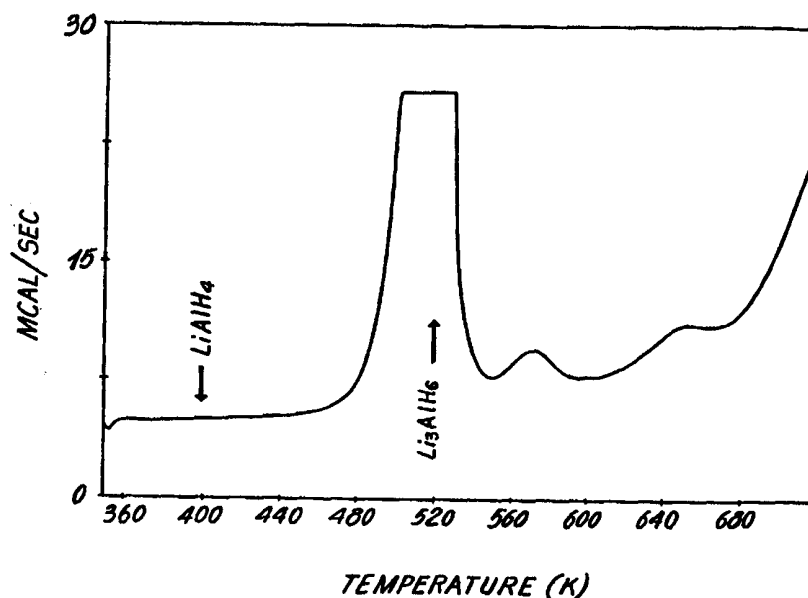




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(21) International Application Number: PCT/CA98/00927 (22) International Filing Date: 30 September 1998 (30.09.98) (30) Priority Data: 2,218,271 10 October 1997 (10.10.97) CA (71) Applicant (for all designated States except US): MCGILL UNIVERSITY [CA/CA]; 845 Sherbrooke Street West, Montreal, Quebec H3A 1B1 (CA). (72) Inventors; and (75) Inventors/Applicants (for US only): ZALUSKA, Alicja [CA/CA]; 3510 Paul-Emile-Cote, Montreal, Quebec H4E 1Z6 (CA). ZALUSKI, Leszek [CA/CA]; 3510 Paul-Emile-Cote, Montreal, Quebec H4E 1Z6 (CA). STRÖM-OLSEN, John, Olaf [CA/CA]; 443 Lansdowne Avenue, Montreal, Quebec H3Y 2V4 (CA). (74) Agents: MURPHY, Kevin, P. et al.; Swabey Ogilvy Renault, Suite 1600, 1981 McGill College Avenue, Montreal, Quebec H3A 2Y3 (CA).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>

(54) Title: METHOD OF FABRICATION OF COMPLEX ALKALI METAL HYDRIDES



(57) Abstract

Novel hydrides are produced by mechanically alloying at least two different hydrides, preferably at least one simple alkali metal hydride and at least one complex alkali metal hydride such as an alkali metal aluminum hydride; the method of production is simple and can be carried out at room temperature; the novel hydrides are useful as a source of hydrogen and have the particular advantage that after liberation of the hydrogen, the hydride is readily regenerated from the dehydrogenated hydride.

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METHOD OF FABRICATION OF COMPLEX ALKALI METAL HYDRIDES

TECHNICAL FIELD

5 This invention relates to a method of producing a complex hydride; the invention also relates to novel complex hydrides; still further the invention relates to a method of providing a source of hydrogen.

BACKGROUND ART

10 Alkali metals (lithium, sodium and potassium) form a large variety of hydrides: simple hydrides (LiH , NaH , KH) and complex hydrides with other elements, for example, boron or aluminum. Many of these compounds are commonly used in various processes of organic chemistry, acting as reducing agents.

15 Because of the high reactivity of alkali metals, simple hydrides of Li , Na and K are produced in a direct reaction between molten alkali metal and hydrogen at very high pressures and temperatures. For complex hydrides, in each case a special process of fabrication has been developed.

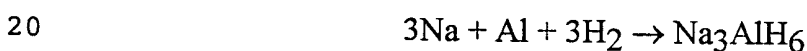
20 Lithium aluminum hydride (LiAlH_4) was discovered about four decades ago [1] and since then it has become the most common reducing agent in many chemical reactions. Sodium aluminum hydride (NaAlH_4) was first synthesized in the early sixties [2], but it has never been used as widely, because of its more difficult fabrication as compared with LiAlH_4 . Another complex hydride: sodium boron hydride (NaBH_4) [3] is
25 also known as having good reduction ability in many organo-chemical reactions [4].

30 In the search for even better reducing agents, many other complex hydrides based on lithium, sodium or potassium have been synthesized as, for example, LiBH_4 , $\text{LiAl}_4\text{H}_{13}$, LiAl_2H_7 , Li_3AlH_6 , KAlH_4 , KBH_4 , $\text{K}(\text{Al}(\text{BH}_4)_4)$.

35 In all cases the synthesis of these complex hydrides is performed through a chemical reaction under special conditions. For example, commercial fabrication of LiAlH_4 involves the reaction of LiH with AlCl_3 in diethyl ether [5]. In the early sixties Ashby and co-workers developed an alternative route of production of complex metal hydrides by

direct synthesis [2, 6, 7]. This method can be applied to the production of LiAlH_4 , NaAlH_4 , KAlH_4 and CsAlH_4 . According to Ashby, synthesis of, for instance, NaAlH_4 can be performed in the following way: "one charges either the alkali metal or its hydride to an autoclave with activated
5 aluminum powder in a solvent such as tetrahydrofuran. The mixture then is subjected to a pressure of 2000 p.s.i. (about 140 bar) with hydrogen and heated to 150°C for several hours. After the absorption is complete the mixture is cooled and the complex aluminum hydride is separated from excess of aluminum by filtration. NaAlH_4 can be isolated by addition of a
10 hydrocarbon such as toluene to the tetrahydrofuran solution, followed by vacuum distillation of the tetrahydrofuran" [2].

Another complex alkali metal hydride, Na_3AlH_6 , was primarily fabricated by Zakharin et al [8] in the reaction of NaH and NaAlH_4 at 160°C in heptane. However, due to insolubility of the product
15 in all solvents tested, it could not be purified. In response to the drawbacks of the above method Ashby et al proposed again a direct method for the synthesis of Na_3AlH_6 hydride [2, 7]. According to the direct method, Na_3AlH_6 can be synthesized by the following reaction:



This reaction should be performed in toluene at 165°C and at 5000 p.s.i. (about 350 bar) of hydrogen pressure.

