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Magnesium Ethylenediamine Borohydride as Solid-State Electrolyte for Magnesium Batteries

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Solid-state magnesium ion conductors with exceptionally high ionic conductivity at low temperatures, $5 \times 10^{-8} \text{ Scm}^{-1}$ at 30°C and $6 \times 10^{-5} \text{ Scm}^{-1}$ at 70°C , are prepared by mechanochemical reaction of magnesium borohydride and ethylenediamine. The coordination complexes are crystalline, support cycling in a potential window of 1.2V, and allow magnesium plating/stripping. While the electrochemical stability, limited by the ethylenediamine ligand, must be improved to reach competitive energy densities, our results demonstrate that partially chelated Mg^{2+} complexes represent a promising platform for the development of an all-solid-state magnesium battery.

Magnesium metal is an attractive anode material for rechargeable batteries, owing to its high volumetric capacity (3833 mAhcm^{-3} compared to 2036 mAhcm^{-3} for Li), negative redox potential of -2.36 V vs. SHE, and abundance. In contrast to metallic Li, Mg can be electroplated dendrite-free and promises consequently high energy density and safe operation¹⁻⁴. The development of effective Mg electrolytes is a crucial step towards the realization of a competitive reversible Mg battery and significant progress has been made particularly for liquid electrolytes, but safety and stability concerns remain^{2,5}.

Simple Mg electrolytes using solvents such as propylene carbonate and diethyl carbonate, analogous to typical Li electrolytes, do not allow for reversible electrodeposition of Mg metal, due to the formation of an ion blocking layer on the Mg surface⁵. Most reported Mg electrolytes are therefore based on organometallics, often Grignard reagents, or Mg salts dissolved in etheral solvents or glymes^{1,5,6}. In particular electrolytes based on magnesium borohydride, $\text{Mg}(\text{BH}_4)_2$, have received significant attention since 2012, when Mohtadi *et al.* demonstrated the first fully inorganic and halide-free Mg electrolytes enabling reversible Mg plating and stripping^{7,8}. In an effort to improve the safety, attempts have been made to replace the flammable organic solvents by ionic liquids⁹⁻¹¹.

By replacing the liquid electrolytes altogether, all-solid-state batteries employing solid-state electrolytes promise to be safer, *e.g.* in terms of heat and mechanical shock resistance. However, designing solid-state electrolyte materials with sufficiently high magnesium ion conductivity represents a major challenge owing to the divalent positive charge carried by the Mg^{2+} ion. Polymer-electrolyte systems, typically based on poly(ethylene oxide) (PEO) and Mg salts were considered as solid-state electrolytes, but the initially investigated salts $\text{Mg}(\text{SO}_3\text{CF}_3)_2$ or $\text{Mg}(\text{N}(\text{SO}_2\text{CF}_3)_2)_2$ are known to be incompatible with the Mg metal anode¹². In inorganic solids, Mg ion conductivity has been reported for few systems, but typically only at high temperatures, *e.g.* phosphate based polycrystalline solid Mg conductors, such as $\text{MgZr}_4(\text{PO}_4)_6$ and $\text{Mg}_{1.4}\text{Zr}_4\text{P}_6\text{O}_{24.4} + 0.4\text{Zr}_2\text{O}(\text{PO}_4)_2$ reach high conductivity above 10^{-5} Scm^{-1} at temperatures above 400°C ¹³⁻¹⁵.

$\text{Mg}(\text{BH}_4)_2$ derived materials, successfully used as liquid electrolytes thanks to the reductive stability of the BH_4^- anion, are attractive also as solid-state Mg ion conductors. However, the conductivity of pristine $\text{Mg}(\text{BH}_4)_2$ in the solid state is extremely low ($< 10^{-12} \text{ Scm}^{-1}$ at 30°C) which has been attributed to its structure, in which the Mg ions are located in firm tetrahedral cages composed of four BH_4^- units¹⁶. Higher ionic conductivity of $1 \times 10^{-6} \text{ Scm}^{-1}$ at 150°C and Mg plating/stripping was demonstrated for $\text{Mg}(\text{BH}_4)(\text{NH}_2)$ ¹⁷, in which Mg^{2+} is tetrahedrally coordinated by two BH_4^- and two NH_2^- . Reversible Mg plating/stripping was also demonstrated for PEO/ $\text{Mg}(\text{BH}_4)_2$ composites, but no values for the ionic conductivity were reported¹⁸.

