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Study on the thermal conversion of scheelite-type $\text{ABO}_4$ into perovskite-type $\text{AB(O,N)}_3$

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Phase-pure scheelite $\text{AMoO}_4$ and $\text{AWO}_4$ ($\text{A} = \text{Ba, Sr, Ca}$) were thermally treated in ammonia atmosphere at 400 to 900 °C. SrMoO$_4$ and SrWO$_4$ were shown to convert into cubic perovskite SrMoO$_2\text{N}$ and SrWO$_{1.5}\text{N}_{1.5}$, at 700 °C and 900 °C respectively, and to form metastable intermediate phases (scheelite SrMoO$_{4-x}\text{N}_x$ and SrWO$_{4-x}\text{N}_x$), as revealed by X-ray diffraction (XRD), elemental analysis and FTIR spectroscopy. High-temperature oxide melt solution calorimetry reveals that the enthalpy of formation for 10SrM(O,N)$_3$ ($\text{M} = \text{Mo, W}$) perovskites is less negative than that of the corresponding scheelite oxides, though the conversion of the scheelite oxides into perovskite oxynitrides is thermodynamically favorable at moderate temperatures. The reaction of BaMoO$_4$ with ammonia leads to the formation of rhombohedral Ba$_3\text{M}_2\text{(O,N)}_8$ and corresponding binary metal nitrides Mo$_3\text{N}_2$ and W$_{4.6}\text{N}_4$. Similar behavior was observed for CaMO$_4$, which converted upon ammonolysis into individual oxides and nitrides. Thus, BaMoO$_4$ and CaMoO$_4$ were shown to not provide access to perovskite oxynitrides. The influence of the starting scheelite oxide precursor, the structure distortion and the degree of covalency of the B-site-N bond are discussed within the context of the formability of perovskite oxynitrides.

1 INTRODUCTION

Perovskite oxynitrides $\text{AB(O,N)}_3$ are typically synthesized via ammonolysis of oxide precursors; thus they can be formally represented as nitrogen-substituted perovskite-type oxides,$^1, 2$ which exhibit an emerging class of materials suitable for novel applications in the fields of energy conversion, storage, non-toxic pigments, dielectrics, etc.$^3$. Most perovskite-type oxynitrides are synthesized via conversion of scheelite-type $\text{ABO}_4$ and pyrochlore-type $\text{A}_2\text{B}_2\text{O}_7$ upon thermal treatment in ammonia atmosphere. However, not all scheelite- and pyrochlore-type oxides are able to access perovskite oxynitrides. For example, pyrochlore-type La$_2\text{Zr}_2\text{O}_7$, as well as scheelite-type EuMO$_4$ ($\text{M} = \text{Nb and Ta}$) and SrMoO$_4$, provide access to perovskite-type LaZrO$_2\text{N}_4$ as well as EuMO$_2\text{N}_5$ and SrMoO$_2\text{N}_6$, respectively; whereas other precursor oxides such as scheelite-type ATaO$_4$ ($\text{A} = \text{Nd, Sm, Gd, Dy}$) and A$_2\text{W}_2\text{O}_9$ ($\text{A} = \text{Pr, Nd, Sm, Gd, Dy}$) convert upon ammonolysis into pyrochlore-type A$_2\text{Ta}_2\text{O}_6\text{N}_7$ and scheelite-type AWO$_3\text{N}$, respectively.

According to our previous work, only a limited number of perovskite-type oxynitrides are formable.$^9$ For instance, SrMoO$_2\text{N}$, SrWO$_2\text{N}$, CaMoO$_2\text{N}$ and CaWO$_2\text{N}$ appear to be feasible; while, BaMoO$_2\text{N}$ and BaWO$_2\text{N}$ are not stable in the perovskite-type structure. Although perovskite-type SrMo(O,N)$_3$, SrW(O,N)$_3$ and CaMo(O,N)$_3$,$^6, 10-14$ have been reported in the literature (consistent with our prediction based on tolerance and octahedral factors)$^9$, details of the structure evolution of the oxides into perovskite-type oxynitrides are scarce. Furthermore, the existence of perovskite-type BaMo(O,N)$_3$ is questionable,$^{11, 12}$ whereas, perovskite-type BaW(O,N)$_3$ and CaW(O,N)$_3$ have not yet been synthesized.