Synthesis of Li_3AlH_6 was first discovered by Ehrlich et al
25 [9,10]. Later on, Mayet and co-workers [11, 12, 13, 14] developed another method of fabrication of Li_3AlH_6 , which provided better reproducibility and higher purity of the hydride. In this method a solution of LiAlH_4 in ether is added drop by drop into a suspension of LiH in toluene heated up to the temperature of 50°C. The mixture is kept for several hours at 50°C to
30 eliminate ether and is subsequently heated up to 95°C to complete the reaction. Hot catalyst: $\text{Al}(\text{C}_2\text{H}_5)_3$ in toluene or etherate of triethylaluminum, is added during the first step of the process.

All the above methods for the production of complex alkali metal hydrides suffer from many drawbacks, i.e. the need to use solvents or

dispersing liquids (hydrocarbons) with activators, multi-step character and relatively poor yield and reproducibility.

In order to overcome these problems, another method of fabrication of complex alkali metal hydrides was developed more recently
5 by Dymova and co-workers [15, 16, 17, 18]. In this method the solvents were eliminated, but instead a reaction at a temperature above the melting point of the alkali metal was proposed. The reaction of molten alkali metals (Li, Na, K, Cs) with aluminum was performed at a temperatures of 200-400°C and at hydrogen pressure of 100-400 bar.

10 In conclusion, all the previous methods of fabrication of complex alkali metal hydrides have three main disadvantages:

- i) inconvenience of the use of toxic and flammable solvents
such as toluene and tetrahydrofuran;
- ii) very high hydrogen pressures (100 - 400 bar); and
- 15 iii) high temperatures (100°C - 400°C).

Alkali-metal-based complex hydrides were developed with a clear purpose to serve as reducing agents in chemical reactions, mainly in organic chemistry. However, other applications of these hydrides have also been considered. Most of these hydrides undergo decomposition at high
20 temperatures. The decomposition releases hydrogen and therefore alkali-metal hydrides can be used in some cases as an immediate source of hydrogen [19]. For example, LiAlH_4 decomposes when heated up to a temperature of 125°C and releases gaseous hydrogen. This phenomenon has been exploited in equipment for hydrogen storage. It should be stressed
25 however that these applications use alkali metal hydrides for a single, irreversible hydrogen release. There is no way to reverse the dehydrogenation reaction in these prior systems, without repeating the whole chemical procedure used in the production of the hydride, which obviously cannot be accomplished inside the hydrogen storage tank.

30 The present invention seeks to develop materials which can be used as a reversible source of hydrogen, i.e., which can be reversibly hydrided and dehydrided in subsequent cycles of hydrogen admission and evacuation, without any other treatment. The only prior method of yielding reversibility of hydrogenation in alkali-metal-based hydrides was reported
35 in a recently published paper of Bogdanovic and Schwickardi [20]. The

authors studied traditional alkali metal aluminum hydrides (NaAlH_4 and Na_3AlH_4) and state that "the reverse reaction has not been accomplished" until their method of doping with special Ti-based catalysts was developed [21]. The authors fabricated alkali metal hydrides in a conventional way (following the process described by Zakharin [8]). For example, Na_3AlH was prepared from NaAlH_4 and NaH in heptane under hydrogen. The suspended reagents were intensively stirred at 162°C for 72h under a hydrogen pressure of 140 bar. The reversible hydrogenation was achieved when the materials were treated with 2 mol% of $\beta\text{-TiCl}_3$ in ether or with 3 mol% $\text{Ti}(\text{OBu})_4$ in ether.

Although the results showed significant improvement of the hydride performance as compared to the undoped materials, the authors indicate that the kinetics of the process were still unsatisfactorily slow (for example, one absorption cycle took five to twenty or even 100 hours) and suffered from cyclic instability.

DISCLOSURE OF THE INVENTION

In the present invention a totally different method of fabrication of complex alkali metal hydrides has been developed which is based on a mechano-chemical reaction. The resulting hydrides exhibit outstanding, reversible hydrogenation properties, which remarkably surpass any previously reported performance.

In accordance with one aspect of the invention there is provided a method of producing a complex hydride comprising: mechanically alloying a first particulate hydride material with a second particulate hydride material, said first and second particulate hydride materials each being selected from the group consisting of alkali metal hydrides, alkali metal borohydrides, alkali metal aluminum hydrides and mixtures thereof, said first and second particulate hydride materials being different.

In accordance with the invention complex alkali-metal hydrides are fabricated in a solid-state reaction, i.e., a mechano-chemical reaction or mechanical alloying. Instead of using ion-carriers necessary for the chemical reaction to proceed (in the form of solvent or activators), and instead of applying high hydrogen pressures, a physical contact between the

reagents is provided by means of mechanical treatment in the course of the reaction or alloying, which is the essence of the process.

Mechanical treatment during the reaction provides enhanced local reactivity of the reagents, by means of the continuous creation of fresh surfaces unaffected by oxides and hydroxides, and introduces local stress and deformation which is believed to enhance the rate of reaction. The method is performed on dry powders of the components (being not in the form of a slurry and without any solvents or additions). The method does not require high hydrogen pressures or elevated temperatures and can be performed under inert gas atmosphere or a hydrogen gas atmosphere, at normal pressure and at room temperature (20°C). Also, no special catalyst, reaction agents or other activators are required. The method can be easily accomplished by grinding, agitating or ball milling of the appropriate reagents, and this can be carried out at a wide range of impact energies and grinding, agitating or milling times. The method may be carried out in the absence of a solvent.

In another aspect of the invention there is provided novel complex hydrides which function as a recyclable store of hydrogen. In the present invention the novel complex hydrides include hydrides which are a novel combination of elements as well as hydrides of known chemical composition but which have a novel physical structure.

In a further aspect of the invention there is provided a method of providing a source of hydrogen gas comprising: liberating hydrogen from a complex alkali metal hydride derived from the mechanical alloying of at least two different hydrides of alkali metals, with formation of a supply of hydrogen gas and a dehydrogenated form of the complex hydride, removing the liberated hydrogen, and regenerating the complex hydride as a future source of hydrogen by exposing the dehydrogenated form to hydrogen gas and absorbing the hydrogen gas in the dehydrogenated form.