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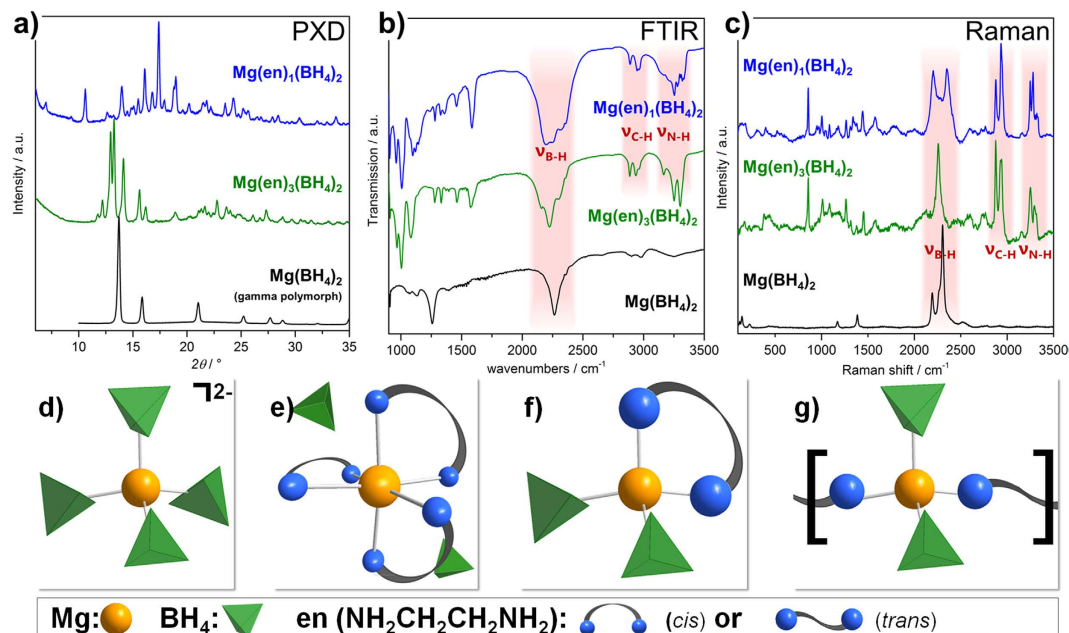
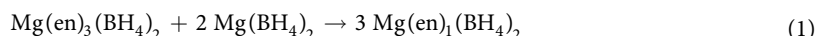


Figure 1. (a) Powder X-ray diffractograms, (b) FTIR and (c) Raman spectra of $\text{Mg}(\text{BH}_4)_2$, $\text{Mg}(\text{en})_3(\text{BH}_4)_2$, and $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ along with a conceptual sketch of the local coordination of Mg^{2+} in (d) $\text{Mg}(\text{BH}_4)_2$, (e) $\text{Mg}(\text{en})_3(\text{BH}_4)_2$, and two likely possibilities in $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ (f,g).

Here we report that high ionic conductivity can be achieved in $\text{Mg}(\text{BH}_4)_2$ based materials already near room temperature by coordinating Mg^{2+} by a neutral bidentate ethylenediamine ligand ($\text{NH}_2\text{CH}_2\text{CH}_2\text{NH}_2$, abbreviated as en). Using only one bidentate ligand per metal atom leads to a partially chelated, mixed coordination of Mg^{2+} , which renders it mobile. Our approach is compatible with Mg plating/stripping and could be extended to other ligands and Mg salts, widening the portfolio of electrolytes for the development of an all-solid-state Mg battery.

