sub>2</sub>Ni(Fe)H<sub>4</sub>. Consequently, when most of MgH<sub>2</sub> transformed into Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> in the Mg–Ni(Fe)–H powder, Fe in the Ni(Fe) particles transformed into α-Fe nano particles with smaller grain sizes surrounding the Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> particles (see Fig. 6d).

### 3.2 Hydrogen sorption behaviors of the Mg–Ni(Fe) powder

Fig. 7a shows the PCT curves of the hydrogen absorption–desorption processes for the Mg–Ni(Fe) samples at 598, 623, 648 and 673 K. The data obtained from PCT curves are summarized in Table 2. During H<sub>2</sub> absorption, two plateaus are clearly visible



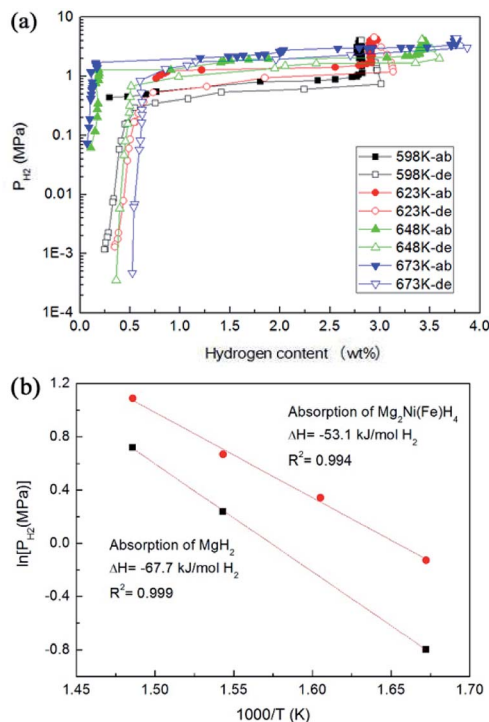


Fig. 7 PCT curves at various temperatures of the Mg–Ni(Fe) sample (a), the corresponding van't Hoff plots for MgH<sub>2</sub> and Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> in the Mg–Ni(Fe)–H sample (b).

on each profile. According to the XRD results given above, two types of hydrides, MgH<sub>2</sub> and Mg<sub>2</sub>Ni(Fe)H<sub>4</sub>, formed after hydrogenation. It is worth noting that Mg<sub>2</sub>NiH<sub>4</sub> has higher equilibrium pressure than MgH<sub>2</sub> due to its lower thermodynamic stability.<sup>29,30</sup> The maximum hydrogen absorption capacities at 598, 623, 648 and 673 K are 3.0, 3.1, 3.6 and 3.9 wt%, respectively. It is noticed that the capacities at 598 and 623 K are lower than the theoretical value of 3.6 wt% for Mg<sub>2</sub>Ni. This is due to the existence of MgO, the residual Mg in the hydride Mg–Ni(Fe) composites, as shown in Fig. 4b. However, when the hydrogenated temperature is above 648 K, the capacity is higher than 3.6 wt%. Since the original molar ratio of Mg and Ni(Fe) in the Mg–Ni(Fe) is 2 : 1 and the ICP analysis revealed that Ni content in the Fe(Ni) powder is 61.2 wt%, the molar ratio of Mg to Ni in the Mg–Ni(Fe) is actually 3.3 : 1, higher than 2 : 1. Therefore, after complete hydrogenation, the maximum hydrogen absorption capacities of the Mg–Ni(Fe) can be higher than 3.6 wt%.

The van't Hoff plot ( $\ln P$  versus  $1/T$ ) is used to estimate the thermal stability of hydrides, as shown in Fig. 7b. Table 3

Table 2 Data obtained from pressure–composition isotherms of the Mg–Ni(Fe) sample at various temperatures