DESCRIPTION OF PREFERRED EMBODIMENTS

i) Hydrides

The mechano-chemical method of the invention for the production of complex alkali metal hydrides is universal to such an extent that it can be effectively applied to almost any required composition of the

complex hydride. Therefore the method is not limited to the production of known complex hydrides, but can be also used to develop new complex hydrides with new hydrogenation properties.

Complex hydrides in the context of the present invention
5 contemplates hydride compounds containing at least two different metals, at least one of which is an alkali metal.

In especially preferred embodiments the first particulate
hydride material comprises at least one hydride selected from the group
consisting of alkali metal hydrides (LiH, NaH, KH) and alkali metal
10 borohydrides (LiBH₄, NaBH₄, KBH₄) and the second particulate hydride
material comprises at least one alkali metal aluminum hydride (LiAlH₄,
NaAlH₄, KAlH₄).

The first and second particulate hydride materials suitably
have a particle size of less than 100 μ m.

15 By varying the proportions of the first and second hydride
materials and the composition of the respective materials, a wide variety of
different complex hydrides are readily formed.

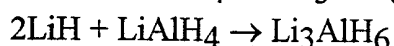
Suitably the molar ratio of the first hydride material to the
second hydride material is 5:1 to 1:5 and preferably 3:1 to 1:2.

20 In a particular embodiment, fabrication of the complex alkali-
metal based hydrides comprises mixing powders of simple hydrides (LiH,
NaH, KH) with other hydride complexes (e.g. AlH₃) or other alkali metal
hydrides (LiAlH₄, LiBH₄, NaAlH₄, NaBH₄, KAlH₄, KBH₄ etc.) in a
desired proportion and applying mechanical treatment at high impact
25 energy, for example, by grinding, agitating or ball milling, in an inert
atmosphere, for instance of argon or a hydrogen atmosphere. The
fabrication can be carried out at a temperature below 100°C and in
particular proceeds efficiently at room temperature (20°C).

By way of example the powders may be ball milled in a high
30 energy ball mill such as that available under the Trade-mark SPEX 8000 or
Model SP2100 (from SCP. Science of St-Laurent, Quebec). These ball
mills typically employ steel balls or tungsten carbide balls. Suitable
operating parameters include a weight ratio of mill balls to hydrides of 30:1
to 2:1 and a milling time of 0.25 to 20 hours, more usually 2 to 5 hours,

however, a wide range of grinding, agitating or ball milling conditions can be employed to effect the desired mechanical alloying.

Production of, for example, Li_3AlH_6 or Na_3AlH_6 can be easily and effectively performed according to the reactions previously used in other methods, but employing the simpler method of the invention:



Other new complex hydrides can be produced by the substitution of certain alkali hydride complexes by others in the initial mixture of hydrides. This leads to the formation of complex hydrides which have not been synthesized until now, due to the limitations of the chemical methods. In the present invention a great number of such complex hydrides have been synthesized, being a combination of simple hydride complexes, for example:



In the case of the above and analogous reactions (which have not been previously established by chemical methods) the reaction routes are still not fully determined and are under investigation. Results demonstrate that these reactions often take place with formation of new, unknown crystallographic structures. More importantly, these new phases exhibit new hydrogenation performance, which leads to reversibility of hydrogen storage in alkaline metal hydrides.

The novel hydrides of the invention are useful as reducing agents, the method of the invention providing a simpler procedure than heretofore available for the production of such reducing agents. The hydrides of the invention are also useful as a recyclable store of hydrogen. After liberating the hydrogen from the hydrides of the invention with formation of a dehydrogenated form of the hydride, the hydride is readily regenerated on exposure of the dehydrogenated form to hydrogen.

In a particular embodiment of the invention there is provided a complex particulate hydride produced by mechanical alloying of at least two different hydrides and which is characterized as a recyclable store of hydrogen which liberates hydrogen at an elevated temperature within 60 minutes, and having a dehydrogenated state which absorbs hydrogen within 60 minutes.

In a further particular embodiment of the invention there is provided the dehydrogenated form of the complex hydride of the invention characterized by an ability to absorb hydrogen gas with generation of said hydride.

ii) Reversible Hydrogenation

Complex alkali metal hydrides fabricated by the mechano-chemical process exhibit unique properties in terms of reactivity in hydriding/dehydriding reactions. No other alkali metal hydrides undergo reversible cycles of hydrogenation/dehydrogenation under gaseous hydrogen at medium pressures (20 - 50 bars), when being in the solid powder form, with no solvents, catalysts or activators. Excellent hydrogenation/dehydrogenation cyclability occurs in the novel complex hydrides synthesized in this invention. Results show that reversibility of hydrogenation in, for example, Na_3AlH_6 , produced by mechano-chemical reaction, occurs with fast kinetics, even without any catalytic treatment. Reaction rates are significantly faster or similar to these presented in ref. [20] for hydrides treated by the special Ti-doping method [21]. Moreover, reversible hydrogen capacities are much higher than these reported in [20].

Newly designed and fabricated complex alkali metal hydrides (consisting of either novel forms of previously known compositions or of new compositions) represent a whole spectrum of materials with controllable hydrogenation properties. It has been established that, in some cases, lithium-based complexes decrease plateau pressures. Therefore in order to design materials with lower operational hydrogen pressures more lithium complexes are introduced in the course of the mechano-chemical reaction into the structure of the complex hydride as, for example, in $\text{Li}_2\text{NaAlH}_6$. On the other hand, boron-based complexes can increase the plateau pressure in some complex hydrides and therefore boron-based complexes are advantageous in the hydride to allow lower operational

temperatures, if this is required (e.g. $\text{Na}_2\text{BA1H}_6$). Reversible hydrogen capacity is obviously dependent on the relative ratios of alkali metals (Li, Na, K) and other elements (aluminum, boron) in the complex hydride. Therefore mechano-chemical reactions permit design of an optimum composition of the complex alkali metal hydrides depending on the required operational conditions for reversible hydrogen storage (hydrogen pressure, temperature, hydrogen capacity).