The synthesis of the highly conducting $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ proceeds in two steps: First, ball milling of $\text{Mg}(\text{BH}_4)_2$ with three equivalents of en produces a $\text{Mg}(\text{en})_3(\text{BH}_4)_2$ complex. The synthesis, structure, and thermal stability of this compound have been reported recently¹⁹. In order to have good control over the stoichiometry, ca. 3.2 equivalents of en were used during ball milling and excess en was removed under vacuum at elevated temperatures, see the experimental section in the Supplementary Information for details.

In a second step, the produced $\text{Mg}(\text{en})_3(\text{BH}_4)_2$ was ball-milled with $\text{Mg}(\text{BH}_4)_2$ according to reaction 1 to produce $\text{Mg}(\text{en})_1(\text{BH}_4)_2$.



Other ratios of $\text{Mg}(\text{en})_3(\text{BH}_4)_2$ and $\text{Mg}(\text{BH}_4)_2$ were also investigated, however the product with the nominal chemical formula $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ showed the highest ionic conductivity.

Powder X-ray diffraction (PXD) shows that the synthesized coordination complex is a polycrystalline solid. Figure 1a compares the diffractograms of $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ to those of the reactants $\text{Mg}(\text{BH}_4)_2$ and $\text{Mg}(\text{en})_3(\text{BH}_4)_2$. From comparison of the Bragg reflections, we conclude that $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ contains no remaining traces of the reactants $\text{Mg}(\text{en})_3(\text{BH}_4)_2$ and $\text{Mg}(\text{BH}_4)_2$. The first reflection for $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ is observed at $q = 0.49 \text{ \AA}^{-1}$, indicative of a large unit cell. Indexing of the PXD did not yield a plausible unit cell. Single crystals would facilitate structural analysis, but are not available.

$\text{Mg}(\text{en})_1(\text{BH}_4)_2$ is thermally stable up to 75°C , where a structural phase transition occurs as observed by X-ray diffraction (Figure S1a). The transition is accompanied by an endothermic event observed by differential scanning calorimetry (Figure S1b). A slight weight loss, measured by thermal gravimetry, associated with the decomposition of $\text{Mg}(\text{en})_1(\text{BH}_4)_2$, begins above 100°C (Figure S1b). The thermal stability of $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ up to 75°C is thus comparable to the thermal stability of liquid electrolytes used in standard Li ion batteries.

To shed light onto the local structure and coordination of $\text{Mg}(\text{en})_1(\text{BH}_4)_2$, we used Fourier transform infrared (FTIR) and Raman spectroscopy (Fig. 1b,c). We first discuss the coordination of Mg^{2+} in $\text{Mg}(\text{en})_3(\text{BH}_4)_2$ for which a crystal structure model has been reported¹⁹. A conceptual sketch of the local coordination is shown in Fig. 1e. Mg^{2+} is octahedrally coordinated by three chelating bidentate en ligands, resulting in an effectively larger cation $[\text{Mg}(\text{en})_3]^{2+}$ which is charge balanced by the BH_4^- anions.

Going from $\text{Mg}(\text{en})_3(\text{BH}_4)_2$ to $\text{Mg}(\text{en})_1(\text{BH}_4)_2$, the number of en ligands per Mg^{2+} ion reduces from three to one. Consequently, Mg^{2+} is necessarily forced into a mixed coordination with both en and BH_4^- in the first coordination sphere. In $\text{Mg}(\text{en})_1(\text{BH}_4)_2$, two likely possibilities exist for the coordination of Mg^{2+} by the en ligand and BH_4^- , as sketched in Fig. 1f and g: (i) A chelating coordination of en as in $\text{Mg}(\text{en})_3(\text{BH}_4)_2$, whereby the en ligand is non-bridging in the lower symmetry cis configuration as in Fig. 1f, and the complex could be formulated as