Temperature (K)	Low plateau of absorption (MPa)	High plateau of absorption (MPa)	Maximum hydrogen content (wt%)
598	0.45	0.88	3.0
623	—	1.41	3.1
648	1.27	1.95	3.6
673	2.05	2.97	3.9

presents the value of the formation enthalpy ( $\Delta H$ ) for the Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> in this work and the ones for the Mg<sub>2</sub>NiH<sub>4</sub> reported in literature. According to the linear fittings of  $\ln P$  versus  $1000/T$  in Fig. 8b, the van't Hoff equations for the hydrogenation are  $\ln(P_{\text{low}}) = -6.392 \times 10^3/T + 10.57$  for MgH<sub>2</sub> and  $\ln(P_{\text{high}}) = -8.138 \times 10^3/T + 12.81$  for Mg<sub>2</sub>Ni(Fe)H<sub>4</sub>. The hydrogenation enthalpies ( $\Delta H_{\text{ab}}$ ) for MgH<sub>2</sub> and Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> are therefore calculated to be  $-67.7$  and  $-53.1$  kJ mol<sup>-1</sup> H<sub>2</sub>, respectively. In contrast, Pourabdoli *et al.* reported that the integral heat of H<sub>2</sub> desorption for the MgH<sub>2</sub>-10 wt% (9Ni-2Mg-Y) nano-composite was about 78 kJ mol<sup>-1</sup> H<sub>2</sub> measured by using adsorption microcalorimetry.<sup>31</sup> Therefore, the formation enthalpy of Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> is obviously higher than those reported in literature. Van Setten *et al.* have studied the influence of transition metals (Cu, Fe and Co) on the structural and hydrogen sorption properties of Mg<sub>2</sub>NiH<sub>4</sub> using first-principle based calculations.<sup>18</sup> Doping with Co or Cu leads to octahedron is distorted. The hydrogen tetrahedra around such Ni atoms are distorted with Ni–H distances from 1.51 to 1.80 Å, whereas in undoped Mg<sub>2</sub>NiH<sub>4</sub> they are between 1.56 and 1.58 Å. Therefore, the highest hydriding enthalpy,  $-55.5$  kJ mol<sup>-1</sup> H<sub>2</sub>, is found for the Fe doped Mg<sub>2</sub>NiH<sub>4</sub>.<sup>18</sup> This value is higher than the simulated value for Mg<sub>2</sub>NiH<sub>4</sub> ( $-63.5$  kJ mol<sup>-1</sup> H<sub>2</sub>). Therefore, the results present in this work also confirm that doping with Fe would remarkably destabilize Mg<sub>2</sub>NiH<sub>4</sub> through increasing the formation enthalpy. In addition, the thermodynamics of marginally stable compounds, whereas doping with Fe leads to an unstable compound. In the Fe doped Mg<sub>2</sub>NiH<sub>4</sub>, the hydrogenation of MgH<sub>2</sub> in the Mg–Ni(Fe) is also slightly improved by the addition of Ni(Fe) nanoparticles, which is in accordance with recent experimental results and theoretical calculations. For instance, the hydriding enthalpies of Mg–Ni nanocomposite coprecipitated from solution and MgH<sub>2</sub>–Ni/Ti prepared by ball milling were reduced to  $-70.0$  and  $-67.8$  kJ mol<sup>-1</sup> H<sub>2</sub>.<sup>32,33</sup> Dai and Shevlin carried out theoretical calculations and found that the significant electronic density donation from the H<sup>-</sup> ions to an empty d-state of the Ni dopant was able to reduce the H<sup>-</sup> anion charge to weaken the hydrogen bonding.<sup>34,35</sup> Thus, doping with Ni would also remarkably destabilize MgH<sub>2</sub> to improve the thermodynamics of MgH<sub>2</sub>.

Fig. 8a shows DSC curves obtained by heating the Mg–Ni(Fe)–H powder up to 723 K at heating rates of 3, 5 and 10 K min<sup>-1</sup> under Ar. These curves suggest a two-step desorption behavior by showing two endothermic peaks. The first one is a strong and broad endothermic peak in the lower temperature range, while the second one is a relatively weaker peak in the higher temperature range. Clearly, the strong peak corresponds well to the decomposition of Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> while the weak one results from the decomposition of the MgH<sub>2</sub>. The peak dehydrogenation temperatures of the Mg<sub>2</sub>Ni(Fe)H<sub>4</sub> in the Mg–Ni(Fe)–H powder at heating rates of 3, 5 and 10 K min<sup>-1</sup> are 501.3, 510.6 and 526.1 K, respectively. Zou *et al.* prepared the Mg-rich Mg–Ni ultrafine particles through the arc plasma method.<sup>27</sup> DSC analyses showed that there are three endothermic peaks appeared at 513, 643 and 668 K, which correspond to the phase transformation of Mg<sub>2</sub>NiH<sub>4</sub> from its low temperature form to the high temperature form, the dehydrogenation of Mg<sub>2</sub>NiH<sub>4</sub> and MgH<sub>2</sub>, respectively. Therefore, the