2 EXPERIMENTAL METHODS

2.1 Synthesis

Scheelite-type oxide precursors (i.e., SrMoO$_4$, SrWO$_4$, BaMoO$_4$, BaWO$_4$, CaMoO$_4$ and CaWO$_4$) were synthesized via solvothermal methods. Thus, Sr(NO$_3$)$_2$ (Sigma-Aldrich, >99.0%), Ba(NO$_3$)$_2$ (Sigma-Aldrich, >99.0%) or Ca(NO$_3$)$_2$·4H$_2$O (Sigma-Aldrich, >99.0%), was mixed in an equimolar ratio with...
Na2MoO4•4H2O (Sigma-Aldrich, >99.5%) or Na2WO4•4H2O (Sigma-Aldrich, >99.5%) in ethylenediamine (FLUKA, > 99.5 %) under vigorous stirring. Subsequently, the reaction mixture was transferred into an autoclave with Teflon lining and heated at 200 °C for 24 h. The resulting mixture was rinsed 5 times with deionized water and ethanol alternately. Centrifugation and drying at 60 °C overnight led to the powdered scheelite-type oxides.

The resulting oxides were ground to fine powders (grain size < 500 nm) and placed in a silica crucible. The thermal treatments were carried out in flowing ammonia at temperatures between 400 and 900 °C for 4 – 24 h. The Schlenk system used for thermal ammonolysis is specifically limited to small batch sizes (about 0.3 – 0.5 g) to maximize exposure to flowing NH3 and thus the product homogeneity.

2.2 Sample characterization

The crystalline phase composition of the as-synthesized samples was analyzed by using powder X-ray diffraction (XRD, STOE STADI P) with Mo Kα radiation (wavelength 0.7093 Å). The oxygen and nitrogen contents of the synthesized samples were determined by hot gas extraction using a LECO TC436 analyzer. Fourier Transform Infrared (FT-IR) spectroscopy was performed on a Varian 670-IR Spectrometer. Thermogravimetric analysis (TGA 92, SETARAM) under ammonia atmosphere was done to obtain the weight change of samples. High resolution transmission electron microscope (HRTEM, JEOL JEM-2100F) was used to assess the morphology and the local crystallinity of the samples.

High temperature oxidative -solution calorimetry was used to determine the enthalpies of formation of the prepared oxynitride samples. This method is well developed and has been applied previously to study nitrides and oxynitrides. Using this technique, ~ 5 mg pellets, made by pressing the powders into a die with a diameter of 1mm, were dropped from room temperature into molten sodium molybdate (3Na2O▪4MoO3) with a diameter of 1mm, were dropped from room temperature into molten sodium molybdate (3Na2O▪4MoO3) and 200 °C for 24 h. The resulting mixture was rinsed 5 times with deionized water and ethanol alternately. Centrifugation and drying at 60 °C overnight led to the powdered scheelite-type oxides.

The resulting oxides were ground to fine powders (grain size < 500 nm) and placed in a silica crucible. The thermal treatments were carried out in flowing ammonia at temperatures between 400 and 900 °C for 4 – 24 h. The Schlenk system used for thermal ammonolysis is specifically limited to small batch sizes (about 0.3 – 0.5 g) to maximize exposure to flowing NH3 and thus the product homogeneity.

The crystallographic data and phase compositions of the as-synthesized samples were constrained to the same value for the anions. All refinements were performed with the Fullprof software.