BRIEF DESCRIPTION OF DRAWINGS

FIG. 1 is a DSC scan of a hydride of the invention corresponding to Li_3AlH_6 ;

FIG. 2 is a DSC scan of a hydride of the invention corresponding to Na_3AlH_6 ;

FIG. 3 is an X-ray diffraction pattern of a hydride formed by mechanically alloying LiH, NaH, LiAlH_4 and NaAlH_4 , in accordance with the invention;

FIG. 4 shows the hydrogen desorption of the hydride of Example 2 and Fig.3, and hydrogen absorption of the dehydrogenated form of the hydride with time, at 230°C ; and

FIG. 5 is a plot of pressure-composition-isotherms for three hydrides produced by the method of the invention.

EXAMPLES

Example 1.

Fabrication of Li_3AlH_6 and Na_3AlH_6 was performed by mechanical alloying of the respective amounts of LiH and LiAlH_4 to produce Li_3AlH_6 and of NaH and NaAlH_4 to produce Na_3AlH_6 in a stainless-steel vial with stainless-steel balls. The reaction took place at room temperature under argon atmosphere, with no solvents, catalysts or activators. In each case the mixture of hydrides was ball milled in a commercial ball mill SPEX 8000 (Trade-mark) having stainless steel balls at a weight ratio of balls to hydrides of 16:1, and a milling time of 3 hours at 20°C .

Formation of Li_3AlH_6 and Na_3AlH_6 was confirmed by x-ray diffraction and also by differential scanning calorimetry (DSC).

Fig. 1 shows a DSC scan at a scan rate of 20 deg./min. of the mechanically treated mixture of $2\text{LiH} + \text{LiAlH}_4$. Instead of the endothermic peak of the decomposition of LiAlH_4 occurring at 125°C (398 K) a large endothermic effect was observed at the temperature characteristic for Li_3AlH_6 , which is 240°C - 260°C according to Ref. [11] (decomposition of LiH is beyond the registered temperature range).

Fig. 2 presents a DSC curve at a scan rate of 20 deg./min. for a mechanically alloyed mixture of $2\text{NaH} + \text{NaAlH}_4$, and again there is no endothermic effect of the decomposition of NaAlH_4 (which occurs at 185°C , i.e., 458 K), but instead an endothermic reaction at higher temperature is observed, characteristic for the decomposition of Na_3AlH_6 (i.e. 280°C (553 K) [2]). Additional proof of the mechano-chemical reaction is the second peak seen in the DSC scan of Fig. 2. This endothermic effect can be attributed to the decomposition of NaH . However, the temperature of the peak is significantly shifted (as compared to the decomposition of pure NaH) which indicates that in this case NaH was a product of the previous decomposition of Na_3AlH_6 , which occurred within the first endothermic effect.

Example 2.

New hydride structures have been formed after mechano-chemical reaction of the appropriate mixtures consisting of the following hydrides in various proportions: NiH , NaH , LiAlH_4 , NaAlH_4 , NaBH_4 , KAlH_4 , LiB_4 , etc. The results demonstrate that the mechanical alloying method of the invention has wide applicability in the production of hydrides. Fig. 3 shows the x-ray diffraction pattern of a new, simple structure of the hydride formed as a result of mechano-chemical reaction involving four components: LiH , NaH , LiAlH_4 and NaAlH_4 . Instead of the diffraction reflections characteristic for the mixture of the components or other known complex hydrides, a single phase with simple bcc structure was observed. The milling conditions were the same as for Example 1.

Example 3.

The hydride of Example 2 and Fig. 3 fabricated by mechanical alloying, exhibits reversible hydriding/dehydriding behavior as

a result of the reaction with gaseous hydrogen in a hydrogen storage container. After fabrication, the powders were placed in a reaction chamber of a gas titration system. Hydrogen absorption and desorption was measured as a result of the pressure change of hydrogen in the chamber. As
5 seen in Fig. 4, absorption of about 3 wt. % of hydrogen occurred at 230°C within about 30 min. and desorption within 40 min. Cycles of absorption and desorption were repeated by cyclic admission and evacuation of hydrogen.

10 Example 4.

Materials prepared by mechano-chemical reaction exhibit different thermodynamic properties (equilibrium pressure) in dependence on the initial composition of the mixture of hydrides. As a result, operational conditions of hydrogenation/dehydrogenation cycles can be
15 adjusted by changes of the material composition. Fig. 5 shows pressure-composition-isotherms (taken at 220°C) for three different hydrides.

- (a) - Na_3AlH_6
- (b) - $\text{Na}_{1.8}\text{Li}_{0.6}\text{B}_{0.6}\text{AlH}_6$
- 20 (c) - $\text{Li}_{1.8}\text{Na}_{1.2}\text{AlH}_6$.

Hydride (a) is the hydride of Fig. 2, in Example 1.

References:

- [1] H. Schlesinger and H. Brown, US Pat. No. 2,461,661, 1949.
- [2] E. C. Ashby and P. Kobetz, Inorg. Chem., Vol. 5, No. 9, p. 1616,
25 1963.
- [3] H. Schlesinger and H. Brown, US Pat. No. 2,534,533, 1950.
- [4] J. Osby, S.W. Heinzman and B. Ganem, J. Am. Chem. Soc., Vol. 108.No. 1, p. 67, 1986
- [5] A. E. Finholt, A. C. Bond and H. Schlesinger, J. Am. Chem. Soc.,
30 Vol. 69, p. 1199, 1947.
- [6] E. C. Ashby, G. J. Brendel and H. Redman, Inorg. Chem., Vol.2, p. 449, 1963.
- [7] E. C. Ashby, French Pat. No. 1,235,680, 1960.
- [8] L. Zakharin and V. Gavrilenko, Dokl. Akad. Nauk SSSR, Vol. 1, No.
35 145, p. 793, 1962.