**Table 3** The formation enthalpy value of  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  in the Mg–Ni(Fe)–H sample and the ones of  $\text{Mg}_2\text{NiH}_4$  reported in the literatures<sup>25,27–29</sup>

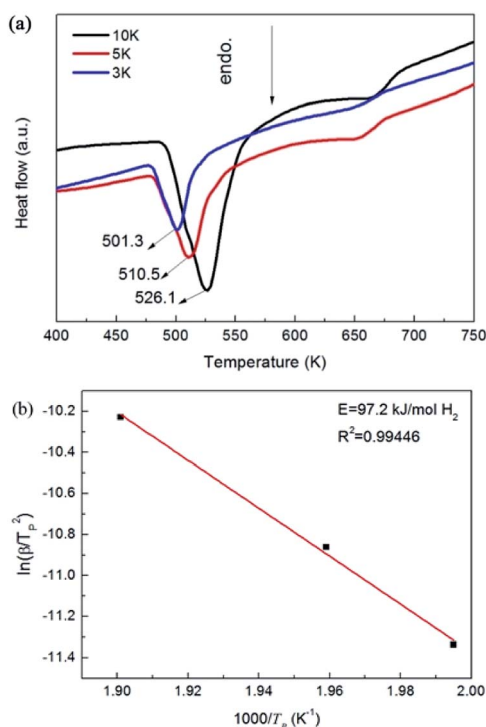
Formation ( $\Delta H$ ) ( $\text{kJ mol}^{-1} \text{H}_2$ )	Reference
–53.1	This work
–67.5	25
–64.6	27
–66.3	28
–63.2	29

peak dehydrogenation temperature of the  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  in the Mg–Fe(Ni)–H powder is 117 K lower than that of the  $\text{Mg}_2\text{NiH}_4$  in the Mg-rich Mg–Ni ultrafine particles. In addition, it is noteworthy that the peak dehydrogenation temperatures of the  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  in the Mg–Ni(Fe)–H powder is indeed in the temperature range of the phase transformation of  $\text{Mg}_2\text{NiH}_4$ .<sup>36,37</sup> Thus, doping with Fe would improve the hydrogen desorption properties of  $\text{Mg}_2\text{NiH}_4$  through promoting the decomposition of the low temperature form of  $\text{Mg}_2\text{NiH}_4$ .

To further study the desorption behavior of the Mg–Ni(Fe)–H powder, the dehydrogenation activation energy ( $E_{\text{de}}$ ) was calculated by using Kissinger's method:<sup>16,20</sup>

$$\frac{d \ln \left( \frac{\beta}{T_p^2} \right)}{d \frac{1}{T_p}} = -\frac{E_{\text{de}}}{R} \quad (4)$$

where  $\beta$  is the heating rate,  $T_p$  is the peak temperature, and  $R$  is the gas constant. Just as expected, the peaks of dehydrogenation shift



**Fig. 8** DSC profiles of the Mg–Ni(Fe)–H sample measured at different heating rates (a) and the corresponding  $\ln(\beta/T_p^2)$ – $1000/T_p$  plot (b).

