Study of Structural, Thermodynamic and Electrochemical Properties of MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75-x}$Fe$_x$ (x = 0 and 0.75) Compounds

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Abstract
The structural, thermodynamic and electrochemical properties of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ compound, used as negative electrode of Ni – MH accumulator, are studied and compared to those of MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ electrode. The thermodynamic results show that the total substitution of cobalt by iron leads to a decrease solid – gas capacity measured at 25°C from 5.5 H/mol to 3.93 H/mol for MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ compounds, respectively. The electrochemical discharge capacity for the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ compound reaches its maximum of 270 mAh/g after 12 cycles and then decreases to 200 mAh/g after 25 cycles. However, the electrochemical discharge capacity of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ compound reaches its maximum value of 200 mAh/g after 9 cycles and then decreases to 180 mAh/g after 25 cycles. The decrease of the discharge capacity for the two alloys is attributed to the corrosion and deprecipitation of the alloys in the KOH aqueous solution. The value of the hydrogen diffusion coefficient $D_H$ determined by the cyclic voltammetry are equal to $5.85 \times 10^{-10}$ cm$^2$ s$^{-1}$ and $3.96 \times 10^{-10}$ cm$^2$ s$^{-1}$ for MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ compounds, respectively. The hydrogen diffusion coefficient determined by electrochemical impedance spectroscopy (EIS), corresponding to 10 and 100 % of the charge state are equal to $6.64 \times 10^{-10}$ cm$^2$ s$^{-1}$ (α phase) and $1.6 \times 10^{-10}$ cm$^2$ s$^{-1}$ (β phase) for the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ alloy. However, for the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$, the hydrogen diffusion coefficient are equal to $7.27 \times 10^{-10}$ cm$^2$ s$^{-1}$ (α phase) and $2.51 \times 10^{-10}$ cm$^2$ s$^{-1}$ (β phase).

1. Introduction
In recent years, hydrogen may constitute the main energy vector in the future. There is multiple ways of storing hydrogen. An alternative to the use of gaseous pressure tanks is to store hydrogen in metal hydrides. Many of applications of hydrogen storage alloys have been widely used as the negative electrode materials in the nickel – metal hydride (Ni / MH) secondary batteries for their higher energy density, high capacity rate and longer life cycle. The high charge / discharge rate of Ni–MH batteries is gaining increasing attention.
The rapid charging performance of the cell is very important for some applications such as mobile phones and electric vehicles. The key to the improvement of the commercial Ni/MH batteries is the improvement of the electrochemical properties of the hydrogen storage alloys including the electrochemical capacity, the life cycle and the high rate of dischargeability.

Most of the negative electrodes of such batteries are made of LaNi$_5$-type alloy [1, 2]. The behavior of these batteries can be changed when modifying the hydrogen storage alloy to achieve a good performance. The LaNi$_5$ compound, one of the AB$_5$ - rare type– earth alloys, is the most promising candidate [3, 4]. Nevertheless, it exhibits a high equilibrium pressure of 1.7 bars at room temperature and its electrochemical capacity decreases rapidly when increasing the charge – discharge cycles. This such latte phenomenon was attributed to the oxidation and the formation of the La(OH)$_3$ film at the surface [5]. To decrease the plateau pressure and inhibit the oxidation phenomenon, some works have been carried out to replace the lanthanum by mischmetal (a mixture of different rare earth such as: Ce, Nd and Pr) and a part of nickel atoms with different transition elements such as Mn, Al and Co [6, 7-10].

These commercial AB$_5$ alloys are usually of the type Mm(Ni,Co,Al,Mn)$_5$, containing typically 10 wt% Co. Cobalt is present in the alloys to guarantee the long cycle life of the negative electrode [11]. However, cobalt is also the most expensive metal among the alloying compounds representing 10 % of the weight but 45 % of the total alloy price. Therefore, it is interesting to investigate low-Co or Co-free AB$_5$-type alloys. For this purpose we have studied the influence of the Fe for Co substitution, especially on the life time of the electrode.

In the present work, we have studied the effect of total substitution of Co by Fe on the structural, thermodynamic and electrochemical properties of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ compound. The value of the hydrogen diffusion coefficient DH was determined by the cyclic voltammetry and electrochemical impedance spectroscopy (EIS) have been evaluated the reversibility and the kinetic of the electrochemical reaction of the both alloys.