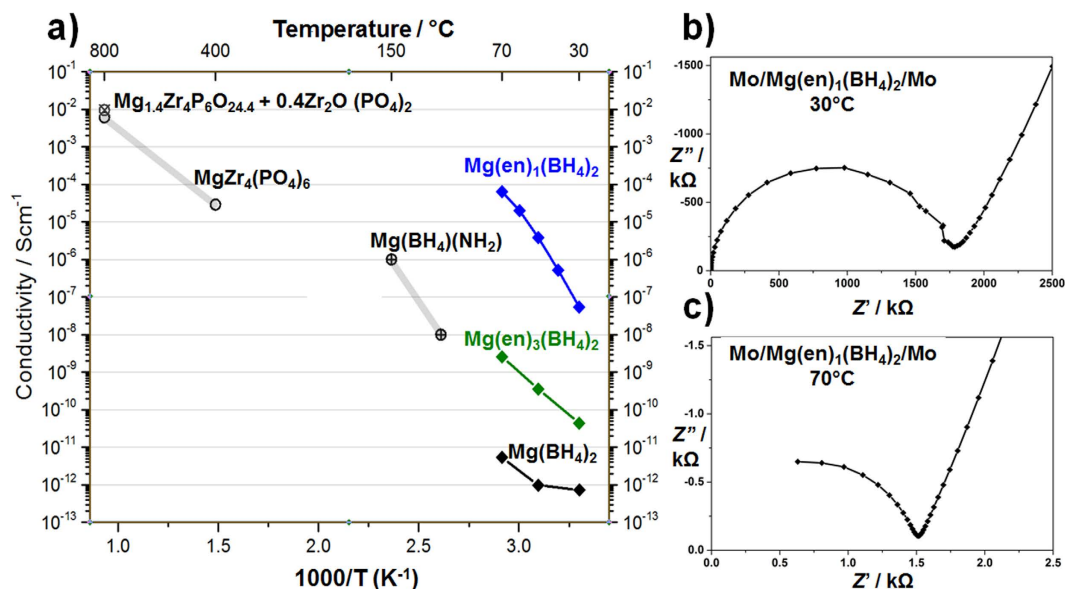


Figure 2. (a) Temperature dependence of ionic conductivity of Mg(BH₄)₂, Mg(en)₃(BH₄)₂ and Mg(en)₁(BH₄)₂, together with literature data for MgZr₄(PO₄)₆¹³, Mg_{1.4}Zr₄P₆O_{24.4} + 0.4Zr₂O(PO₄)₂^{14,15} and Mg(BH₄)(NH₂)¹⁷. (b,c) Nyquist plots of Mo/Mg(en)₁(BH₄)₂/Mo at 30 and 70 °C, respectively.

[Mg(en)(BH₄)₂] or ii) a complex in which en acts as a bridging ligand between different Mg²⁺ ions as in Fig. 1g, thus being in the centrosymmetric trans configuration and forming chains, formulated as [-Mg(BH₄)₂-en-Mg(BH₄)₂-en-].

Evidence favoring the chelating cis configuration comes from FTIR and Raman spectroscopy. The spectra shown in Fig. 1c and d of the starting materials and Mg(en)₁(BH₄)₂ differ significantly. The peak positions are summarized in Table S1 and the descriptions are based on literature assignments of a number of transition metal ethylenediamine complexes studied by vibrational spectroscopy supported by N-deuteration^{20–23}. The FTIR spectrum of Mg(BH₄)₂ is characterized mainly by B-H stretching at 2266 cm⁻¹, and bending at 1259, 1130 and 1070 cm⁻¹. The B-H stretching band splits into several peaks in the FTIR and Raman spectra of Mg(en)₃(BH₄)₂ and Mg(en)₁(BH₄)₂, indicative of the presence of several different types of B-H bonds in the crystal structures, in agreement with four different BH₄ sites in Mg(en)₃(BH₄)₂. When comparing the spectra of Mg(en)₃(BH₄)₂ and Mg(en)₁(BH₄)₂, the bands assigned to the C-N and C-H backbone of the en ligand in Mg(en)₃(BH₄)₂ and Mg(en)₁(BH₄)₂, remain almost unchanged, namely C-H stretching at 2888 ± 1, 2941 ± 4 and 2963 ± 1 cm⁻¹, C-H bending at 1460 ± 2 cm⁻¹, CH₂ twist at 1279 ± 1 cm⁻¹ and C-N and C-C stretching at 1094 ± 3 cm⁻¹ and 1005 ± 1 cm⁻¹, respectively. Additional peaks are present in the spectrum for Mg(en)₁(BH₄)₂ at 3320, 3280 and 3205 cm⁻¹ in the N-H stretching region as well as between 1240 and 1120 cm⁻¹. A bridging en ligand in trans C_{2h} configuration results in higher symmetry and thus less peaks in the spectra²¹. This is not observed for the Raman and FTIR spectra of Mg(en)₁(BH₄)₂, where the same peaks are present as in Mg(en)₃(BH₄)₂, evidencing a chelating coordination of en around the Mg ion, as shown in Fig. 1f.

