Synthesis, structure, and polymorphic transitions of praseodymium(III) and neodymium(III) borohydride, Pr(BH₄)₃ and Nd(BH₄)₃

SeyedHosein Payandeh GharibDoust,* Michael Heere, Carlo Nervi, Magnus H. Sørby, Bjørn C. Hauback and Torben R. Jensen*

In this work, praseodymium(III) borohydride, Pr(BH₄)₃, and an isotopically enriched analogue, Pr(¹⁴BD₄)₃, are prepared by a new route via a solvate complex, Pr(¹⁴BD₄)₃S(CH₃)₂.

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1. Introduction

One of the greatest challenges in the 21st century is the transition towards a sustainable and environmentally friendly energy system.\(^1\)\(^-\)\(^2\) Hydrogen is the most abundant and lightest element and has the highest gravimetric energy density for any known substance. Hydrogen is therefore an ideal (renewable) energy carrier.\(^3\)\(^-\)\(^5\) Metal borohydrides have received considerable attention as potential hydrogen storage materials due to their high hydrogen densities.\(^6\)\(^-\)\(^7\) A wide variety of metal borohydrides have been synthesized and characterized, and an inverse correlation has been found between the decomposition temperature of the metal borohydride and the Pauling electronegativity of the metal.\(^7\)\(^-\)\(^8\) In recent years, rare-earth (RE) metal borohydrides, in particular, have received considerable interest due to their various properties.\(^9\)\(^-\)\(^14\) RE(BH\(_4\))\(_3\)(THF)\(_2\) (RE = Eu and Yb, THF = tetrahydrofuran) have shown high luminescence properties,\(^15\) and other lanthanide borohydride complexes such as RE(BH\(_4\))\(_3\)(THF)\(_3\) (RE = Nd, Sm) have been used extensively for the catalysis of organic polymerization reactions.\(^16\)\(^-\)\(^18\) These compounds have a higher solubility in apolar solvents compared to that of traditional RECl\(_3\)(THF)\(_3\) compounds, which makes them promising candidates as catalysts.\(^16\)\(^-\)\(^18\) The hydrogen capacities of rare-earth metal borohydrides, RE(BH\(_4\))\(_n\), \(n = 2, 3\), vary between 9.07 wt% H\(_2\) for Y(BH\(_4\))\(_3\) to 5.56 wt% for Yb(BH\(_4\))\(_3\),\(^19\)\(^-\)\(^24\) and are within the range of MgH\(_2\) (7.65 wt% H\(_2\)) and NaAlH\(_4\) (7.46 wt% H\(_2\)), which have previously received considerable attention.\(^25\)\(^-\)\(^26\) Bimetallic rare-earth borohydrides such as MY(BH\(_4\))\(_3\), M = Li, Na, and LiRE(BH\(_4\))Cl, RE = La, Ce, Gd have moderate to high Li and Na ion conductivity.\(^12\)\(^-\)\(^13\)\(^-\)\(^27\)\(^-\)\(^31\) Magnetocaloric properties have also been observed for K\(_2\)Gd(BH\(_4\))\(_5\) and Cs\(_3\)Gd(BH\(_4\))\(_6\).\(^32\)
The crystal chemistry of rare-earth metal borohydrides, \( \text{RE(BH}_4\text{)}_3 \), is diverse and interesting.\(^5\)\(^,\)\(^{23}\) Metal borohydrides with the largest cations, \( \text{RE} = \text{La, Ce} \), crystallize in the trigonal crystal system with space group \( R\bar{3}c \) \((r\text{-RE(BH}_4\text{)}_3)\).\(^{13}\)\(^,\)\(^{34}\) Smaller cations, e.g., \( \text{Pr} \) to \( \text{Yb} \), crystallize in the cubic crystal system with space group \( \text{Pa}\bar{3} \) \((\alpha\text{-RE(BH}_4\text{)}_3)\).\(^{19}\)\(^,\)\(^{23}\)\(^,\)\(^{35}\) Most of the trivalent rare-earth borohydrides undergo a transition to a second cubic polymorph with space group symmetry either \( P\bar{m}\bar{3}m \) or \( Pm\bar{3}c \) at elevated temperatures \((\beta\text{-RE(BH}_4\text{)}_3)\).\(^{19}\)\(^,\)\(^{23}\)\(^,\)\(^{35}\)\(^,\)\(^{39}\) All known RE \((\text{RE} = \text{Pr})\) with space group symmetry \( \text{Pa}\bar{3}, R\bar{3}c \) or \( P\bar{m}\bar{3}c \) have crystal structures related to polymorphs of rhenium trioxide, \( \text{ReO}_3\).\(^{33}\)\(^,\)\(^{34}\) Some rare-earth metals form stable borohydrides in oxidation state\(\text{II} \), e.g., \( \text{Sm}^{2+} \) and \( \text{Eu}^{2+} \), which crystallize in the orthorhombic crystal system \((\text{space group } Pbcn)\) at RT. Upon heating, europium \((\text{Eu})\) borohydride, \( \text{Eu(BH}_4\text{)}_3 \), transforms into tetragonal and cubic polymorphs with space group symmetry \( \text{either } P4_12_12 \text{ or } Pm\bar{3}m \).\(^21\)\(^,\)\(^{24}\)

Previous methods for the synthesis of trivalent rare-earth metal borohydrides were based on the reaction of \( \text{LiBH}_4 \) with \( \text{RECl}_3 \),\(^{23}\)\(^,\)\(^{34}\) but the products obtained using this method were contaminated with amorphous \( \text{LiBH}_4 \). The presence of amorphous \( \text{LiBH}_4 \) has consequences for thermal properties and chemical reactivity of samples prepared by this method. For example, \( \text{La(BH}_4\text{)}_3 \) synthesized according to this method formed \( \text{Li}_3\text{K}_3\text{La}_2\text{(BH}_4\text{)}_{12} \) upon reaction with \( \text{KBH}_4 \).\(^{13}\) Therefore, our focus has been on new synthetic strategies and investigation of chemical, physical, and structural properties of the pure compounds. Here we present a new method to obtain solvate complexes, \( \text{RE(BH}_4\text{)}_3\text{S(CH}_3\text{)}_2 \), \( \text{RE} = \text{Pr, Nd} \), and the corresponding borohydrides, \( \text{Pr(BH}_4\text{)}_3 \) and \( \text{Nd(BH}_4\text{)}_3 \), which allows detailed investigation of the polymorphic transformations.