2. Experimental Detail

The MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ and MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ alloys were prepared by UHF induction melting (UHF) of the pure elements followed by an appropriate annealing to ensure a good homogeneity.

They are carefully checked by metallographic examination, electron probe microanalysis (EPMA) and powder X-ray diffraction (XRD) with CuK$_\alpha$ beam.

The pressure-composition -isotherm (P-C-T) was measured using a volumetric (Sievert) method. Each sample was first activated by several hydrating -dehydrtating cycles to reduce the grain size and to increase the kinetic.

2.1. Preparation of the M H Electrode and Ni – MH Cell

These ingot alloys were ground mechanically and saved to less than 63µm in a glove box under an argon atmosphere. The “latex” technology has been used for the electrode preparation. Ninety percent of the alloy powder was mixed with 5% of black carbon (to obtain a good conductivity) and 5% of polytetrafluoroethylen (PTFE). Two pieces of 0.5 cm$^2$ of this latex have been pressed on each side of a nickel grid to prevent the electrode platform breaking into pieces during the charge–discharge cycling [12]. This assembly forms the negative electrode of Ni–MH battery. The counter electrode was formed by the Ni ox hydroxide-Ni(OH)$_2$, whereas the reference electrode was the Hg/HgO 1M KOH electrode. The electrolyte was 1M KOH solution, prepared with deionized water.

2.2. Electrochemical Measurements

All the electrochemical measurements were performed at room temperature in a conventional three electrode-open-air cell using a computer- as a controlled response frequency analyser coupled directly to a VMP Biologic potentiostat–galvanostat. The discharge capacity of the electrode was determined by a galvanostatic charging-discharging, respectively, at C/3 and D/6 regime. Every cycle was carried out by charging fully at 120 mA/g for 3 h (this time majored by 50 % due to the efficiency of the charging reaction) and discharging at 60 mA/h/g at room temperature. After activating the electrode for 35 cycles, the cyclic voltammetry was applied at scan rates of 10, 20, 30, 40 and 50 µV s$^{-1}$ between −1.1 and −0.5V versus Hg/HgO. Impedance spectroscopy measurements were realized after 13 cycles. The chronoamperometry was applied after 12 charge-discharge cycles. This method consists of fully charging and discharging the electrode at a constant potential (- 0.6 V versus Hg/HgO).

The electrode was charged to a state of charge (SOC) of 10% at C/3 regime. After that, a rest of 3 h was applied to the open-circuit potential to get stabilized. The electrochemical impedance spectrum of the electrode was measured by the AC perturbation of ± 5 mV as amplitude and 1 mHz to 50 KHz frequency range for 10 points per decade.

3. Resultats and Discussions

3.1. Structural Properties

EPMA analysis indicates that the composition of these compounds MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ and MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ are a single phase and has a homogenous composition in agreement with the nominal one. The diffraction patterns were refined with a Rietveld analysis using the program fullproff represented in Fig. 1 and Fig. 2. The structural characterisation shows that these compounds are indexed in CaCu5 hexagonal crystalline structure (P6/mmm space group). The results of the metallographic
examination, the electron probe microanalysis (EPMA) and the structure characterisation by X-ray diffraction (XRD) of these compounds are given in Table 1. In fact, the lattice parameters and lattice volume increases progressively by the totaly substitution of iron for cobalt, which indicate that the capacities of the as-quenched alloys decrease greatly with the increase of the Fe content [13].

Figure 1. Refined diffraction pattern of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$.

Figure 2. Refined diffraction pattern of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ [25].

Table 1. Lattice parameters and lattice volume of the substituted compound.

<table>
<thead>
<tr>
<th>Compound</th>
<th>a (Å)</th>
<th>c (Å)</th>
<th>V (Å$^3$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>LaNi$_5$</td>
<td>5.012</td>
<td>3.984</td>
<td>86.68</td>
</tr>
<tr>
<td>MmNi$<em>{3.55}$Mn$</em>{0.4}$Al$<em>{0.3}$Co$</em>{0.75}$ [24]</td>
<td>5.019</td>
<td>4.049</td>
<td>88.32</td>
</tr>
<tr>
<td>MmNi$<em>{3.55}$Mn$</em>{0.4}$Al$<em>{0.3}$Fe$</em>{0.75}$ [25, 26]</td>
<td>5.0422</td>
<td>4.067</td>
<td>89.54</td>
</tr>
</tbody>
</table>