toward higher temperature as the heating rate increases. According to eqn (4), the  $E_{\text{de}}$  value of the  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  in the Mg–Ni(Fe)–H is estimated to be  $97.2 \text{ kJ mol}^{-1} \text{H}_2$ . It is obviously lower than those reported by Hur and Cermak,<sup>38,39</sup> which are 109 and  $113 \text{ kJ mol}^{-1} \text{H}_2$ , respectively. Giusepponi *et al.* found that the higher coordination of iron compared to magnesium in the hydride was able to destabilize the crystalline structure and raise the probability of the H diffusion toward the interface.<sup>40</sup> Recent research works have clearly shown that Fe nanoparticles play the role of catalyst for the hydrogen absorption–desorption in Mg. Polanski *et al.* have reported that the  $2\text{Mg} + \text{Fe}$  nanocomposite can absorb 1.1 wt% hydrogen at 303 K due to the catalytic effect of nano Fe particles on the hydrogen absorption in Mg.<sup>41</sup> Bassetti *et al.* have prepared  $\text{MgH}_2$ –Fe nanocomposites, which can release about 5 wt% hydrogen in about 600 s at 573 K due also to the catalytic effect from Fe.<sup>42</sup> Therefore, a portion of Fe in the Ni(Fe) nano particles transformed into  $\alpha$ -Fe nano particles in the Mg–Ni(Fe)–H, which also improved the dehydrogenation kinetics of  $\text{Mg}_2\text{NiH}_4$ .

## 4 Conclusions

In the present work, the formation of  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  in the mixed precursors of the coarse-grained Mg powder and the Ni(Fe) nano powder prepared through arc plasma method was investigated. The microstructure, composition, phase components and hydrogen storage properties of the Mg–Ni(Fe) powder were characterized and the main results are as follows.

(1) After sintering of the Mg–Ni(Fe) powder at 673 K under 3.7 MPa  $\text{H}_2$  for 48 h, the yield of  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  reached 61.2 wt% and a portion of Fe atoms in the Ni(Fe) nano particles transformed into  $\alpha$ -Fe nano particles.

(2)  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  shows lower hydrogen desorption temperature and better desorption kinetics when compared to the  $\text{Mg}_2\text{NiH}_4$  obtained from arc plasma evaporated Mg–Ni powder. The peak dehydrogenation temperature of  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  is reduced to 501.3 K and the dehydrogenation activation energy is reduced to  $97.2 \text{ kJ mol}^{-1} \text{H}_2$ .