The temperature dependence of the ionic conductivity of Mg(BH₄)₂, Mg(en)₃(BH₄)₂ and Mg(en)₁(BH₄)₂ was determined by electrochemical impedance spectroscopy (EIS) in the temperature range between room temperature and 70 °C in symmetric cells with Mg-blocking molybdenum electrodes, Mo/Mg(en)_x(BH₄)₂/Mo (Fig. 2a). Typical Nyquist plots are presented in Fig. 2b and c for Mg(en)₁(BH₄)₂ at 30 and 70 °C, respectively and show standard ionic conduction behavior with a single semicircle. The ionic conductivity was extracted from the higher frequency intercept of the extrapolated semicircle with the horizontal axis, and taking into account the geometry of the sample. Ionic conductivities measured as a function of temperature are summarized in Fig. 2a for Mg(BH₄)₂, Mg(en)₃(BH₄)₂ and Mg(en)₁(BH₄)₂, together with selected literature data for Mg(BH₄)(NH₂), MgZr₄(PO₄)₆ and Mg_{1.4}Zr₄P₆O_{24.4} + 0.4Zr₂O(PO₄)₂.

The ionic conductivity of Mg(en)₁(BH₄)₂ is 5 × 10⁻⁸ Scm⁻¹ at 30 °C, exceeding that of Mg(BH₄)₂ by more than four and that of Mg(en)₃(BH₄)₂ by three orders of magnitude and further increases to above 6 × 10⁻⁵ Scm⁻¹ at 70 °C. This is so far the highest conductivity measured for solid-state Mg conductors in this temperature range. The temperature dependence of the conductivity of Mg(en)₃(BH₄)₂ and Mg(en)₁(BH₄)₂ shows Arrhenius behavior from 30 to 70 °C, with an apparent activation energy of 0.9 and 1.6 eV, respectively. These activation energy values are relatively high but comparable to the other Mg solid-state electrolyte systems reported in Fig. 2a.

Our impedance results show that introducing the chelating en ligand significantly improves the mobility of the Mg²⁺ ion. Of fundamental importance is the reduction of the number of en ligands coordinating the Mg²⁺ cation from three in Mg(en)₃(BH₄)₂ to one in Mg(en)₁(BH₄)₂, leading to an asymmetric mixed coordination of Mg²⁺ by one en and two BH₄⁻. A related observation was also made for liquid electrolytes, in which Mg(BH₄)₂ is solvated by glymes, and where cations which were only partially solvated by the solvent and partially coordinated by the

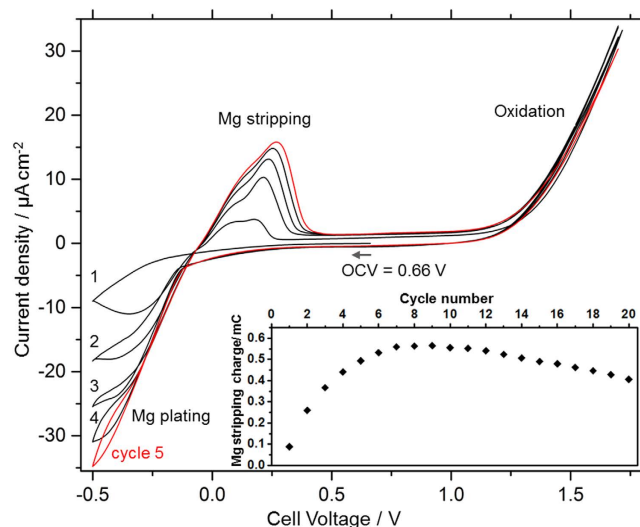


Figure 3. Cyclic voltammogram of a Pt/Mg(en)₁(BH₄)₂/Mg cell at 60 °C at a scan rate of 10 mVs⁻¹. Inset: Mg stripping charge (=area of the Mg stripping peak) vs. cycle number.

anion were found to be more mobile¹⁸. However, the findings from liquid chelated systems cannot be translated directly to the solid, since in liquid electrolytes the ion moves with its solvation shell.