3.2. Thermodynamic Properties

The pressure – composition – isotherm (P-C-T) with MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ measured at 25, 40 and 80°C is represented in Fig. 3 and Fig. 4,
where the hydrogen desorption pressure is plotted as a function of an amount of desorbed hydrogen. These values of solid-gas capacity and pressure equilibrium at 25, 40 and 80°C are given in Table 2. The solid-gas capacity measured at 25°C from 5.5 H/mol to 3.93 H/mol for MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ compounds, respectively. The result shows that the substitution of cobalt by iron increases the stability of the compound.

Figure 3. Isotherm of desorption at 25, 40 and 80 °C of compound MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$.  

![Figure 3](image1.png)

The pressure equilibrium decrease from 0.065 to 0.024 bars for MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ compounds, respectively, shows that the substitution of cobalt by iron increases the stability of the compound and can be used as negative electrodes in NiMH batteries or for the hydrogen storage. Referring to the cell volumes listed in Table 1, it can be found that a large cell volume corresponds to a low plateau pressure. It is believed that the expansion of the cell...
volume leads to a decrease of the equilibrium pressure. For both compounds the capacity decreases slightly when the temperature increases. So, for MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Co\textsubscript{0.75} compound, the capacity is of 5.5 H/moles at 25°C, and decreases to reach the value of 4.96 H/moles at 80°C. While for the MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Fe\textsubscript{0.75} the capacity is of 3.93 H/mole at 25°C, and decreases to reach the value of 3.4 H/mole at 80°C.

The enthalpy and entropy of these hydrides formation, calculated using the equilibrium pressure at 25, 40 and 80°C, are given in Table 3. It is obvious from the data that the absolute value of the alloy enthalpy increases when the cobalt is substituted by iron. This leads to an improved stability of the accordingly prepared hydrides. These results show also that the value of enthalpy formation of the hydride is negative. So, it can be concluded that the reaction of the hyduration is exothermic.

Table 2. Solid-gas capacity and pressure equilibrium at 25, 40 and 80°C temperature of MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Co\textsubscript{0.75} and MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Fe\textsubscript{0.75} compounds.

<table>
<thead>
<tr>
<th>P(bar)</th>
<th>Capacity (H/mole)</th>
<th>P(bar)</th>
<th>Capacity (H/mole)</th>
<th>P(bar)</th>
<th>Capacity (H/mole)</th>
</tr>
</thead>
<tbody>
<tr>
<td>25°C</td>
<td>0.065</td>
<td>5.50</td>
<td>0.138</td>
<td>5.15</td>
<td>0.82</td>
</tr>
<tr>
<td>40°C</td>
<td>0.024</td>
<td>3.93</td>
<td>0.085</td>
<td>3.74</td>
<td>0.582</td>
</tr>
</tbody>
</table>

Table 3. The enthalpy and entropy of the formation of these hydrides.

<table>
<thead>
<tr>
<th>Compounds</th>
<th>(\Delta H) (Kcal/mole H\textsubscript{2})</th>
<th>(\Delta S) (cal/K/mole H\textsubscript{2})</th>
</tr>
</thead>
<tbody>
<tr>
<td>MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Co\textsubscript{0.75}</td>
<td>-9.72</td>
<td>27.09</td>
</tr>
<tr>
<td>MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Fe\textsubscript{0.75}</td>
<td>-10.38</td>
<td>28.53</td>
</tr>
</tbody>
</table>