3 RESULTS AND DISCUSSION

3.1 Ammonolysis of scheelite-type oxides

BaMoO4 and BaWO4. The ammonolysis of the scheelite-type oxides was performed using a neutron wavelength of λ=1.494 Å at SINQ and λ=1.308 Å at HZB. Crystallographic parameters were confirmed by the individual Rietveld refinements of the XRD and ND patterns. The peak shapes were modeled with pseudo-Voigt function for XRD and Thompson-Cox-Hastings pseudo-Voigt function for ND patterns. Isotropic thermal parameters of O/N were constrained to the same value for the anions. All refinements were performed with the Fullprof software.

A noticeable reaction between BaWO4 and NH3 occurs at 700 °C (Figure S3). Compared to BaMoO4, BaWO4 seems to be rather more inert against ammonia, thus more than 50 wt% of BaWO4 still remained after ammonolysis at temperatures up to 850 °C (Table S2). Hence, the ammonolysis of BaWO4 at 700 and 850 °C leads to a mixture consisting of BaWO4, Ba3W2O6N2 and Ba3W2N2. The lattice parameters of Ba3W2O6N2 and Ba3W2N2 were determined as 5.9706 (5) and 21.5020 (6) Å respectively. The ammonolysis of the scheelite-type oxides was performed at 600, 700 and 900 °C for 6 h. The sample treated at 600 °C already formed small amounts of Ba3Mo2O6N2 oxynitride phase (structure identical to Ba3Mo2O6N2). The absence of the perovskite-type BaMoO4 is consistent with the experimental work of Liu et al. and our previously reported values. No cubic perovskite BaMoO3 was observed. Up to 900 °C, only small amounts of BaMoO4 and BaWO4 were detected besides the main phase Ba3Mo2O6(N2).
The samples obtained upon ammonolysis of BaMoO$_4$ and BaWO$_4$ were also investigated by FTIR spectroscopy; both show an absorption band around 975 cm$^{-1}$ for oxy nitride (Figure S5), which was assigned to a stretching mode (ν(M-N)) in (WO$_3$N)$_3^-$ / (MoO$_3$N)$_3^-$, having W$^{6+}$ / Mo$^{6+}$ in tetrahedral coordination, as reported by Herle et al.$^{30}$

Thus ammonolysis of the scheelite-type oxides BaWO$_4$ and BaMoO$_4$ leads to non-perovskite oxynitrides products following the paths proposed in the equations (1)-(3):

10$\text{BaMoO}_4 + \frac{38}{3} \text{NH}_3 \xrightarrow{700 \degree C} 3\text{Ba}_3\text{Mo}_2(\text{O,N})_8 + \text{BaMoO}_3 + \text{Mo}_2\text{N}_2 + 19\text{H}_2\text{O} + \frac{14}{3} \text{N}_2$

$9\text{BaMoO}_4 + 12\text{NH}_3 \xrightarrow{900 \degree C} 3\text{Ba}_3\text{Mo}_2(\text{O,N})_8 + \text{Mo}_2\text{N}_2 + 18\text{H}_2\text{O} + 2\text{N}_2$

$69\text{BaWO}_4 + 92\text{NH}_3 \xrightarrow{850 \degree C} 23\text{Ba}_3\text{W}_2(\text{O,N})_8 + 5\text{W}_{1.6}\text{N}_4 + 138\text{H}_2\text{O} + 13\text{N}_2$

Thus ammonolysis of scheelite-type SrMoO$_4$ was found to proceed in a different way, leading to the formation of CaO and various molybdenum nitrides (including Mo$_2$N, Mo$_3$N$_2$ and MoN, depending on the temperature, time and ammonia flow) (Figure S6) and consequently CaMoO$_4$ was not considered further as a precursor for the corresponding perovskite-type oxynitrides.