2. Experimental

2.1 Sample preparation

\( \text{Pr(BH}_4\text{)}_3 \) and \( \text{Nd(BH}_4\text{)}_3 \) compounds were synthesized using a new method based on the reaction of \( \text{Pr}_2 \) or \( \text{Nd}_2 \) with a dimethyl sulfide borane complex, \( \text{S(CH}_3\text{)}_2\text{BH}_3 \) (DMS). \( \text{RE} \) metal ingots were scrapped with sand paper to remove the oxide layer, placed in an autoclave, and heated up to 380 °C for 2 hours under hydrogen pressure \((\text{H}_2) = 100 \text{ bar}\) to form \( \text{REH}_3 \) \((\text{RE} = \text{Pr, Nd})\). The formation of \( \text{REH}_3 \) was confirmed by X-ray powder diffraction (XRPD). However, the hydrogenation was repeated for the second time in order to prevent contamination of the sample with the \( \text{RE} \) metal. In the next step, the \( \text{REH}_3 \) \((\text{RE} = \text{Pr, Nd}) \) powder was ball milled for two hours under an argon atmosphere \((10 \text{ min milling, 2 min break, 12 repetitions}) \) with a powder-to-ball ratio of 1 : 20. A Fritsch Pulverisette 6 planetary ball mill was used, which was equipped with a tungsten carbide vial \((80 \text{ mL}) \) and balls \((\text{o.d. } 10 \text{ mm})\).

The ball-milled powder was transferred to a reaction flask and mixed with \( \text{S(CH}_3\text{)}_2\text{BH}_3 \) in a toluene solvent \((10 \text{ M}) \) at a molar ratio of 1 : 4.5. The excess of the \( \text{S(CH}_3\text{)}_2\text{BH}_3 \) solvent was used to completely consume the \( \text{REH}_3 \), and the solution was diluted to half-concentration by adding the same volume of toluene. The mixture was stirred for 2 days at 45 °C. Then the temperature was decreased to 35 °C and dimethyl sulfide was added to the mixture \((50 \text{ mL per gram of } \text{REH}_3\) in order to dissolve \( \text{RE(BH}_4\text{)}_3\text{S(CH}_3\text{)}_2 \), allowing the removal of possible remaining \( \text{REH}_3 \) in the sample. After one day of stirring at 35 °C, the solution was filtered, and the solvent was removed using a rotary evaporator at 70 °C.

At this stage, \( \text{RE(BH}_4\text{)}_3\text{S(CH}_3\text{)}_2 \) \((\text{RE} = \text{Pr, Nd})\) was obtained. In order to remove the coordinated solvent, the powder was transferred to a glass tube and annealed at 180 °C under vacuum for 1 h. The sample was then ground with a mortar and pestle in a glovebox. The heat treatment was repeated once more in order to completely remove the DMS solvent from the sample.

An isotopically enriched sample of praseodymium \((\text{iii}) \) borohydride, \( \text{Pr}^{(11}\text{BD}_3) \), was prepared using the same procedure with different reagents. For deuteration, the \( \text{Pr} \) ingots were transferred to an autoclave placed in liquid nitrogen and deuterium, \( \text{D}_2 \), gas was added \((p(\text{D}_2) = 8 \text{ bar}) \) and the pressure increased to \( p(\text{D}_2) = 14 \text{ bar} \) upon heating the autoclave from room temperature \((\text{RT})\). The autoclave was then heated further to 390 °C for 2 h. The deuteration procedure was repeated three times in order to fully convert the \( \text{Pr} \) metal to \( \text{PrD}_3 \). The solvated praseodymium \((\text{iii}) \) borohydride was prepared using \( \text{S(CH}_3\text{)}_2^{11}\text{BD}_3 \) and using the same procedures as described above.

\( \text{LiBH}_4 \) \((95\%), \text{ toluene, } C_6H_5CH_3 \) \((\text{anhydrous, } 99.9\%) \), dimethyl sulfide, DMS, \( \text{S(CH}_3\text{)}_2 \) \((\text{anhydrous, } 99.9\%) \), and praseodymium, \( \text{Pr} \) \((99.9\%) \), and neodymium, \( \text{Nd} \) \((99.9\%) \), ingots were purchased from Sigma-Aldrich. \( \text{S(CH}_3\text{)}_2\text{BH}_3 \) and \( \text{S(CH}_3\text{)}_2^{11}\text{BD}_3 \) \((10.0 \text{ M in toluene}) \) were purchased from Katchem. All chemicals were used as received and sample manipulation was performed in an argon-filled glove box with a circulation purifier, \((O_2, H_2O) < 0.5 \text{ ppm})\).

2.2 In situ synchrotron radiation X-ray powder diffraction

\( \text{In situ} \) synchrotron radiation X-ray powder diffraction (SR-XRPD) data of the samples were collected at the Swiss-Norwegian Beam Line \((\text{SNBL, BM01A}) \) at ESRF, Grenoble, France, and at the P02 beamline at Petra III, Desy, Hamburg, Germany. At SNBL, a Dexcela-PerkinElmer 2923 CMOS pixel detector\(^{40}\) was used, and the wavelength was 0.7129 Å. At Petra III, a PerkinElmer XRD1621 detector was used, and the selected X-ray wavelength was 0.2072 Å. The samples were mounted in boron silicate capillaries \((\text{o.d. } 0.5 \text{ mm}) \) under argon and sealed with glue to prevent contact with air. For the \( \text{in situ} \) XRPD experiment under pressure which was performed in ESRF, the sample was loaded in a quartz capillary \((\text{o.d. } 0.5 \text{ mm}) \) and attached to a specially designed sample holder from ESRF, and hydrogen gas was applied, \( p(\text{H}_2) = 40 \text{ bar}) \). For the high-pressure experiments performed in the P02 beamline at Petra III, samples were packed in sapphire tubes \((\text{i.d. } 0.8 \text{ mm}) \) and attached to a specially designed sample cell for studying solid–gas reactions, and hydrogen gas was applied, \( p(\text{H}_2) = 98 \text{ bar} \).
2.3 Neutron powder diffraction

Neutron powder diffraction (NPD) data of the Pr(11BD4)3 sample were collected at two different places, the spallation neutron source SINQ at the Paul Scherrer Institute in Villigen, Switzerland and the JEEP II reactor at the Institute for Energy Technology in Kjeller, Norway. At SINQ, NPD data were collected at a high-resolution powder neutron diffractometer (HRPT) from 2θ = 10° to 130°.43 The powder sample was enclosed in a vanadium can (8 mm diameter), and data were collected with a wavelength of λ = 1.494 Å. At the JEEP II reactor, data were collected using a PUS instrument and neutrons with λ = 1.5583 Å were obtained from a Ge (511) focusing monochromator. Data were collected from 10° to 130° in steps of 0.05° in 2θ.44