3.3. Electrochemical Properties

3.3.1. Life Time of the Electrode

Fig. 5 shows the variation of the discharge capacity of the MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Fe\textsubscript{0.75} and the MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Co\textsubscript{0.75} electrodes as a function of the number of cycles. The discharge capacity of the MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Co\textsubscript{0.75} alloy reaches a maximum value of 270 mAh/g after 12 cycles, decreases to about 200 mAh/g after 25 cycles and remains constant after that. However, the discharge capacity of the MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Fe\textsubscript{0.75} Compound reaches a maximum value of 200 mAh/g after 9 cycles, decreases to about 180 mAh/g after 25 cycles and then remains constant. The capacity decrease is mainly due to the surface layer oxidation and corrosion phenomena. Pan et al. [14] and Geng et al. [15] assume that the loss in discharge capacity is due to the deterioration of the negative electrode material with increasing the number of charge-discharge cycles. In fact, during the cycling, the rare earth elements, such as La or the transition metal such as Mn segregate, to the grain boundaries, where they were subject to corrosion. The corrosion is disposed on the surface of the grain particles as a needle-shaped La(OH)$_3$ or as Mn$_3$O$_4$. This decrease may be also attributed to the oxidation of iron on the alloy surface which limits the hydrogen transfer from the surface to the bulk of the alloy and consequently to a loss in the amount of hydrogen stored in the material. The cobalt-containing alloy has a higher capacity and a better stability than the iron-containing alloys. The negative electrode charging efficiency was also improved by the Co additive. With respect to the relative discharge capacities of these alloys, it is easy to conclude that the substitution of the cobalt by iron decreases the stability and the life time cycle.

![Figure 5. Variation of the electrochemical discharge capacity of the MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Fe\textsubscript{0.75} and MnNi\textsubscript{3.55}Mn\textsubscript{0.4}Al\textsubscript{0.3}Co\textsubscript{0.75} compounds as a function of the cycle number [24].](image)
3.3.2. Cyclic Voltammetry

The cyclic voltammograms of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ compounds at different scan rates (10, 20, 30, 40 and 50 µV/s) after activation are illustrated in Fig. 6 and 7, respectively. The observed anodic peak is attributed to the oxidation of the absorbed hydrogen atoms on the surface. These figures show that the anodic current peak increases and its potential slightly shift to the positive direction with increasing the scan rate. Fig. 8 shows that the variation of the anodic peak potential ($E_{a,p}$) of this electrode has a linear dependence on log ($v$). So, it can be concluded that the electrode formed by these alloys form an irreversible system [16]. In the case, the anodic peak potential can be written by the Equation 1 as [16].

\[
\frac{d E_{a,p}}{d \log(v)} = \frac{2.3 R T}{2 \alpha n F}
\]  

where $v$ is the scan rate, $R$ is the constant of rare gas, $T$ is the temperature of the electrochemical cell, $\alpha$ is the charge transfer coefficient, $n$ the number of exchanged electron and $F$ the faraday constant. So, this formula allows the determination of the value of the charge transfer coefficient $\alpha$.

![Figure 6](image1.png)

**Figure 6.** Cyclic voltammograms of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ compound obtained at potential scan rates: 10, 20, 30, 40 and 50 µV s$^{-1}$ [24].

![Figure 7](image2.png)

**Figure 7.** Cyclic voltammograms of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ compound obtained at potential scan rates: 10, 20, 30, 40 and 50 µV s$^{-1}$ [26].
Fig. 8. Variation of the anodic peak potential of cyclic voltammograms of the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ electrodes as a function of log(v) \([24, 26]\).

Fig. 9 shows the variation of the anodic peak current versus the square root of the scan rate \((v^{1/2})\) is linear. So, for a semi-infinite diffusion and an irreversible transfer can be expressed by Equation 2 as \([16]\):

\[
I_{ap} = 0.496 \alpha^{1/2} (nF)^{3/2} s C_0 \left[\frac{VD}{RT}\right]^{1/2}
\]  

Where \(\alpha\) is the charge transfer coefficient, \(n\) is the number of the exchanged electron, \(F\) is the Faraday constant, \(s\) is the geometrical surface of the working electrode \((\text{cm}^2)\), \(C_0\) is the concentration of the diffusing species \((\text{mol cm}^{-3})\), \(D\) is the hydrogen diffusion coefficient \((\text{cm}^2\text{s}^{-1})\) and \(V\) is the potential scan rate \((\text{V s}^{-1})\). The concentration of diffusing species \(C_0\) is determined by Equation 3 as:

\[
C_0 = \frac{Q M}{m F V_M}
\]  

Where, \(M\), \(m\), \(F\) and \(V_M\) are respectively, the molecular mass, the effective mass of material, the Faraday constant and the molecular volume of material. \(Q\) is the amount of the hydrogen anodic oxidation during the anodic sweep of cyclic voltammetry \([17]\). It is expressed by equation 4 as:

\[
Q = \int_{t_1}^{t_2} i dt = \frac{1}{V} \int_{E_1}^{E_2} dE
\]  

Where, \(t_1\), \(t_2\) and \(t\) are times and \(E_1\), \(E_2\) and \(E\) are potentials, \(i\) is the anodic current and \(V\) is the scan rate. According to Equation 2, the hydrogen diffusion coefficient in this metal hydride electrode can be determined using the slope of the curve \(I_{ap} = f(v^{1/2})\).

