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Phase Stability and Fast Ion Conductivity in the Hexagonal LiBH₄-LiBr-LiCl Solid Solution

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Supporting Information Description

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- Figure S3: Rietveld refinement of sample s1.
- Figure S4: HP-DSC signal of sample s3.
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- Figure S7: Density as a function of composition.
- Figure S8: Activation energy as a function of volume.

Samples preparation

Sample Name	Composition (Molar Fraction)			Synthesis & Treatments
	LiBH ₄	LiBr	LiCl	
s1	0.33	0.33	0.33	BM 1.5 h + AN 2h 250 °C
s2	0.38	0.33	0.29	BM 1.5 h + AN 2h 250 °C
s3	0.38	0.33	0.29	BM 1, 2, 4, 6, 22, 24, 41 h
s4		0.50	0.50	BM 1.5 h
s5	0.80	0.20		BM 1.5 h + AN 4h 250 °C
s6	0.70	0.30		BM 1.5 h + AN 2h 250 °C
s7	0.60	0.40		BM 1.5 h + AN 2h 250 °C
s8	0.50	0.50		BM 1.5 h + AN 2h 250 °C
s9	0.40	0.60		BM 1.5 h + AN 2h 250 °C
s10	0.60	0.20	0.20	BM 1.5 h + AN 2h 250 °C
s11	0.22	0.58	0.20	BM 1.5 h + AN 2h 250 °C
s12	0.22	0.19	0.59	BM 1.5 h + AN 2h 250 °C
s13	0.60	0.10	0.30	BM 1.5 h + AN 2h 250 °C
s14	0.21	0.69	0.10	BM 1.5 h + AN 2h 250 °C
s15	0.40	0.40	0.20	BM 1.5 h + AN 2h 250 °C
s16	0.40	0.50	0.10	BM 1.5 h + AN 2h 250 °C
s17	0.49	0.41	0.10	BM 1.5 h + AN 2h 250 °C
s18	0.50	0.30	0.20	BM 1.5 h + AN 2h 250 °C
s19	0.50	0.20	0.30	BM 1.5 h + AN 2h 250 °C
s20	0.60	0.30	0.10	BM 1.5 h + AN 2h 250 °C
s21	0.70	0.20	0.10	BM 1.5 h + AN 2h 250 °C

In the table and in Figure S1 the samples prepared and the conditions of the synthesis are reported.



Figure S1. Composition plot for all samples prepared. Preparation methods and compositions of the investigated samples are indicated in the table. Ball milled (BM) time is reported, together with annealing (AN) time and temperature.

Verification of the Vegard's law for the LiBr-LiCl cubic solid solution

The LiBr-LiCl phase diagram reported in literature,¹ presents a complete solid solution with no miscibility gap and a liquidus curve with a thermal minimum at 522 °C, $y_{LiCl} = 0.36$.

An equimolar mixture of LiBr and LiCl (sample *s4*) was prepared to verify the Vegard's law in the LiBr-LiCl binary system. An *in situ* high-temperature X-ray Powder Diffraction (XPD) measurement was performed on $(\text{LiBr})_{0.5}(\text{LiCl})_{0.5}$ (Bragg-Brentano geometry) into Aanton-Paar XRK 900 reactive chamber, under dynamic vacuum. Patterns were collected in a temperature range RT < T < 400 °C with a heating/cooling rate of 5 °C/min. In order to remove the presence of water in the powder, a preliminary annealing at 120 °C for 10 h was performed. Calcium Oxide (CaO, purity >99% from Sigma-Aldrich) was used as internal standard.

Analysing by X-ray diffraction the $(\text{LiBr})_{0.5}(\text{LiCl})_{0.5}$ (sample *s4*) as function of temperature (**Figure S2**), the cell parameter of $\text{Li}(\text{Br})_{1-x}(\text{Cl})_x$ solid solution has been determined. A cubic solid solution is formed at 250 °C, resulting stable down to *RT*, confirming the calculated phase diagram.¹ The cell parameter obtained by Rietveld refinement does follow the Vegard's law.



Figure S2. X-ray diffraction pattern as a function of temperature for sample s4. Symbols: blue circles CaO and green squares $Li(Br)_{1-x}(Cl)_x$.

Rietveld refinement of sample s1 after AN

Using the solution of equation (2) it was possible to refine the pattern. The sample after AN contains 14 wt.% of solid solution $LiBr_{0.28}Cl_{0.72}$ and 86 wt.% of h-Li(BH₄)_{0.39}Br_{0.33}Cl_{0.2}.



Figure S3. Results of the Rietveld refinement of XPD pattern of sample s1 after AN. (Rwp 11.66 %, χ 2.10).

HP-DSC signal of sample s2

The HP-DSC signal confirms that all $LiBH_4$ is in the hexagonal phase as solid solution, considering that the transition peak of $LiBH_4$ is not present.



Figure S4. HP-DSC signal of sample s2 after ball milling. The ramp used was form 30 °*C to 300* °*C at 5* °*C/min under 10 bar of H*₂.

Stability of the solid solution evaluated over time

The stability of the solid solution of sample s2 was evaluated over time, i.e. after about one-year. The pattern obtained after one year corresponds to that of the as-prepared sample.



Figure S5. X-ray diffraction pattern of sample s2 after AN preparation (bottom) and after one year (top). Symbols indicate the cubic and hexagonal solid solutions.

Effect of the milling time on the formation of the solid solution.

Different patterns were collected for different time of milling for the sample s3, with the same composition as the sample s2. t0 indicates the pattern of the hand-mixed sample.



Figure S6. Effect of the milling time on sample s3. Symbols: triangles LiBr, cross LiCl, circles h-Li(BH_4)_{1- α - β}(Br)_{α}(Cl)_{β}, squares Li(Br)_{1- ε}(Cl)_{ε}.

Density map as a function of composition

The density (ρ) of the samples was evaluated using the occupancy of the 2b site of h-LiBH₄, the weight of the anion and the obtained cell volume. Figure S7b shows conductivity at *RT* as a function of $1/\rho$. The samples highlighted have the best proprieties for possible technological applications.



Figure S7. (a) Contour map of density of samples in the hexagonal solid solution in the LiBH₄-LiBr-LiCl system as a function of composition. Iso-density dashed lines correspond of the values and lines shown in the legend. (b) plot of Li-ion conductivity at 50 °C with respect to the inverse of the density. The oval highlights the samples having a low density and, at the same time, a high conductivity.

Activation energy as a function of volume

Figure S8 shows the activation energy as a function of the obtained volume for h-Li(BH4)_{1- α -} $_{\beta}(Br)_{\alpha}(Cl)_{\beta}$ monophasic samples. The activation energy increases with volume decreases, indicating that also the dimension of the lattice is influence on the transport proprieties of the system.}



Figure S8. Activation energy of h-Li(BH4)_{1- α - β}(Br)_{α}(Cl)_{β} monophasic samples, as a function of the unit cell volume.

References

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