2.4 Structure solution and refinement

A new solvate compound and new polymorphs of Pr(BH4)3, rhombohedral r-Pr(BH4)3 (R3c) and cubic β-Pr(BH4)3 (Fm3c), were identified and structurally characterized in this investigation, see Table 1 and Tables S2, S3.† Moreover, a new high-temperature polymorph of Nd(BH4)3 (Fm3c) was identified. Structural models for the new compounds were derived from isostructurally known compounds as discussed in the next sections. The structure models were refined using X-ray and neutron powder diffraction data by the Rietveld method implemented in the program Fullprof.45 Data collected for samples s1 were used to refine the structural models of Pr(11BD4)3S(CH3)2 and r-Pr(11BD4)3, resulting in Rwp = 6.43 and 4.66% (not corrected for background), respectively (Fig. S7 and S6†). In order to confirm the crystal structure of r-Pr(BH4)3, and to obtain the exact position of H atoms, Pr(11BD4)3 was synthesized, and NPD of sample s2 was used to refine the structure model with Rwp = 3.95% (not corrected for background), χ2 = 4.12 (Fig. S11†). Moreover, the crystal structure of Pr(11BD4)3S(CH3)2 was confirmed by Rietveld refinement based on NPD data with Rwp = 2.00% (not corrected for background), χ2 = 2.18 (Fig. S9†). NPD data of Pr(BD4)3S(CH3)2 and r-Pr(BD4)3 were collected from 2θ 10° to 130°; however, Rietveld refinements of these structures were performed in the 2θ range of 10° to 110° and 10° to 70°, respectively. This was due to the low quality of the neutron diffraction data at high 2θ angles.

The crystal structure of β-Pr(BH4)3 and β-Nd(BH4)3 was refined based on the XRPD pattern of samples s3 and s5 resulting in Rwp = 2.22% and 4.59% and χ2 = 301 and 5970, respectively (see Fig. S5 and S12†). A high χ2 value is due to extremely high counting statistics of the powder diffraction data obtained from modern 2D detectors. The crystal structures obtained by powder X-ray diffraction were refined based on the fixed positions of the elements in the (known) structure models. Therefore, H–H or D–D distances in these data were obtained by considering the original coordinates of the H/D atoms. The structural model of Y(BH4)3 that was used to refine the structure of the two polymorphs (Pa3 and Fm3c) had been optimized by DFT and the H/D coordinates are considered reliable.37 The R3c structural model for La(BH4)3 was not optimized by DFT;34 however, the original coordinates of the H/D atoms were used for refinement. In the next step, the H–H/D–D distances were compared to the refinement based on NPD data. In the NPD data refinement, the coordinates of D atoms were refined and for this purpose, additional strains were added to the structures, and B–D and D–D distances were fixed at 1.19 and 1.90 Å, respectively. The D–D distances obtained by NPD were in agreement with the values obtained by XRPD. The backgrounds were described by linear interpolation between selected points. Unit cell parameters, scale factors, zero-point, peak shape mixing parameters (pseudo-Voigt function), and three profile parameters (U, V, W) were refined. The crystallographic data of the compounds are provided in Table 2.

2.5 Thermal analysis and mass spectrometry

The thermal behavior of the samples was studied by combined thermogravimetric analysis (TGA), differential scanning calorimetry (DSC), and mass spectrometry (MS). DSC and TGA were carried out using a PerkinElmer STA 6000 apparatus, which was attached to a Hiden Analytical HPR-20 QMS mass spectrometer. Approximately 10 mg of the sample was placed in an Al2O3 crucible and heated from 40 to 500 °C (ΔT/Δt = 5 °C min⁻¹) in an argon flow of 40 mL min⁻¹. The evolved gases were transported to a MS and analyzed for H2 (m/z = 2) and B2H6 (m/z = 27).

2.6 DFT calculations

Quantum Espresso version 6.146 was employed for DFT periodic lattice calculations using the Generalized Gradient Approximation (GGA) functional PW86PBE,47,48 with the

Table 1 Overview of investigated samples, molar ratio of the used reactants, synthesis procedure, and the sample composition determined by Rietveld refinement of XRPD or NPD data, the latter indicated by (*)

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3. Results and discussion

3.1 Synthesis

A new approach to synthesize pure rare-earth metal borohydrides is presented using metal hydrides, REH₃, RE = Pr, Nd, and a borane donating complex, S(CH₃)₂BH₃ in solution, forming solvated complexes, see reaction (1).

\[
\text{REH}_3(s) + 3\text{S(CH}_3)_2\text{BH}_3(\text{toluene}) \rightarrow \text{RE(BH}_4)_3\text{S(CH}_3)_2(s) + 2\text{S(CH}_3)_2(\text{toluene})
\]

Using this method, Pr(BH₄)₃S(CH₃)₂ and Nd(BH₄)₃S(CH₃)₂ are synthesized and Pr(BH₄)₃ and Nd(BH₄)₃ are obtained by heating the solvate complexes under vacuum at 180 °C, see reaction (2).

\[
\text{RE(BH}_4)_3\text{S(CH}_3)_2(s) \rightarrow \alpha\text{-RE(BH}_4)_3\text{S(CH}_3)_2(s) + \text{S(CH}_3)_2(g), \quad \text{RE = Pr, Nd}
\]

Pr(¹¹BD₄)₃S(CH₃)₂ and Pr(¹¹BD₄)₃ were synthesized using a similar approach, reaction (1) and (2), and PrD₃ and S(CH₃)₂¹¹BD₃ as reactants.

3.2. Structure analysis

3.2.1. Crystal structure of RE(BH₄)₃S(CH₃)₂, RE = Pr, Nd.

The diffraction patterns of the praseodymium(III) and neodymium(III) borohydride dimethyl sulfide complex, RE(BH₄)₃S(CH₃)₂, RE = Pr, Nd clearly resembled the Bragg reflections of Y(BH₄)₃S(CH₃)₂ and Gd(BH₄)₃S(CH₃)₂. Therefore, the Y(BH₄)₃S(CH₃)₂ structure (monoclinic, P2₁/c) was used as an initial structural model for Rietveld refinements, see Table 2.