- [9] R. Ehrlich, A. Young, G. Rice, J. Dvorak, P. Shapiro and H. F. Smith, J. Amer. Chem. Soc., Vol. 88, No. 4, p. 858, 1996.
- [10] J. J. Dvorak and R. Ehrling, US Pat. No. 3,357,806, 1967.
- 5 [11] J. Mayet, S. Kovacevic and J. Tranchant, Bull. Soc. Chim. Fr., Vol. 22, p. 504, 1973.
- [12] J. Mayet, S. Kovacevic and J. Tranchant, Bull. Soc. Chim. Fr., Vol. 2, p. 506, 1973.
- [13] J. Tranchant and J. Mayet, French Pat. No. 7020279, 1970.
- [14] J. Tranchant and J. Mayet, French Pat. No. 6914185, 1969.
- 10 [15] T. N. Dymova, N. G. Eliseeva, S. Bakum and Y. M. Dergachev, Dokl. Nauk SSSR, Vol. 215, p. 1369, 1974.
- [16] T. N. Dymova, Y. M. Dergachev, V. A. Sokolov and N. A. Grechanaya, Dokl. Akad. Nauk SSR, Vol 224, No. 3, P. 591, 1975.
- [17] T. N. Dymova, Russ. Pat. No. 265868, 1968.
- 15 [18] T. N. Dymova, S. I. Bakum and V. M. Bakulina, Russ. Pat. No. 223788, 1967.
- [19] C. Ward, D. Stanga, L. Pataki and R. Venter, J. Power Sources, Vol. 41, p. 335, 1993.
- [20] B. Bogdanovic and M. Schwickardi, J. Alloys and Comp., Vol. 253, p.1, 1997.
- 20 [21] B. Bogdanovic, German Pat. Appln. No. 195 26 434.7, 1995.

CLAIMS

1. A method of producing a complex hydride comprising:
mechanically alloying a first particulate hydride material with a
second particulate hydride material, said first and second particulate
5 hydride materials each being selected from the group consisting of alkali
metal hydrides, alkali metal borohydrides, alkali metal aluminum hydrides
and mixtures thereof, said first and second particulate hydride materials
being different.
- 10 2. A method according to claim 1, wherein the mechanical
alloying is carried out in an inert atmosphere.
3. A method according to claim 1, wherein the mechanical
alloying is carried out in an atmosphere of hydrogen.
- 15 4. A method according to claim 1, 2 or 3, wherein the
mechanical alloying is carried out at a temperature below 100°C in the
absence of a solvent.
- 20 5. A method according to claim 1, 2, 3 or 4, wherein said first
particulate hydride material comprises at least one hydride selected from
the group consisting of alkali metal hydrides and alkali metal borohydrides;
and said second particulate hydride is an alkali metal aluminum hydride.
- 25 6. A method according to claim 1, 2, 3, 4 or 5, wherein said first
and second particulate hydride materials have a particle size of less than
100µm.
7. A method according to any one of claims 1 to 6, in which the
30 molar ratio of the first hydride material to the second hydride material is 5:1
to 1:5.
8. A method according to claim 7, wherein said molar ratio is
3:1 to 1:2.
- 35

9. A method according to any one of claims 1 to 8, wherein said alloying comprises ball milling said first and second particulate hydride materials at a weight ratio of mill balls to said particulate hydride materials of 30:1 to 2:1 for a time of 0.25 to 20 hours.
- 5 10. Novel complex hydrides produced by the method of any one of claims 1 to 9.
- 10 11. A hydride selected from $\text{NaLi}_2\text{AlH}_6$, Na_2BAlH_6 and $\text{Na}_2\text{LiAlH}_6$.
12. A hydride Li_3AlH_6 having a DSC scan as shown in Fig. 1.
- 15 13. A hydride Na_3AlH_6 having a DSC scan as shown in Fig. 2.
- 14 A hydride produced by the method of claim 1, 2, 3 or 4, in which said first and second materials comprise LiH , NaH , LiAlH_4 and NaAlH_4 and having an x-ray diffraction pattern as shown in Fig. 3.
- 20 15. A complex particulate hydride produced by mechanical alloying of at least two different hydrides and which is characterized as a recyclable store of hydrogen which liberates hydrogen at an elevated temperature within 60 minutes and having dehydrogenated state which absorbs hydrogen within 60 minutes.
- 25 16. A hydride according to claim 15, wherein said at least two different hydrides are selected from the group consisting of alkali metal hydrides, alkali metal borohydrides, alkali metal aluminum hydrides and mixtures thereof.
- 30 17. A hydride according to claim 15, wherein said at least two different hydrides comprise a first particulate hydride material and a second particulate hydride material and said first particulate hydride material comprises at least one hydride selected from the group consisting of alkali

metal hydrides and alkali metal borohydrides; and said second particulate hydride is an alkali metal aluminum hydride.

18. A method of providing a source of hydrogen gas comprising:
5 liberating hydrogen from a complex alkali metal hydride derived from the mechanical alloying of at least two different hydrides of alkali metals, with formation of a supply of hydrogen gas and a dehydrogenated form of said complex hydride,
removing said liberated hydrogen, and
10 regenerating said complex hydride as a future source of hydrogen by exposing said dehydrogenated form to hydrogen gas and absorbing said hydrogen in said dehydrogenated form.