The values of the charge transfer coefficient \(\alpha\) and the hydrogen diffusion coefficient \(D_{H\text{diff}}\), calculated from the slope of the curve \(E_{ap} = f(\log (v))\) and \(I_{ap} = f(v^{1/2})\) as explained above, are given in Table 4. The values of the charge transfer coefficient are close to 0.5. This shows symmetry of the reaction of insertion and desinsertion of hydrogen in the bulk of material in terms of reversibility and diffusivity of hydrogen.
Based on this table, it can be seen that the value of the hydrogen diffusion coefficient in the Fe-containing alloy is less than that in the Co-containing alloy. The probable reason for this decrease is the reduction of the number of available sites of hydrogen when Co is substituted by Fe. In fact, when the alloy has a higher hydrogen storage capacity and lifetime cycle, it exhibits a good diffusivity of hydrogen. On the other hand, the iron facilitates the oxidation of the surface of the alloy through the combination with oxygen to form an oxide film which in turn inhibits the hydrogen diffusion and leads to a loss in the discharge capacity.

### 3.3.3. Chronoamperometry

When we assume that the hydride alloy particles have a spherical form [15-18], the diffusion of hydrogen in the bulk of these particles can be, in the spherical coordinate, given by Eq 5 as:

$$\frac{\partial (r c)}{\partial t} = \frac{D \partial^2 (r c)}{\partial r^2}$$  

(5)

Where \( c \) is the hydrogen concentration in the alloy, \( t \) is time, \( \overline{D} \) is the average diffusion coefficient of hydrogen in the bulk and \( r \) is a distance from the center of the sphere. Eq 6 gives the solution of the diffusion equation, given by Crank [19] under different boundaries, as:

$$\frac{c - c_0}{c_s - c_0} = 1 + \frac{2a}{\pi r} \sum_{n=1}^{\infty} \frac{(-1)^n}{n} \sin \left( \frac{n\pi r}{a} \right) \exp \left( -\frac{\overline{D} \mu}{a^2} n^2 \pi^2 t \right)$$  

(6)

Where “a” is the sphere radius and \( C_0 \) and \( C_s \) are respectively the initial hydrogen concentration in the bulk and the surface hydrogen concentration of the alloy.

Where, \( F \) and \( d \) are the faraday constant and the density of material respectively.

For a large time, Eq 7 can be rewritten by equation 3 as:

$$\log(i) = \log \left( \frac{6 F \overline{D}}{d a^2} (C_0 - C_s) \right) - \frac{\pi^2 \overline{D}}{2.303 a^2} t$$  

(7)

The ± sign in Equation 3 indicates the charge state for the sign (-) and the discharge state for the sign (+). According to Equation 3, \( \overline{D} \) can be evaluated from the slope of the plot of \( \log(i) \) vs. time “\( t \)”. Taking into account that the diffusion coefficient of hydrogen \( \overline{D} \) is determined by the electrochemical impedance spectroscopy, the radius “a” of the particles, supposed as spherical, concerned by the electrochemical reaction can be calculated from the value of the slope \( \frac{\overline{D}}{a^2} \).
Fig. 10 and 11, shows that log(i) vs. time “t” may be divided into two time domains. For short time (t < 2000 s), the current decreases rapidly and is under charge-transfer control [9]. At long time (t >2000 s), the diffusion current decreases linearly with time. This reflects that the diffusion current obeys (complies with) equation 3. For the hydrogen diffusion coefficient, $D$ is determined by the electrochemical impedance spectroscopy, the variation of the average radius of the particles participating in the electrochemical reaction of the $\text{MmNi}_{3.55}\text{Mn}_{0.4}\text{Al}_{0.3}\text{Co}_{0.75}$ and $\text{MmNi}_{3.55}\text{Mn}_{0.4}\text{Al}_{0.3}\text{Fe}_{0.75}$ alloys of number of cycle are given in table 5, show the average radius decrease of the number of cycles from 100 µm before cycling to 7 µm after 13 cycles. So, we assume that the decrease of the particle size is due to the microcracking of the electrode powder alloys, during the hydrogen absorption and desorption process [14]. Before the charge-discharge process, the alloy ingot were ground mechanically and saved to less than 63 µm in glove box in order to prepare the negative electrode. So, we assume that the decrease of particle size during the cycling and the hydriding is due to the pulverization of powder induced through the expansion of the lattice volume reaching 25 % [14].
3.3.4. Impedance Spectroscopy