In this structure, the rare-earth cation is coordinated to five BH₄⁻ units and a sulfur atom from the dimethyl sulfide molecule, forming a distorted octahedral geometry. Four BH₄⁻ groups bridge between the RE³⁺ centers by edge sharing, bidentate (η²) coordination, and the remaining BH₄⁻ group is a terminal ligand. The S(CH₃)₂ molecule coordinates to RE³⁺ via the S atom of S(CH₃)₂ (Fig. 1). The structure of Pr(¹¹BD₄)₃S(CH₃)₂ was refined using the Rietveld method using both X-ray and neutron powder diffraction data, see Fig. S7 and S8. The refined structural data are provided in Table 2.

**Table 2. Structural data extracted from Rietveld refinements of the XRPD and NPD data for Pr(BH₄)₃S(CH₃)₂, different polymorphs of Pr(BH₄) and Pr⁵⁻Nd(BH₄)₃.**

<table>
<thead>
<tr>
<th>Sample</th>
<th>Pr(¹¹BD₄)₃S(CH₃)₂</th>
<th>α-Pr(BH₄)₃</th>
<th>β⁵⁻-Pr(BH₄)₃</th>
<th>r-Pr(¹¹BD₄)₃</th>
<th>β⁵⁻-Nd(BH₄)₃</th>
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</thead>
<tbody>
<tr>
<td>Crystal system</td>
<td>Monoclinic</td>
<td>Cubic</td>
<td>Cubic</td>
<td>Trigonal</td>
<td>Cubic</td>
</tr>
<tr>
<td>Space group</td>
<td>P2₁/c</td>
<td>Pn₃</td>
<td>Fm₃c</td>
<td>R₃c</td>
<td>Fm₃c</td>
</tr>
<tr>
<td>T(°C)</td>
<td>RT</td>
<td>RT</td>
<td>190</td>
<td>160</td>
<td>260</td>
</tr>
<tr>
<td>ρ(At, H₂) [bar]</td>
<td>1</td>
<td>1</td>
<td>40</td>
<td>1</td>
<td>98</td>
</tr>
<tr>
<td>a (Å)</td>
<td>5.6950(2)</td>
<td>11.2941(5)</td>
<td>11.1438(7)</td>
<td>7.373(6)</td>
<td>11.1386(1)</td>
</tr>
<tr>
<td>b (Å)</td>
<td>22.9167(1)</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>c (Å)</td>
<td>8.2400(4)</td>
<td>—</td>
<td>—</td>
<td>19.89(2)</td>
<td>—</td>
</tr>
<tr>
<td>β(°)</td>
<td>100.68(0)</td>
<td>90</td>
<td>90</td>
<td>120</td>
<td>90</td>
</tr>
<tr>
<td>Space group</td>
<td>4</td>
<td>8</td>
<td>8</td>
<td>6</td>
<td>8</td>
</tr>
<tr>
<td>V/(Å³)</td>
<td>1056.78(60)</td>
<td>1440.64(11)</td>
<td>1383.88(15)</td>
<td>936.38(14)</td>
<td>1381.95(21)</td>
</tr>
<tr>
<td>Z</td>
<td>4</td>
<td>8</td>
<td>8</td>
<td>6</td>
<td>8</td>
</tr>
<tr>
<td>ρ (g cm⁻³)</td>
<td>1.6318</td>
<td>1.7098</td>
<td>1.7801</td>
<td>2.1015</td>
<td>1.8146</td>
</tr>
<tr>
<td>ρ_{m} (H₂) [wt%]</td>
<td>114.0427</td>
<td>111.5413</td>
<td>116.1161</td>
<td>257.1704</td>
<td>116.2783</td>
</tr>
<tr>
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<td>X-ray</td>
<td>X-ray</td>
<td>X-ray</td>
<td>Neutron</td>
<td>X-ray</td>
</tr>
</tbody>
</table>

Pr–B distance for α-Pr(BH₄)₃ is calculated based on the average of two Pr–B distances in the structure.
and are isostructural to RT polymorphs of α-Y(BH₄)₃ and α-Gd(BH₄)₃.²⁰,²⁶,²⁸ The rare-earth atoms coordinate to six BH₄⁻ units with two different RE–B distances, i.e. Pr–B of 2.8535(1) and 2.8679(1) Å, and Nd–B of 2.8306(1) and 2.8449(1) Å, respectively, forming a distorted octahedral geometry. BH₄⁻ complexes bridge the RE³⁺ centers with bidentate (η²) coordination resulting in a 12-fold coordination of RE to hydrogen (Fig. 2, middle).

### 3.2.3 Crystal structure of β-RE(BH₄)₃

Two different space groups of Fm̅3c and Pm̅3m have been reported for β-RE(BH₄)₃ as a high-temperature polymorph. The Fm̅3c structure has a fully ordered array of [BH₄]⁻ groups, with 8 formula units in the unit cell, while the Pm̅3m structure has half the lattice parameter and the unit cell contains one formula unit.²⁰,²⁷,²⁸ The BH₄⁻ complexes randomly take either of the two possible orientations in the Pm̅3m structure. The X-ray diffraction patterns of these two structural models are similar and cannot be distinguished. However, neutron powder diffraction of Y(BH₄)₃ has shown that the Fm̅3c structure is the correct space group.²⁷,²⁸

High temperature polymorphs, β-RE(BH₄)₃, RE = Pr, Nd (Fm̅3c), are formed when α-RE(BH₄)₃ is heated under high hydrogen pressures. The β- polymorphs are isostructural to β-RE(BH₄)₃, RE = Ce, Sm, Ho, Y, Er, Tm, Yb, with straight RE–BH₄ RE coordination (η²), which is isostructural to the ideal cubic ReO₃ structure (Fig. 2, bottom). The RE atoms coordinate to six BH₄⁻ units with the same distance of Pr–B = 2.7859(2) Å (T = 190 °C) and Nd–B = 2.7847(3) Å (T = 269 °C), forming an ideal octahedral geometry in comparison with the distorted octahedron observed for the α-RE(BH₄)₃.²⁷ The β- polymorphs also have a bidentate (η²) coordination of RE to BH₄⁻ forming a 12-fold coordination of RE to hydrogen. This structure contains large unoccupied voids at (1/8, 1/8, 3/8) coordinate with a distance of ~3.35 Å from the center to the nearest hydrogen atom (Fig. 2, bottom).³⁴

### 3.2.4 Crystal structure of r-RE(BH₄)₃

A rhombohedral deformation of cubic β-Pr(BH₄)₃, denoted as r-RE(BH₄)₃, that crystallizes in the trigonal crystal system with space group R̅3c (Z = 6) is also observed, which is isostructural to a high-pressure polymorph of rhenium trioxide, ReO₃.⁵⁴ r-RE(BH₄)₃ has been observed as the room temperature polymorph for RE = La, Ce. However, r-Pr(BH₄)₃ forms at T > 190 °C and the structure has been confirmed by Rietveld refinement of NPD data of r-Pr(BH₄)₃ (Fig. S11†). Six BH₄⁻ complexes are coordinated to each Pr³⁺ ion forming a regular octahedra, Pr–B distance of 2.8395(16) Å, similar to that observed in β-Pr (BH₄)₃.