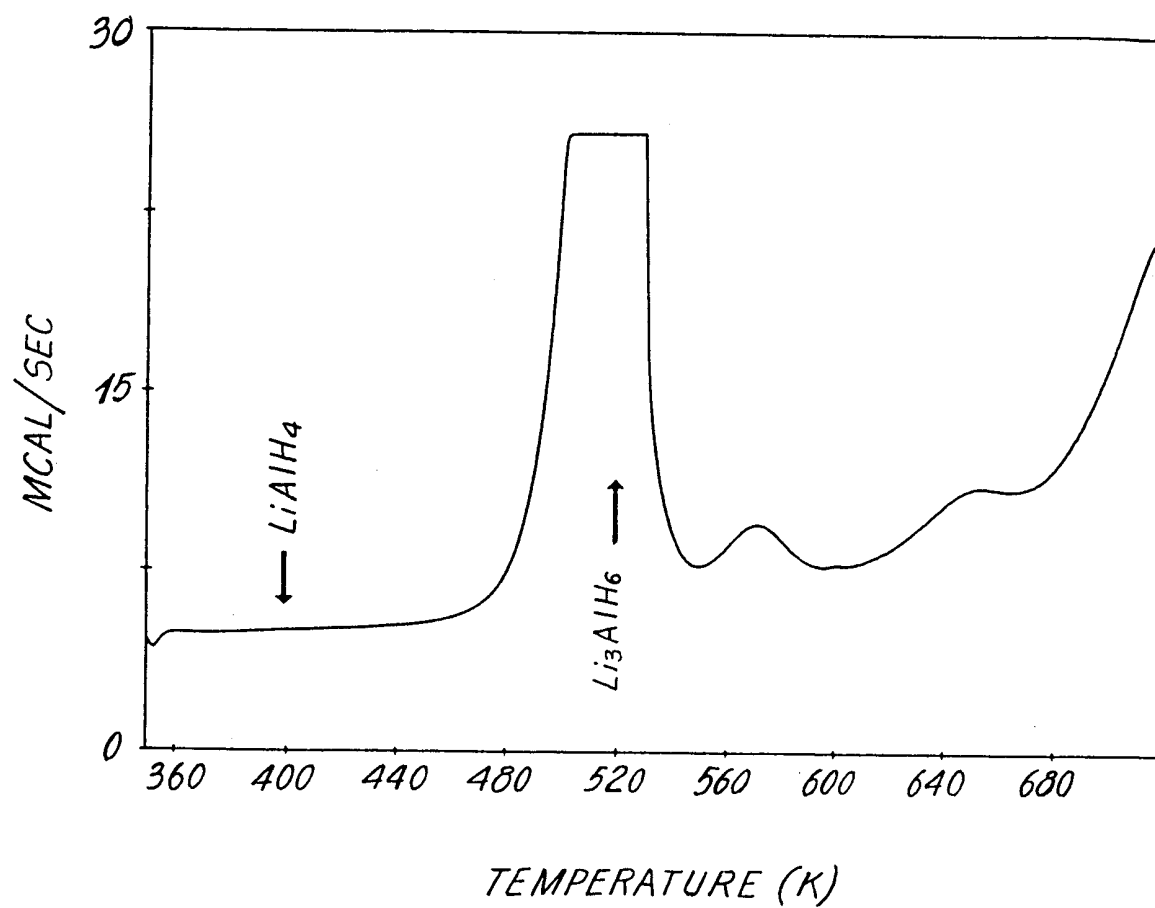
19. A method according to claim 18, wherein said at least two
15 different hydrides are selected from the group consisting of alkali metal hydrides, alkali metal borohydrides, alkali metal aluminum hydrides and mixtures thereof.

20. A method according to claim 18, wherein said at least two
20 different hydrides comprise a first particulate hydride material and a second particulate hydride material and said first particulate hydride material comprises at least one hydride selected from the group consisting of alkali metal hydrides and alkali metal borohydrides; and said second particulate hydride is an alkali metal aluminum hydride.

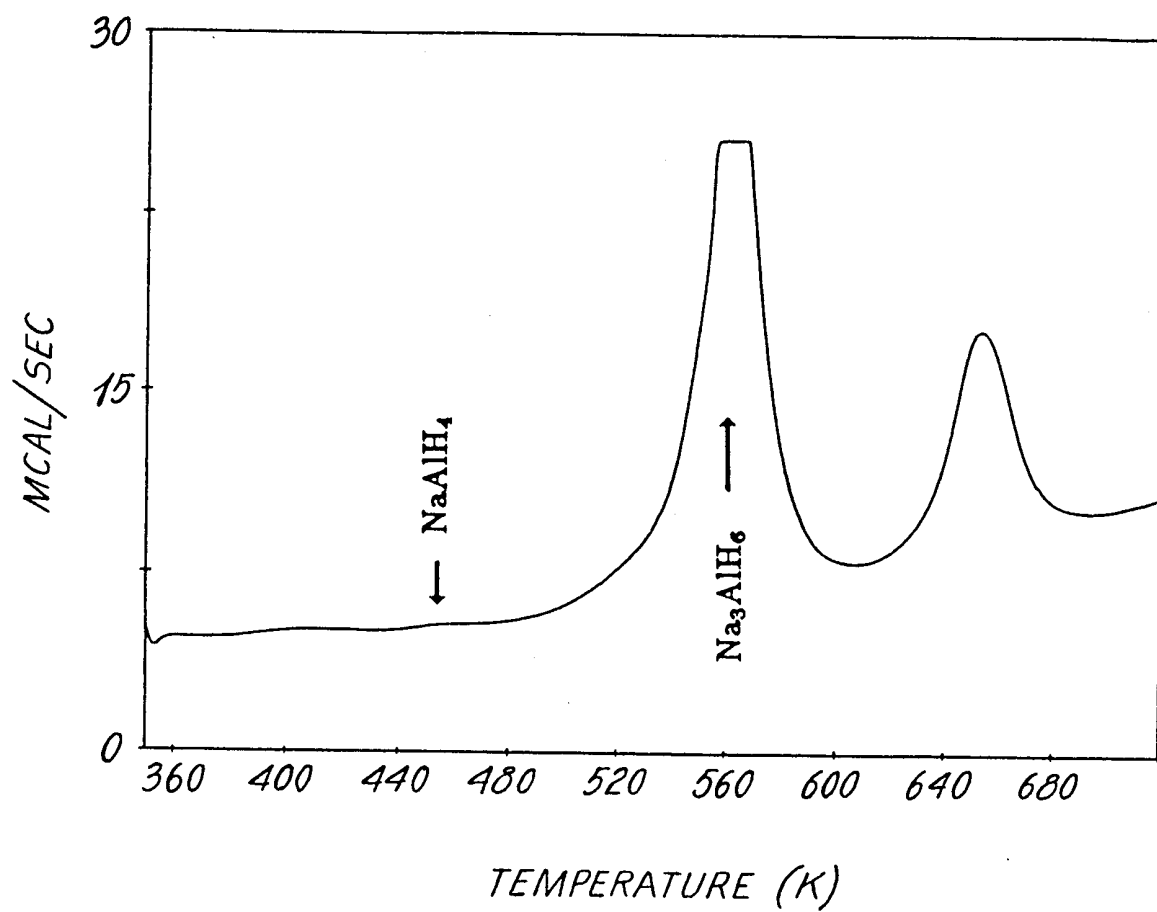
21. A method according to claim 18, wherein said hydride is
25 selected from $\text{NaLi}_2\text{AlH}_6$, Na_2BAlH_6 and $\text{Na}_2\text{LiAlH}_6$.