Ammonolysis of CaWO$_4$ at 900 $\degree$C for 6 h leads to complete decomposition into Ca$_3$WO$_6$ and W$_4$N$_4$ and no oxynitride phase was observed (Figure S6). For both CaMoO$_4$ and CaWO$_4$, the corresponding perovskite oxynitrides did not form and thus their conversion into oxide/nitride mixture is assumed to occur as follows:

$6\text{CaMoO}_4 + 12\text{NH}_3 \xrightarrow{700 \degree C} 6\text{CaO} + \text{Mo}_2\text{N} + \text{Mo}_3\text{N}_2 + \text{MoN} + 18\text{H}_2\text{O} + 4\text{N}_2$

$6\text{CaWO}_4 + 9.2\text{NH}_3 \xrightarrow{900 \degree C} 11.5\text{Ca}_3\text{WO}_6 + \text{W}_{4.6}\text{N}_4 + 13.8\text{H}_2\text{O} + 2.6\text{N}_2$

**3.2 Intermediate oxynitride phase during the conversion of SrMoO$_4$ into perovskite-type SrMoO$_2$N**

An interesting phenomenon during the ammonolysis of SrMoO$_4$ at 600 $\degree$C relates to the incorporation of 2.23 wt % nitrogen without the formation of any new crystalline phase; thus, the color of the sample changed from white to light-grayish and the FTIR spectrum showed a new absorption band at 978 cm$^{-1}$ related to (MoO$_3$N)$_3^-$ units in tetrahedral coordination (Figure 3a),$^{30}$ as observed also in Ba$_3$Mo$_3$(O,N)$_8$. Tetra-coordinated Mo$^{6+}$ in scheelite-type SrMoO$_4$ can be identified by FTIR spectroscopy via very broad band around 822 cm$^{-1}$ representing the
antisymmetric stretching vibrations of Mo-O in (MoO\textsubscript{4})\textsuperscript{2-} tetrahedral units.\textsuperscript{35} Thus, the formation of (MoO\textsubscript{3}N)\textsuperscript{3-} is considered to be a result of the substitution of one oxygen with nitrogen in (MoO\textsubscript{4})\textsuperscript{2-} tetrahedra. Therefore, we assume that an intermediate scheelite-type oxynitride phase SrMoO\textsubscript{4}\textsubscript{x}N\textsubscript{y} (x = 0.39 in our experiment, as obtained from elemental analysis and Rietveld refinement, Figure 4) forms at 600 °C, which subsequently rearranges into the perovskite structure while taking up more nitrogen. The absence of the absorption band of (MoO\textsubscript{3}N)\textsuperscript{3-} in the samples obtained upon ammonolysis at temperatures above 700 °C might be related to the strong absorption of the black sample.

Figure 3 FTIR spectrum of the as-synthesized scheelite oxide (a) SrMoO\textsubscript{4}, (b) SrWO\textsubscript{4} and the resulting oxynitrides from ammonolysis at different temperatures (400, 600, 700 and 900 °C) for 6h..

Figure 4 Rietveld patterns of the X-ray powder diffraction data of the sample obtained upon ammonolysis of the SrMoO\textsubscript{4} at 600 °C for 4h. Blue tick marks are Bragg peak positions of related phase as SrMoO\textsubscript{3.61(3)}N\textsubscript{0.39(3)} (the ratio of O/N was fixed based on the results of the elemental analysis). The green line at the bottom denotes the difference intensities between the observed and calculated profiles. Table S3 summarizes the results of the structure refinement.