### 3.2.5 DFT optimization of the Pr(BH₄)₃ polymorphs

Period plane wave DFT calculations of the three polymorphs were attempted in order to shed light on their properties. The structure of α-Pr(BH₄)₃, (the Pa₃ form), has been fully optimized with no constraints (128 atoms; for this large cell, Z = 8, a Monkhorst–Pack grid of 1 × 1 × 1 was selected). The calculations were performed adopting an antiferromagnetic structure, in which half of the Pr atoms have positive magnetization and half negative, with a total magnetization equal to zero. Calculations with different magnetization schemes failed to converge. By this approach, the optimized cell volume is larger than the experimental one by only 1.27%, and final geometry and ion positions in the cell are very similar to the starting one. Similar calculations on the Fm̅3c and the R̅3c structures resulted in a significantly larger unit cell, +12.3% and +20.5%, respectively. The overestimation of the unit cell size is likely due to neglecting phonon interactions, which become significant at higher temperatures required to convert Pr(BH₄)₃ to the Fm̅3c and R̅3c polymorphs. Therefore, a simplified approach was adopted, which allows comparing the energy of the three

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Fig. 2 Crystal structures of (top) r-Pr(BH₄)₃, R̅3c, (middle) α-RE(BH₄)₃, Pa₃, and (bottom) β-RE(BH₄)₃, Fm̅3c. One of the empty voids in the β-Pr (BH₄)₃ structure is shown with a (reddish/violet) sphere.
different structures: only the atom positions within the unit cell were optimized with unit cell parameters fixed to the experimental values. By this method, the $P$α$3$ structure computed by DFT is the most stable, with the $Fm\overline{3}c$ and the $R\overline{3}c$ structure higher in energy by 4.3 and 6.4 kcal mol$^{-1}$, respectively, corresponding to decreasing stability of $\alpha$-Pr(BH$_4$)$_3$ > $\beta$-Pr(BH$_4$)$_3$ > $r$-Pr(BH$_4$)$_3$. Moreover, DFT calculations of the $r$-Pr($^{11}$BD$_4$)$_3$ structure obtained by NPD and with fixed unit cell parameters show 7.4 kcal mol$^{-1}$ higher energy than that of the $P$α$3$ structure. However, this relative energy order could be easily modified at higher temperatures, because the energy values of the three structures are close to each other.

3.3 In situ synchrotron radiation X-ray powder diffraction

3.3.1 Investigation of Pr(BH$_4$)$_3$ polymorphism as a function of $p, T$. The thermal properties of Pr($^{11}$BD$_4$)$_3$S(CH$_3$)$_2$ in $p$(Ar) = 1 bar are investigated by in situ SR-XRPD, see Fig. 3. Only the Bragg reflections of the solvate compound are observed at RT. At $T > 130$ °C, Pr($^{11}$BD$_4$)$_3$S(CH$_3$)$_2$ decomposes with the release of S(CH$_3$)$_2$, and formation of two different polymorphs of praseodymium(III) borohydride, $\alpha$- and $\beta$-Pr($^{11}$BD$_4$)$_3$, 66.8(3) and 33.2(3) wt%, respectively (Fig. S1†). Further heating of the sample leads to a decreasing degree of the long range order of the $\beta$ polymorphs (i.e. decreasing diffracted intensity of $\beta$) and an apparent increasing amount of $\alpha$-Pr($^{11}$BD$_4$)$_3$. This occurs because the sample composition is normalized to 1 (notice that the diffracted intensity of $\alpha$ appears constant in the temperature range of ~150 to ~180 °C). The Bragg reflections of $\beta$-Pr($^{11}$BD$_4$)$_3$ are abruptly shifted to higher 2$\theta$ angles at ~176 °C with a similar abrupt increase of the diffracted intensity, revealing a contraction of the unit cell and the formation of a new polymorph denoted as $\beta'$-Pr(BH$_4$)$_3$ (Fig. S3c) (not shown in Fig. 3b because of the overlap of the Bragg reflections of $\alpha$ and $\beta'$ phases at $20 = 7.2^\circ$ and 10.2$^\circ$). At the same time, the fraction of this polymorph, $\beta'$, increases and reaches a maximum of 33.5(9) wt% at 185 °C (Fig. 3b). At $T = 190$ °C, $\alpha$-Pr($^{11}$BD$_4$)$_3$ disappears, and $r$-Pr($^{11}$BD$_4$)$_3$, with the smallest $V/Z$ forming as the major polymorph, 89(1) wt%. Simultaneously, the unit cell volume of $\beta$-Pr($^{11}$BD$_4$)$_3$ shrinks for the second time upon formation of $\beta''$-Pr($^{11}$BD$_4$)$_3$ (displayed by arrows in Fig. 3a). The structural mechanism for formation of the four polymorphs of praseodymium(III) borohydride is discussed in section 3.5.

Upon cooling, the polymorphs $r$- and $\beta''$-Pr($^{11}$BD$_4$)$_3$ are stable at 119 °C where $\alpha$-Pr($^{11}$BD$_4$)$_3$ starts to form at the expense of $r$-Pr($^{11}$BD$_4$)$_3$. Further cooling of the sample fully transforms the $r$-Pr($^{11}$BD$_4$)$_3$ to $\alpha$-Pr($^{11}$BD$_4$)$_3$, and the sample composition at 59 °C is $\alpha$-Pr($^{11}$BD$_4$)$_3$ 83(1) wt% and $\beta''$-Pr($^{11}$BD$_4$)$_3$, 17.5(7) wt%. The in situ SR-XRPD data of Pr(BH$_4$)$_3$ heated under $p$(Ar) = 1 bar (see Fig. S2†) are similar to those presented in Fig. 3 of Pr($^{11}$BD$_4$)$_3$S(CH$_3$)$_2$.