22. A method according to claim 18, wherein said hydride is as
30 defined in claim 11, 12, 13 or 14.

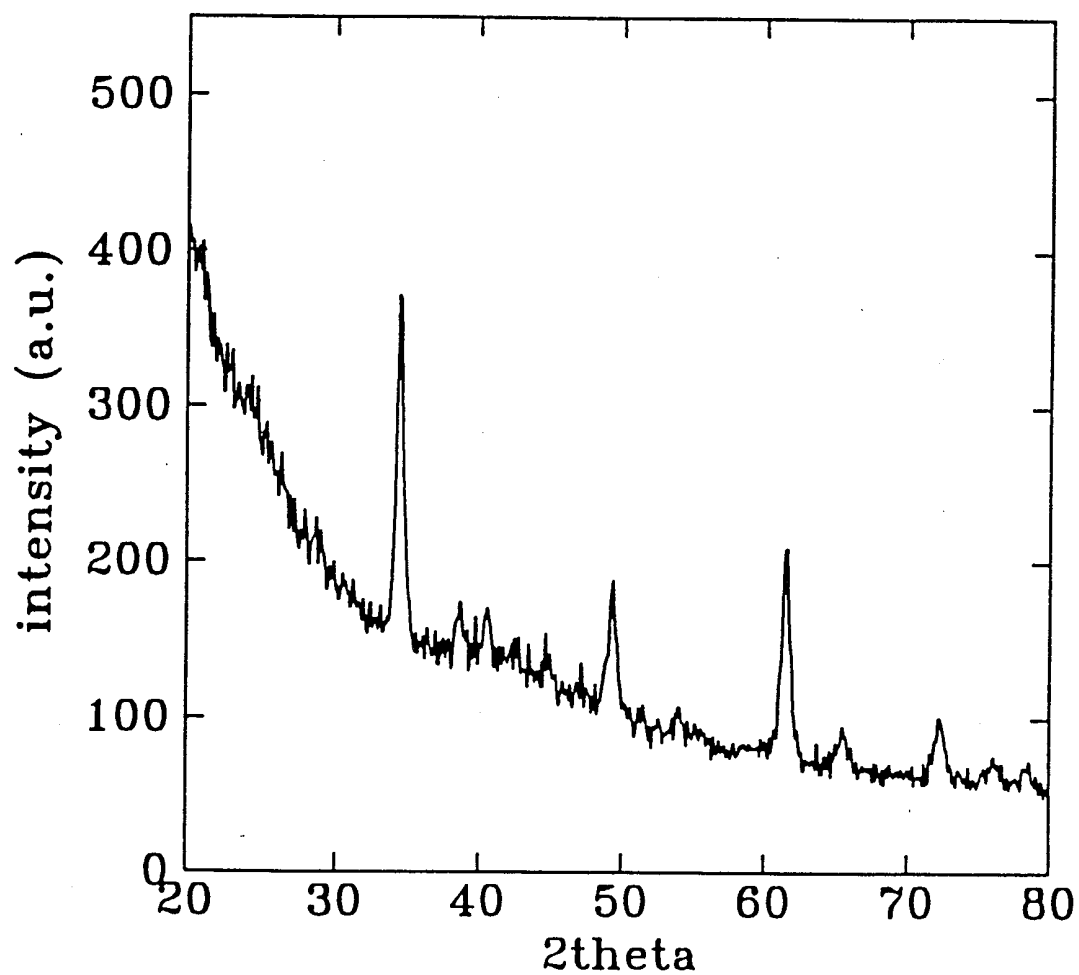
23. A dehydrogenated form of the complex particulate hydride defined in claim 15, 16 or 17, characterized by an ability to absorb hydrogen gas with generation of said hydride.

*Fig. 1*

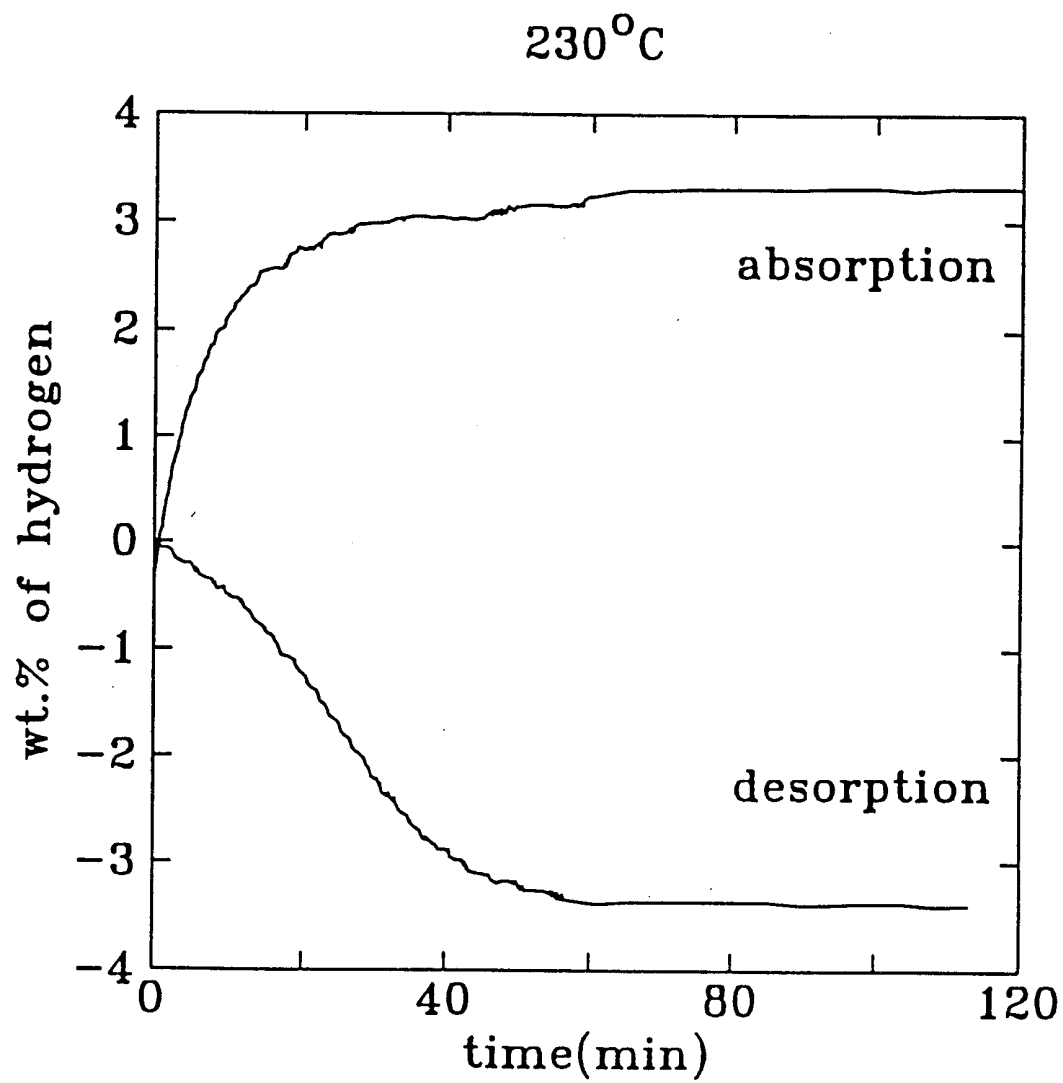
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*Fig. 2*