The kinetic of hydrogen transfer and hydrogen diffusion in the alloys has been investigated using electrochemical impedance spectroscopy. Typical impedance diagrams for the MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ electrodes after 13 cycles of charge-discharge at, respectively, 10 and 100 % state of charge are reported in Fig. 12a and Fig. 13a, respectively. These states of charge correspond to the solid solution α phase (100 %) and the hydride β phase (100 %). All these spectra consist of a very small semicircle in the high frequency (5 KHz to 400 Hz), a slope of about 45° related to the Warburg impedance, in the medium frequency region (300 Hz-100 Hz) and a large semicircle in the low frequency region (100 Hz-1 mHz), which marked dependence on the state of charge. The offset of all the curves corresponds to the electrolyte resistance ($R_e \approx 0.52 - 0.79 \, \Omega$). For these two alloys, this value is almost constant whatever the state of charge.

Based on the modeling of the kinetics of the metal hydride electrode [9, 20-21], the semicircle at a high frequency is attributed to the charge transfer reaction. So, based on this semicircle, we can determine the charge transfer resistance $R_{ct}$, the double layer capacitance $C_{dl}$ and the exchange current density $I_o$. The double layer capacitance and the current density are given, respectively, by Eqs. (8) and (9) as:

$$C_{dl} = \frac{1}{2 \pi f^* S R_{ct}}$$

$$I_o = \frac{RT}{FSR_{ct}}$$

Where $f^*$ is the proper frequency, $T$ the working temperature, $n$ the number of exchanged electrons, $S$ the geometric area of the electrode, $F$ the Faraday constant and $R_{ct}$ is the charge transfer resistance. The values of these parameters are given in Table 6.

The enlarged view of the Warburg region for the MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ and MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ electrodes after 13 cycles of charge-discharge at, respectively, 10 and 100 % state of charge are reported in Fig. 12b and Fig. 13b, respectively.

![Figure 12. a) Typical Nyquist plots for the MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ electrode at 10 and 100 % state of charge after 13 cycles b) Enlarged view of the Warburg region for MnNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ at 10 and 100 % state of charge [24].](image-url)
Figure 13. a) Typical Nyquist plots for the MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ electrode at 10 and 100 % state of charge after 13 cycles b) Enlarged view of the Warburg region for MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Fe$_{0.75}$ at 10 and 100 % state of charge.

These figures show that the - Im(Z) has a linear variation versus Re(Z) with an angle of 45° corresponding to a diffusion process of hydrogen in the bulk material. Fig. 14 shows the variation of Re(Z) versus $W^{-1/2}$ for the of MmNi$_{3.55}$Mn$_{0.4}$Al$_{0.3}$Co$_{0.75}$ compound in the Warburg region at 10 and 100 % state of charge, where $W = 2 \pi f$. Under semi-infinite diffusion conditions, the coefficient of the hydrogen diffusion can be calculated from the slope of this plot. In fact, the theoretical slope of this plot is given by Eq. (10) as:
The following conclusions could be drawn up:

(1) The values of pressure equilibrium at room temperature of MnNi₃.₅₅Mn₀.₄₄Al₀.₃Co₀.₇₅ and MnNi₃.₃₅Mn₀.₷₅Al₀.₃Fe₀.₇₅ are, respectively, equal to 0.065 and 0.024 bar, which indicates total substitution of cobalt by iron decreases the hydride stability.

(2) The MnNi₃.₅₅Mn₀.₄₄Al₀.₃Fe₀.₇₅ compound exhibits a maximum discharge capacity of 200 mAh g⁻¹ after 9 cycles. However, the MnNi₃.₅₅Mn₀.₄₄Al₀.₃Co₀.₇₅ compound exhibits a higher discharge capacity of 270 mAh g⁻¹ after 12 cycles. So, the cobalt-containing alloy has a higher capacity and good stability than the iron-containing alloy.