A second series of in situ XRPD experiments were performed under a hydrogen atmosphere, $p$(H$_2$) = 40 bar, in order to investigate the possible effect of gas and partial pressure on the structural evolution, and the data are presented in Fig. 4a and S3.† In situ SR-XRPD of Pr(BH$_4$)$_3$ at RT reveals diffractions from the polymorphs $\alpha$- (94.7(3) wt%) and $\beta$-Pr(BH$_4$)$_3$ (5.3(2) wt%), see Fig. 4b. Prior to this experiment, Pr(BH$_4$)$_3$ has been annealed at 180 °C and the observed sample composition is similar to the one observed in the previous experiment at 174 °C using Pr($^{11}$BD$_4$)$_3$S(CH$_3$)$_2$ (Fig. 3b). An abrupt displacement of the $\beta$-Pr(BH$_4$)$_3$ Bragg reflections towards lower 2$\theta$ angles is also observed in this case, at 174 °C, and is assigned to the formation of $\beta''$-Pr(BH$_4$)$_3$. Moreover, the unit cell volume of $\beta''$-Pr(BH$_4$)$_3$ shrinks for the second time upon formation of $\beta''$-Pr(BH$_4$)$_3$. However, it is noteworthy that $\beta''$-Pr(BH$_4$)$_3$, 66.6(8) wt%, is now the main component in the sample at 190 °C. The stabilization of the cubic and porous $\beta''$-Pr(BH$_4$)$_3$ in larger
amounts here contrasts the formation of $r$-Pr(BH$_4$)$_3$, 89(1) wt%, at 190 °C, in the previous measurement (Fig. 3b). At $T = 190 \, \text{°C}$, the Bragg reflections of $\alpha$-Pr(BH$_4$)$_3$ disappear and $r$-Pr(BH$_4$)$_3$ forms in smaller amounts. This clearly shows the strong influence of gas and partial pressures on the polymorphism of Pr(BH$_4$)$_3$. Upon cooling, $\beta''$-Pr(BH$_4$)$_3$ remains as the major polymorph at 157 °C where transformation to $\alpha$-Pr(BH$_4$)$_3$ starts.

The extracted unit cell volumes ($V/Z$) as a function of temperature (Fig. 4c) show that the $\beta''$-Pr(BH$_4$)$_3$ structure is $\sim 10 \, \text{Å}^3$ more expanded as compared to that of $r$-Pr(BH$_4$)$_3$ upon cooling in the temperature range of 190 to $\sim 100 \, \text{°C}$, for both experiments (Fig. 3c and 4c).

3.3.2 Investigation of Nd(BH$_4$)$_3$ polymorphism as a function of $p$,$T$. The neodymium borohydride solvate, Nd(BD$_4$)$_3$S(CH$_3$)$_2$ (s4), was also investigated by in situ SR-XRPD under $p$(Ar) = 1 bar, see Fig. S3,† and Bragg reflections of this compound are observed in the temperature range of RT to 165 °C. Nd(BD$_4$)$_3$S(CH$_3$)$_2$ decomposes with release of S(CH$_3$)$_2$ and formation of $\alpha$-Nd(BD$_4$)$_3$, at $T > 165 \, \text{°C}$. However, no polymorphic transitions are observed for Nd(BH$_4$)$_3$ by further heating, and the sample starts to decompose at $T \sim 213 \, \text{°C}$.

In situ SR-XRPD of Nd(BH$_4$)$_3$ under $p$(H$_2$) = 98 bar has also been measured, see Fig. S4.† At RT, the sample contains $\alpha$-Nd(BH$_4$)$_3$, 93(1) wt%, and $\beta$-Nd(BH$_4$)$_3$, 6.8(5) wt%. Similar to Pr(BH$_4$)$_3$, when the sample is heated under hydrogen pressure, a stepwise polymorphic transition is observed $\beta \rightarrow \beta' \rightarrow \beta''$-Nd(BH$_4$)$_3$, i.e. another rare example of negative thermal expansion. This sample fully transforms to $\beta''$-Nd(BH$_4$)$_3$ at 270 °C, which is in contrast to the formation of only 66.6(8) wt% of $\beta''$-Pr(BH$_4$)$_3$ in the previous experiment. These polymorphic transitions are reversible, and Bragg reflections of $\alpha$-Nd(BH$_4$)$_3$ start to appear upon cooling at $T \sim 230 \, \text{°C}$. The observation of the polymorphic transition at higher temperatures for Nd(BH$_4$)$_3$ ($T = 270 \, \text{°C}$) and the suppression of decomposition are clearly due to the elevated hydrogen pressure used in this experiment.

3.4 Atomic distances of praseodymium(m) borohydride as a function of $p$,$T$

The Pr–B distances and weight fractions of praseodymium(m) borohydride polymorphs obtained from Rietveld refinements of in situ SR-XRPD data at $p$(Ar) = 1 bar and selected temperatures are shown in Fig. 5. The $\alpha$-Pr($^{11}$BD$_4$)$_3$ polymorph (67.8(3) wt%) has a Pr–B distance of 2.861(1) Å, at $T = 134 \, \text{°C}$. The major component at 190 °C, $r$-Pr($^{11}$BD$_4$)$_3$ (95(2) wt%) has a longer Pr–B distance of 2.923(4) Å due to transverse rotation of Pr(BD$_4$)$_3$ octahedra. In this process, the B atom in the middle of the Pr–B–Pr linkage is displaced transversely, the Pr–B distance is increased, and the two Pr atoms are pulled together, inducing a contraction in the unit cell volume. Upon cooling to 76 °C, $\alpha$-Pr($^{11}$BD$_4$)$_3$ (73(1) wt%) forms with a shorter Pr–B distance of 2.860(1) Å as compared to the $r$-polymorph.

The Pr–B distances of the different polymorphs of Pr(BH$_4$)$_3$, during heat treatment under $p$(H$_2$) = 40 bar are shown in Fig. 6. At 134 °C, $\alpha$-Pr(BH$_4$)$_3$, 88.7(5) wt%, is the dominant polymorph with Pr–B distances of 2.865(1) Å. Increasing the temperature facilitates the formation of the Pr–B polymorphs with distinct Pr–B distances of 2.872(1), 2.832(2) and 2.786(2) Å for $\beta$, $\beta'$ and $\beta''$ measured at 134, 185 and 190 °C, respectively.

These results reveal that the type of gas (Ar or H$_2$) and partial pressures influence the Pr–B bond lengths. When experiments are performed in $p$(Ar) = 1 bar, complete rotation...
of Pr(BH₄)₆ octahedra occurs and r-Pr(BH₄)₃ with a longer Pr–B distance forms. However, in p(H₂) = 40 bar, the porous β poly morphs stabilize with straight Pr–BH₄–Pr coordination and shorter Pr–B distances.