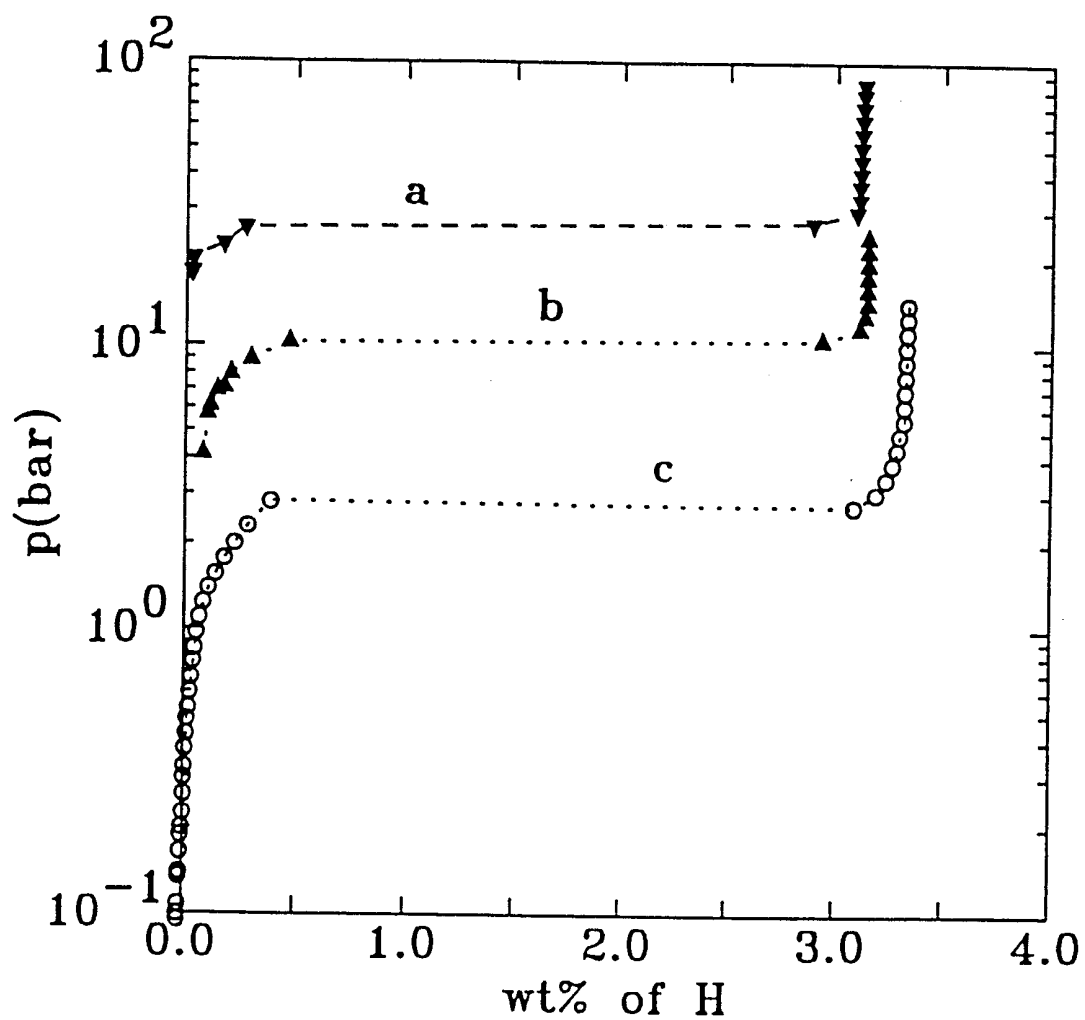
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*Fig. 3*

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*Fig. 4*

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*Fig. 5*

INTERNATIONAL SEARCH REPORT

In national Application No

PCT/CA 98/00927

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C01B6/24 C01B3/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C01B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	CHEMICAL ABSTRACTS, vol. 119, no. 22, 29 November 1993 Columbus, Ohio, US; abstract no. 240428, DYMOVA, T. N. ET AL: "Solid-phase reaction of crystalline aluminum trihydride. Formation of alkali metal tetra- and hexahydroaluminates" XP002089013 see abstract & KOORD. KHIM. (1993), 19(7), 529-34 CODEN: KOKHDC;ISSN: 0132-344X,1993, ----	1-5,7,8, 10,15-17
A	FR 2 067 844 A (FRANCE ETAT) 20 August 1971 cited in the application see the whole document ----- -/--	1,2,10, 15,16

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

4 January 1999

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/CA 98/00927

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DE 195 26 434 A (STUDIENGESELLSCHAFT KOHLE MBH) 23 January 1997 cited in the application see page 2, line 41 - line 60 see page 3, line 30 - page 4, line 6 see example 3	11
A	IDEM	15,18
X	--- BOGDANOVIC B ET AL: "Ti-doped alkali metal aluminium hydrides as potential novel reversible hydrogen storage materials" JOURNAL OF ALLOYS AND COMPOUNDS, vol. 253-254, 20 May 1997, page 1-9 XP004126498 see the whole document	11
A	IDEM	1,10,15,18
X	--- CLAUDY, PIERRE ET AL: "Reactions of lithium and sodium aluminum hydride with sodium or lithium hydride. Preparation of a new alumino-hydride of lithium and sodium (LiNa ₂ AlH ₆)" MATER. RES. BULL. (1982), 17(12), 1499-504 CODEN: MRBUAC;ISSN: 0025-5408,1982, XP002089012 see the whole document	11
A	IDEM	1-5,8

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/CA 98/00927

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
FR 2067844 A	20-08-1971	NONE	
DE 19526434 A	23-01-1997	WO 9703919 A	06-02-1997
		EP 0840707 A	13-05-1998