(3) The value of the charge transfer coefficient of the Fe and Co-containing compounds are, respectively, equal to 0.3 and 0.35. This indicates a good electrochemical reversibility reaction for the Fe-containing compounds.

(4) The hydrogen diffusion coefficient D₁ values determined by the cyclic voltammetry for the Fe and Co-containing compounds are respectively equal to, 3.96 10⁻¹⁰ cm² s⁻¹ and 5.85 10⁻¹⁰ cm² s⁻¹.

References


Table 6. Kinetic parameters of electrodes at 10 et 100 % state of charge after activation.

<table>
<thead>
<tr>
<th>MnNi₃.₅₅Mn₀.₄₄Al₀.₃Co₀.₇₅</th>
<th>MnNi₃.₃₅Mn₀.₷₅Al₀.₃Fe₀.₇₅</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 %</td>
<td>100 %</td>
</tr>
<tr>
<td>Rₑ(Ω)</td>
<td>0.76</td>
</tr>
<tr>
<td>Rₑ(Ω)</td>
<td>1.65</td>
</tr>
<tr>
<td>Cₑ(µF)</td>
<td>24</td>
</tr>
<tr>
<td>Cₑ(mF)</td>
<td>56</td>
</tr>
<tr>
<td>Rₑ(Ω)</td>
<td>84.07</td>
</tr>
<tr>
<td>Iₑ(mA)</td>
<td>15.75</td>
</tr>
<tr>
<td>Aₑ(Ω S⁻¹)</td>
<td>0.353</td>
</tr>
<tr>
<td>D(cm²s⁻¹)</td>
<td>6.64 10⁻¹⁰</td>
</tr>
</tbody>
</table>

For these tow compounds, the hydrogen diffusion coefficient is higher for the α phase and lower for the β phase than the mean value determined by the cyclic voltammetry. We assume that the value of the hydrogen diffusion coefficient depends on the number of interstitial sites susceptible to accept the hydrogen atom. In fact, in the α phase, these sites are more numerous and consequently the hydrogen diffuses easily. EIS measurements indicate that the kinetic properties of the alloy improved with increasing state of charge, mainly due to a fast hydrogen transfer in the bulk of the alloy and the higher electrocatalytic activity of the surface. The transformation of α to β phase is probably a limiting step in the mechanisms of hydrogenation of metal hydride electrode. So, the Dₑ is higher. However, in the β phase, the interstitial sites are almost saturated and the hydrogen diffuses hardly [22]. Yuan and Xu [3] and ValOzen et al. [23] indicated that the apparent hydrogen diffusion coefficient depends on the hydrogen content in the alloy in a way that it increases when the state of charge decreases.

4. Conclusion

The aim of this work is to determine the effect of the substitution of the Co by Fe in the MnNi₃.₅₅Mn₀.₄₄Al₀.₃Co₀.₇₅ compound on the thermodynamic and electrochemical properties. The following conclusions could be drawn up:

(1) The values of pressure equilibrium at room temperature of MnNi₃.₅₅Mn₀.₄₄Al₀.₃Co₀.₇₅ and MnNi₃.₃₅Mn₀.₷₅Al₀.₃Fe₀.₇₅ are, respectively, equal to 0.065 and 0.024 bar, which indicates total substitution of cobalt by iron decreases the hydride stability.

(2) The MnNi₃.₅₅Mn₀.₄₄Al₀.₃Fe₀.₇₅ compound exhibits a maximum discharge capacity of 200 mAh g⁻¹ after 9 cycles. However, the MnNi₃.₅₅Mn₀.₄₄Al₀.₃Co₀.₇₅ compound exhibits a higher discharge capacity of 270 mAh g⁻¹ after 12 cycles. So, the cobalt-containing alloy has a higher capacity and good stability than the iron-containing alloy.

(3) The value of the charge transfer coefficient of the Fe and Co-containing compounds are, respectively, equal to 0.3 and 0.35. This indicates a good electrochemical reversibility reaction for the Fe-containing compounds.

(4) The hydrogen diffusion coefficient Dₑ values determined by the cyclic voltammetry for the Fe and Co-containing compounds are respectively equal to, 3.96 10⁻¹⁰ cm² s⁻¹ and 5.85 10⁻¹⁰ cm² s⁻¹.