3.5 Mechanism for transformation of praseodymium(III) borohydride polymorphs

The three polymorphs α-, β- and r-Pr(BH₄)₃, have the rhenium(vi) trioxide, ReO₃, structure types. In an ideal cubic ReO₃ structure, the Re atoms occupy the corners of the cube and coordinate to six oxygen atoms placed on the centers of the edges. This is also the case for the cubic β-Pr(BH₄)₃ with Pr–B–Pr bond angles of 180°. This structure type is closely related to the perovskite structure, i.e. SrTiO₃, but without the Sr atoms located on the center of the cube leaving an empty void in this position (see Fig. 2).⁵⁵

A wide variety of crystal structures can be derived by coupled rotation of octahedra, i.e. [RE(BH₄)₆], without breaking or forming chemical bonds, such as α- and r-Pr(BH₄)₃, which crystallize in distorted ReO₃ structures. At RT, α-Pr(BH₄)₃ is stable, which unfolds to the ideal ReO₃ structure, β-Pr(BH₄)₃, upon heating, but this polymorph is unstable and transforms to r-Pr(BH₄)₃ (Fig. 7). The transformation of the β-Pr(BH₄)₃ → r-Pr(BH₄)₃ involves a coupled rotation of six ReO₆ octahedra, without breaking or forming chemical bonds, as the empty voids in the Fm3c structure are presented by (blue dashed) empty circles.
to r-polymorph is assigned to the large voids in the β-polymorph that allow bending of the Pr–B–Pr linkage, which increases the Pr–B bond distance, decrease the Pr–Pr distance by rotation of Pr(BH₄)₆ octahedra and consequently contract the unit cell volume. The transformation of β- to r-Pr(BH₄)₃ occurs via two other polymorphs, β' and β″-Pr(BH₄)₃ as further discussed below. The polymorphs β- and r-Pr(BH₄)₃ are only stable at elevated temperatures and transform back to α-Pr(BH₄)₃ at RT, possibly due to H–H repulsion.

The relationship between hexagonal (r) and cubic (β) unit cell parameters in an ideal cubic crystal system is

\[ a(r) = \frac{a(β) \sqrt{2}}{2} \]

and

\[ c(r) = a(β) \sqrt{3}. \]

The values determined from the Rietveld refinement of r- and β″-Pr(BH₄)₃ at 190 °C are \( a(β) = 7.4831(12) \) Å, \( c(β) = 19.995(5) \) Å and \( a(β′′) = 11.1438(7) \) Å, respectively. Therefore, the calculation of the non-deformed hexagonal unit cell from the cubic cell is

\[ a(r) = \frac{11.1438 \sqrt{2}}{2} = 7.8798 \text{ Å} \]

and

\[ c(r) = 11.1314 \sqrt{3} = 19.3016 \text{ Å}. \]

This shows that the rhombohedral structure is compressed along \( c(r) \) and expanded along \( a(r) \). The same deformation has been observed for Ce(BH₄)₃ and La[BH₄]₃ polymorphs.³⁴

The cubic to rhombohedral β- to r-Pr(BH₄)₃ transformation deserves more attention. This transformation clearly occurs via two intermediate polymorphs, β' and β″-Pr(BH₄)₃, with distinct unit cell differences as illustrated in Fig. 3c and 4c. The extracted unit cell volumes (V/Z) as a function of temperature reveal a stepwise decrease of ~8–10 Å³ per transformation in the series: \( β \rightarrow β' \rightarrow β'' \rightarrow r-Pr(BH₄)₃ \). The V/Z values suggest that ~1/3 and ~2/3 of the original cubic structure is contracted to the rhombohedral structure type in \( β' \) and \( β''-Pr(BH₄)₃ \) (\( p(H₂) = 40 \) bar). This may suggest static disorder created by rotation of [Pr(BH₄)₆] octahedra, which reduces the Pr–Pr distance and results in a decrease in the unit cell volume. A dynamic effect, such as transverse vibration of BH₄ complexes normal to the Pr–Pr axis or rotation around this axis, is an alternative explanation. However, this is expected to provide a continuous change of unit cell volume as a function of temperature, which contrasts the stepwise changes clearly observed in Fig. 3c and 4c.

A similar behaviour has also been observed for ScF₃ with an identical cubic ReO₃ structure type responsible for negative thermal expansion (NTE) and zero thermal expansion (ZTE) of (Sc₁₋₀.₀₂Gd₀.₀₂Fe₀.₁)F₃.⁵⁶,⁵⁷ However, unlike Pr(BH₄)₃, continuous contraction of the unit cell volume is observed for ScF₃. For these fluoride, the collapse in the unit cell volume upon heating is attributed to the partial formation of a rhombohedral structure in a cubic-like form rather than a rhombohedral unit cell. In this case, the long-range structure remains cubic, but the local structure is distorted in a similar way to that in the rhombohedral polymorph, which makes it very challenging to identify by XRPD. This occurs due to transverse vibration of fluoride anions, i.e. a dynamic effect.⁵⁶

We observe that ~95 wt% of Pr(BH₄)₃ transforms completely to the rhombohedral polymorph upon heating in argon, \( p(\text{Ar}) = 1 \) bar, i.e. the structure becomes rhombohedral both locally and over a long range. However, a minor amount keeps the cubic long range order and shows a stepwise decrease of the unit cell volume, i.e. \( β' \) (20 wt% at 185 °C) and \( β″-Pr(BH₄)₃ \) (5 wt% at 190 °C). Hydrogen gas, \( p(H₂) = 40 \) bar, clearly facilitates the stepwise polymorphic transitions \( β \rightarrow β' \rightarrow β'' \rightarrow α-Pr(BH₄)₃ \) upon heating/cooling of ~65 wt% of the sample. A similar trend is observed for neodymium(iii) borohydride with no observation of β-polymorphs in \( p(\text{Ar}) = 1 \) bar but increasing amount in \( p(H₂) = 40 \) bar.⁶⁰ ⁶ⁱ ⁶² ⁶³ ⁶⁴ ⁶⁵

3.6 Comparison of the crystal structures of the different RE (BH₄)₃ compounds

The series of rare earth borohydrides show significant structural changes as a function of small changes in the cation ionic radii. The largest rare-earth metal borohydrides, La and Ce, crystallize as the polymorph r-RE(BH₄)₃ at RT and have smaller unit cell volumes and higher densities compared to the cubic structure type, α-RE(BH₄)₃. The smaller RE elements crystallize in cubic structures, e.g. α-Pr(BH₄)₃. This work highlights praseodymium as a ‘border line’ RE element, which exists in five different polymorphs α-, β, β′, β″- and r-Pr(BH₄)₃. This trend, presented in Fig. 8, suggests that the crystal structures of the rare-earth metal borohydrides are dependent on the size of the cation.