Interestingly, impedance measurements on the related halide complexes Mg(en)₁X₂ (X = I, Cl) (not shown here) show much lower ionic conductivity, indicating that the lower symmetry and rotational degrees of freedom of the tetrahedral BH₄⁻ anions play an important role in the conduction mechanism. To understand this in more detail, the crystal structure of Mg(en)₁(BH₄)₂ will need to be determined.

High ionic conductivity is only one of the requirements for an electrolyte for Mg batteries. To test the stability and to confirm that the observed conductivity stems from the transport of Mg ions, cyclic voltammetry measurements were conducted on an asymmetric Pt/Mg(en)₁(BH₄)₂/Mg cell at 60 °C, as shown in Fig. 3. At -0.2 V vs. the Mg counter electrode, we observe the onset of cathodic current corresponding to Mg plating onto the Pt working electrode. The anodic current during the reverse sweep between -0.2 V and 0.5 V is attributed to Mg stripping, strongly indicating that Mg conduction indeed takes place.

For voltages >1.2 V, we observe irreversible oxidation. The interface layer formed during this irreversible oxidation is able to conduct Mg ions, since plating/stripping is also observed in the following cycles. Indeed, the plating/stripping currents increase during the first seven cycles (see Fig. 3 cycles and inset), suggesting that an initialization period improves the interfacial contact between electrodes and electrolyte

To provide direct evidence for Mg²⁺ ion conduction through the electrolyte, we assembled a Cu/Mg(en)₁(BH₄)₂/Mg cell and deposited Mg on the Cu electrode at 60 °C. The cell was subsequently disassembled and the surface of the Cu electrode examined by scanning electron microscopy (SEM) and energy dispersive X-ray (EDX) analysis. The SEM image of the Cu electrode in Figure S3 shows randomly distributed patches, identified as Mg by EDX (also shown in Figure S3), which confirm Mg plating. No such patches were observed for an identically built reference cell to which no plating current was applied (Figure S4). The patchy deposition of Mg provides additional evidence that electrolyte and Cu electrode are not in contact over the full area, but that the contact area increases upon plating, as the plated Mg fills the gaps between electrolyte and electrode. Also for liquid electrolyte systems it has been observed that newly formed interfaces between Mg metal and electrolyte improve the performance²⁴.

In this work, we demonstrate that partially chelated ethylenediamine magnesium borohydride complexes are promising new solid-state Mg conductors with high conductivity of up to $5 \times 10^{-8} \text{ Scm}^{-1}$ at 30 °C and $6 \times 10^{-5} \text{ Scm}^{-1}$ at 70 °C, tackling the challenge of mobility of the divalent Mg²⁺ ion in the solid state. The compounds are easily synthesized by ball milling, no sintering or other energy expensive processes were necessary. While Mg plating/stripping proved feasible, more studies are necessary to understand the mechanism of ionic conductivity on the atomic level in these systems and the interface dynamics. Our approach using partially chelated Mg complexes with a complex anion as solid-state Mg electrolytes can be extended to other ligands and anions, allowing for the tuning of the electrochemical stability. This could open the doors for the development of all solid-state high energy density Mg batteries, conceivably a game-changer in battery technology.

Methods

Sample Synthesis. The synthesis of Mg(en)₃(BH₄)₂ was carried out according to a modified literature procedure¹⁹. Mg(BH₄)₂ was ball-milled with approximately 3.2 equivalents ethylenediamine (en) in a Spex 8000 shaker mill under argon atmosphere for 3×10 min, with 5 min break in between, using a balls-to-sample mass ratio of 10 to 1. An excess of en with respect to the molar ratio for Mg(en)₃(BH₄)₂ was used to ensure a complete reaction. The excess en was removed from Mg(en)₃(BH₄)₂ under dynamic vacuum at 120 °C (1 h). For the synthesis of Mg(en)₁(BH₄)₂, the produced Mg(en)₃(BH₄)₂ was ball-milled with additional Mg(BH₄)₂ in the correct stoichiometric ratio, using the same ball milling procedure.