(3) The formation enthalpy of  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  is determined to be  $-53.1 \text{ kJ mol}^{-1} \text{H}_2$ , higher than the values reported in literature for  $\text{Mg}_2\text{NiH}_4$ . The improvements in hydrogen sorption kinetics and thermodynamics can be attributed to the catalytic effect of  $\alpha$ -Fe nano particles surrounding the  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$  and the destabilization effect caused by Fe substitution for Ni in the  $\text{Mg}_2\text{Ni}(\text{Fe})\text{H}_4$ , respectively.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 F. Cheng, Z. Tao, J. Liang and J. Chen, *Chem. Commun.*, 2012, **48**, 7334.
- 2 H. Shao, G. Xin, J. Zheng and X. Li, *Nano Energy*, 2012, **1**, 590.
- 3 J. J. Reilly and R. H. Wiswall, *Inorg. Chem.*, 1968, **7**, 2254.
- 4 M. Morinaga and H. Yukawa, *Mater. Sci. Eng., A*, 2002, **329**, 268.
- 5 T. B. Massalski, *Binary Alloy Phase Diagrams*, American Society for Metals, 1986.
- 6 A. A. Nayeb-Hashemi and J. B. Clark, *J. Phase Equilib.*, 1985, **6**, 238.
- 7 C. C. Koch, *Nanostruct. Mater.*, 1997, **9**, 13.
- 8 J. Huot, H. Enoki and E. Akiba, *J. Alloys Compd.*, 2008, **453**, 203.
- 9 A. K. Singh and O. N. Srivastava, *J. Alloys Compd.*, 1995, **227**, 63.
- 10 L. Zaluski, A. Zaluska and J. O. Ström-Olsen, *J. Alloys Compd.*, 1995, **217**, 245.
- 11 R. A. Varin and T. Czujko, *Mater. Manuf. Processes*, 2002, **17**, 129.
- 12 M. D. Hampton, J. K. Lomness and L. A. Giannuzzi, *Int. J. Hydrogen Energy*, 2002, **27**, 79.
- 13 H. Shao, T. Liu, X. Li and L. Zhang, *Scr. Mater.*, 2003, **49**, 595.
- 14 L. Xie, H. Shao, Y. Wang, Y. Li and X. Li, *Int. J. Hydrogen Energy*, 2007, **32**, 1949.
- 15 M. Hara, S. Morozumi and K. Watanabe, *J. Alloys Compd.*, 2006, **414**, 207.
- 16 Y. Lu, H. Wang, J. Liu, L. Z. Ouyang, L. Zhu, D. Zhang and M. Zhu, *J. Phys. Chem. C*, 2015, **119**, 26858.
- 17 L. Z. Ouyang, Z. J. Cao, H. Wang, J. W. Liu, D. L. Sun, Q. A. Zhang and M. Zhu, *Int. J. Hydrogen Energy*, 2013, **38**, 8881.
- 18 M. J. VanSetten and G. A. Wijs, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2007, **76**, 075125.
- 19 S. A. Shevlin and Z. X. Guo, *J. Phys. Chem. C*, 2013, **117**, 10883.
- 20 J. X. Zou, X. Q. Zeng, Y. J. Ying, X. Chen, H. Guo, S. Zhou and W. J. Ding, *Int. J. Hydrogen Energy*, 2013, **38**, 2337.
- 21 A. L. Patterson, *X-ray and Neutron Diffraction*, 1966, p. 243.
- 22 S. Ohno and M. Uda, *J. Jpn. Inst. Met.*, 1984, **48**, 640.
- 23 Z. L. Cui, L. F. Dong and C. C. Hao, *Adv. Mater. Res.*, 2013, **734**, 1555.
- 24 Z. Cui, Z. Zhang, C. Hao, L. Dong, Z. Meng and L. Yu, *Thin Solid Films*, 1998, **318**, 76.
- 25 H. Y. Shao, H. Xu, Y. Wang and X. G. Li, *Nanotechnology*, 2004, **15**, 269.
- 26 G. Liang, J. Huot, S. Boily, A. VanNeste and R. Schukz, *J. Alloys Compd.*, 1999, **292**, 247.
- 27 J. X. Zou, H. Q. Sun, X. Q. Zeng, G. Ji and W. J. Ding, *J. Nanomater.*, 2012, 592147.
- 28 E. Rönnebro, J. Jensen, D. Noréus and N. Bjerrum, *J. Alloys Compd.*, 1999, **293**, 146.
- 29 K. Nomura, E. Akiba and S. Ono, *Int. J. Hydrogen Energy*, 1981, **6**, 295.
- 30 G. Liang, S. Boily, J. Huot, A. Neste and R. Schulz, *J. Alloys Compd.*, 1998, **268**, 302.
- 31 M. Pourabdoli, S. Raygan, H. Abdizadeh and D. Uner, *Int. J. Hydrogen Energy*, 2013, **38**, 11910.
- 32 Y. N. Liu, J. X. Zou, X. Q. Zeng, X. M. Wu, D. J. Li and W. J. Ding, *J. Phys. Chem. C*, 2014, **118**, 18401.
- 33 H. B. Lu, C. K. Poh, L. C. Zhang, Z. P. Guo, X. B. Yu and H. K. Liu, *J. Alloys Compd.*, 2009, **481**, 152.
- 34 J. H. Dai, Y. Song and R. Yang, *J. Phys. Chem. C*, 2010, **114**, 11328.
- 35 S. A. Shevlin and Z. X. Guo, *J. Phys. Chem. C*, 2013, **117**, 10883.
- 36 M. L. Post and J. J. Murray, *J. Less-Common Met.*, 1987, **134**, 15.
- 37 L. Q. Li, T. Akiyama and J. Yagi, *Intermetallics*, 1999, **7**, 671.
- 38 T. H. Hur, J. S. Han, J. H. Kim and B. K. Kim, *J. Nanosci. Nanotechnol.*, 2011, **11**, 6499.
- 39 J. Cermak and L. Kral, *Defect Diffus. Forum*, 2009, **283**, 639.
- 40 S. Giusepponi and M. Celino, *Int. J. Hydrogen Energy*, 2013, **38**, 15254.
- 41 M. Polanski, J. Bystrzycki, R. A. Varin and T. Plocinski, *Int. J. Hydrogen Energy*, 2001, **36**, 1059.
- 42 A. Bassetti, E. Bonetti, L. Pasquini, A. Montone, J. Grbovic and M. Vittori Antisari, *Eur. Phys. J. B*, 2005, **43**, 19.