Previous experiments have shown that β-Ce(BH₄)₃ is only formed when Ce(BH₄)₃(S(CH₃)₂) is heated in a closed capillary, and that annealing under dynamic vacuum forms the r-polymorph. Our experimental work reveals that formation of β-Pr(BH₄)₃ and two intermediate polymorphs, β' and β″-Pr(BH₄)₃, can be facilitated by \( p(H₂) = 40 \) bar and \( T = 175 \) and 190 °C, respectively. Thus, the presence of gases clearly facilitates stabilization of the β-polymorphs.

The radius of the neodymium ion is slightly smaller as compared to that of praseodymium, \( n(\text{Nd}^{3⁺}) = 1.123 \) and \( n(\text{Pr}^{3⁺}) = 1.180 \). [See Fig. 8] Unit cell volumes (V) divided by the number of formula units (Z) of the reported rare-earth borohydrides. β-polymorphs of Pr(BH₄)₃ and Nd(BH₄)₃ polymorphs are recorded at RT, while β″-polymorphs are recorded at 190 and 269 °C, respectively. β' polymorphs of Pr(BH₄)₃ and Nd(BH₄)₃ are not shown in this figure but their structural data are presented in Tables 1–3. The ionic radii are taken from ref. 58. Polymorphs stable at RT are shown with a filled symbol (■). Space groups of Fm₃c and Pm₃m are presented by one symbol (■) because of the similarity of their structures.
The thermal analysis, TGA-DSC-MS, of Pr(BH₄)₃ (s3), Fig. 9, reveals a total release of hydrogen of 4.8 wt%, which is lower than the calculated hydrogen content of 5.97 wt% according to the reaction scheme:

$$2\text{Pr(BH}_4\text{)}_3 \rightarrow \text{PrB}_6 + \text{PrH}_2 + 11\text{H}_2.$$

Two endothermic peaks are observed in the DSC data, one at 204 °C is assigned to the first order polymorphic transition of α-Pr(BH₄)₃ to r-Pr(BH₄)₃ and the second at 247 °C to the decomposition of the sample. In order to investigate the reversibility of the polymorphic transition, a second experiment was conducted by heating the sample to 220 °C and cooling back to RT. Fig. 9b shows that the sample releases ~0.5 wt% of hydrogen during the α to r-polymorphic transition temperature. Upon cooling the sample to RT, an exothermic peak is observed in the DSC curve at 125 °C. This event is attributed to the polymorphic transition of r- to α-Pr(BH₄)₃ and is in good agreement with the reformation of α-Pr(BH₄)₃ at 119 °C observed by the in situ SR-XRPD experiment.

Thermal analysis, TGA-DSC-MS, of Nd(BH₄)₃ is shown in Fig. S14,† and reveals one endothermic peak in the DSC data, at T ~ 250 °C, assigned to decomposition. A release of 5.5 wt% of hydrogen is observed in the temperature range 200 to 400 °C, which is slightly lower than the calculated hydrogen content, 5.87 wt% based on the reaction:

$$2\text{Nd(BH}_4\text{)}_3 \rightarrow \text{NdB}_6 + \text{NdH}_2 + 11\text{H}_2.$$

4. Conclusion

In this study, Pr(11BD₄)₃S(CH₃)₂ and four new polymorphs r- and β-, β'- and β''-Pr(BH₄)₃ are presented, which highlights Pr(BH₄)₃ as a ‘border-line’ rare earth metal borohydride with unique crystal chemistry. At RT, α-Pr(BH₄)₃ (Pa3) is formed, and at T = 190 °C the structure transforms to the more dense polymorph r-Pr(BH₄)₃ (R3c). The cubic (β-) to rhombohedral, r-Pr(BH₄)₃ transformation occurs via two intermediate polymorphs, β'- and β''-Pr(BH₄)₃, with a distinct decrease of unit cell volumes, and is a rare example of stepwise negative thermal expansion. This suggests static disorder created by rotation of [Pr(BH₄)₆] octahedra, which reduces the Pr–Pr distance and results in a decrease in the unit cell volume. A similar, but continuous, change of volume is observed for the isostructural ScB₆ with the same cubic ReO₃ structure type responsible for negative thermal expansion (NTE) due to a dynamic structural effect.

We observe that ~95 wt% of Pr(BH₄)₃ transforms completely to the rhombohedral polymorph upon heating in argon (p(Ar) = 1 bar), i.e. the structure becomes rhombohedral both locally and over a long range. However, a minor amount, ~5 wt%, of Pr(BH₄)₃ keeps the cubic long range order and shows a stepwise decrease of the unit cell volume. Hydrogen gas (p(H₂) = 40 bar) clearly facilitates the stepwise polymorphic transitions α-Pr(BH₄)₃ → β- → β’- → β''- upon heating/cooling of ~65 wt% of the sample. These results reveal that the type of gas (Ar or H₂) and partial pressures have little influence on the characteristic bond lengths observed in the five polymorphs, which contrasts the strongly influence on the stability of the different polymorphs and the sample composition.

Neodymium borohydride, Nd(BH₄)₃, is also synthesized and the polymorphic transitions are compared to Pr(BH₄)₃. The smaller voids in Nd(BH₄)₃ are not large enough to allow the rotation of Nd[BH₄]₆ octahedra and formation of a rhombohedral polymorph. However, the stepwise polymorphic transitions β- → β’- → β'' are also observed for this compound in the presence of hydrogen gas (p(H₂) = 98 bar) and the sample completely transforms to β''-Nd(BH₄)₃ at T = 269 °C.
Here we present two very rare examples of stepwise negative thermal expansion. More detailed structural analysis, e.g. also including spectroscopy and advanced neutron scattering, may provide further insight into both static and dynamic structural effects. The in situ neutron powder diffraction experiment at elevated temperature and pressures of different gasses is time consuming but might be feasible with advanced neutron sources. Further structural knowledge of the origin of stepwise negative thermal expansion may allow rational design and synthesis of novel types of technologically important negative or zero thermal expansion materials.

Conflicts of interest

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