The chemical formula of $\text{Mg}(\text{en})_1(\text{BH}_4)_2$ and the stoichiometric reaction according to (1) is corroborated by temperature dependent PXD between 30 and 70 °C, which shows the simultaneous decrease of the intensity of reflections of both reactants, while the new compound is formed (see Figure S2). In the same temperature range, thermogravimetric analysis of the reaction mixture records no weight loss.

All preparations and manipulations of samples were performed in a glove box with a circulation purifier (argon atmosphere, <1 ppm of O_2 and H_2O). The chemicals magnesium borohydride $\text{Mg}(\text{BH}_4)_2$ (Aldrich, 95%) and ethylenediamine (Aldrich, 99.5%) were used as received.

Structural Characterization. Powder X-ray diffraction (PXD) measurements were performed on a Bruker D8 Advance diffractometer equipped with a Goebel mirror selecting $\text{Cu K}\alpha$ radiation ($\lambda = 1.5418 \text{ \AA}$) and a linear detector system (Vantec). Samples were filled and sealed into glass capillaries under argon atmosphere. Simultaneous differential scanning calorimetry and thermogravimetry measurements were performed using a Netzsch STA 449 F3 Jupiter with Al sample pans at $\Delta T/\Delta t = 5 \text{ Kmin}^{-1}$ between 30 and 400 °C under He flow at 80 ml min^{-1} .

FTIR spectra were obtained between 900 and 4000 cm^{-1} with a Bruker Vektor 20 spectrophotometer and a Golden Gate ATR cell in atmospheric conditions, causing short air exposure during the measurements. Raman measurements were performed on samples packed in glass capillaries between 100 and 4000 cm^{-1} , using a Renishaw Raman system equipped with a HeNe Laser, $\lambda = 633 \text{ nm}$.

Electrochemical Characterization. Electrolyte pellets were pressed using a pressing mold with an inner diameter of 12 mm. The material was weighed and loaded into the mold and pressed using a knuckle joint press inside the glovebox. The thickness of the resulting pellet was determined with a caliper (between 0.5 and 0.8 mm). Electrochemical measurements were performed in 2-electrode stainless steel cells, which were assembled inside an argon filled glovebox. A spring was used to maintain mechanical contact between electrodes and electrolyte.

Electrochemical impedance spectroscopy (EIS) was measured using a Zahner IM6ex electrochemical workstation in the frequency range $3 \times 10^6 - 0.1 \text{ Hz}$. Polished molybdenum blocks were used as Mg blocking electrodes. Impedance spectra were obtained at 30, 40, 50, 60 and 70 °C, after thermal equilibration for 2 h at each temperature.

Cyclic voltammetry (CV) measurements were performed at 60 °C after thermal equilibration for 4 h with a Biologic VMP3 electrochemical workstation. An asymmetric cell was prepared with Mg foil (99.95% Mg, 11 mm diameter, 0.05 mm thickness, GalliumSource) as counter electrode and Pt as working electrode. The OCV stabilized at 0.66 V and cyclic voltammetry was carried out between -0.5 and 1.7 V with a scan rate of 10 mVs^{-1} . A similar cell, but with a Cu working electrode was built. Mg was plated at a current of 3 nA for ca. 200 h at overpotentials below -0.1 V at 60 °C. SEM images were taken with a FEI NanoSEM 230 at 25 kV. EDX analysis was performed with an Oxford X-Max SDD EDX system.

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Author Contributions

E.R. und A.R. initiated the study. E.R. prepared the samples and performed XRD and vibrational spectroscopy experiments and analyzed the data. E.R. and R.-S.K. performed electrochemical experiments and analyzed the data. All authors discussed and interpreted the data. The manuscript was written by E.R., A.R. and C.B. All authors discussed and commented on the manuscript.

Additional Information

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