Yang et al.\textsuperscript{35} reported the formation of analogous scheelite-type EuWO\textsubscript{6-x}N\textsubscript{y} oxynitride as intermediate phase during the nitridation from Eu\textsubscript{2}W\textsubscript{2}O\textsubscript{7} to EuWO\textsubscript{14}N\textsubscript{2-x}. However, in their case, the nitrogen substitution is compensated by the partial oxidation of Eu\textsuperscript{2+} to Eu\textsuperscript{3+} (y=0.04 in EuWO\textsubscript{6-x}N\textsubscript{y}, i.e., Eu\textsuperscript{2+1-x}\textsubscript{y}WO\textsubscript{6-x}N\textsubscript{y}.\textsuperscript{35,37}) In our system, Sr\textsuperscript{2+} is not able to be oxidized to Sr\textsuperscript{3+}, so a different mechanism must be responsible for the formation of the nitrogen-containing scheelite-based phase. A likely explanation is that the generation of oxygen vacancies compensates the extra negative charge due to the replacement of oxygen by nitrogen within the pre-formed crystallites, as usually occurs for nitrogen-doped TiO\textsubscript{2} \textsuperscript{38,39} or HfO\textsubscript{2} \textsuperscript{40}. As shown in HRTEM within FFT pattern (Figure 5), the crystalline phase in the sample obtained after ammonolysis of SrMoO\textsubscript{4} at 600 °C for 4h was indexed as tetragonal (I 41/a, i.e., same as scheelite-type SrMoO\textsubscript{4}) and exhibited the presence of pores. Some defect regions with different fringe distance were observed as well probably due to the distortion of lattice. Interestingly, thermogravimetric analysis of the SrMoO\textsubscript{4} in ammonia revealed a slight mass increase of the sample at temperatures up to 600 °C (Figure 6), indicating that the oxygen, which is expected to be released from SrMoO\textsubscript{4} upon ammonolysis, might be stored at intermediate temperature in the pores or interstitially in the structure as molecular oxygen \textsuperscript{35} before being released (as shown by mass loss of SrMoO\textsubscript{4} at temperatures beyond 700 °C, see Figure 6). This was shown to be the reason for anomalous magnetic behavior at T= -219 °C (54 K) as reported by Longvinovich et al.\textsuperscript{35} The sharp weight loss above 650 °C is attributed to the complete conversion from scheelite to perovskite resulting in 1 mol oxygen released. Elemental analyses confirm the expected oxygen loss for samples heated in NH\textsubscript{3} between 600 and 700 °C and are in agreement with the measured mass loss, indicating that nitrogen is already incorporated in the sample at 600 °C. (Table 1 and Table S6)

Based on all these observations, we conclude that the nitridation of SrMoO\textsubscript{4} occurs prior to the reduction of W\textsuperscript{6+} during ammonolysis, thus scheelite-type SrMoO\textsubscript{4-x}N\textsubscript{x} forms as an intermediate phase and decomposes fast according to the following paths (\( \sigma \) stands for amount of oxygen vacancies):

\[
\begin{align*}
\text{SrMoO}_4 + \sigma \text{NH}_3 & \rightarrow \text{SrMoO}^{\sigma \text{V}}\text{O}^{\sigma \text{V}}\text{O}_\sigma \text{N}_\sigma + \text{O}_2 + \sigma \text{H}_2 \text{O} + \frac{\sigma}{2} \text{H}_2 \\
\text{SrMoO}_4^{\sigma \text{V}}\text{O}^{\sigma \text{V}}\text{O}_\sigma \text{N}_\sigma + \text{NH}_3 & \rightarrow \text{SrMoO}_3^{\sigma \text{V}}\text{O}^{\sigma \text{V}}\text{O}_\sigma \text{N}_{\sigma \text{V}} + \text{H}_2 \text{O} + \text{H}_2
\end{align*}
\]

Figure 5 HRTEM micrographs of SrMoO\textsubscript{4} after heating at 600 °C for 4h.

Table 1 Experimental and calculated mass loss of SrMoO\textsubscript{4} upon ammonolysis at 600 and 700 °C. The calculated mass loss relies on the evolution of the chemical composition of the sample upon ammonolysis.

<table>
<thead>
<tr>
<th>Specimens</th>
<th>Experiment</th>
<th>Empirical Formula</th>
<th>Calculated</th>
</tr>
</thead>
<tbody>
<tr>
<td>SrMoO\textsubscript{4} NH\textsubscript{600} 4H</td>
<td>0.4218 g</td>
<td>SrMo\textsubscript{3}O\textsubscript{2}N\textsubscript{2.35}</td>
<td></td>
</tr>
<tr>
<td>SrMoO\textsubscript{4} NH\textsubscript{700} 12H</td>
<td>0.3925 g</td>
<td>SrMo\textsubscript{3}O\textsubscript{3}N\textsubscript{1.04}</td>
<td></td>
</tr>
<tr>
<td>Mass loss (wt %)</td>
<td>6.95</td>
<td>6.97</td>
<td></td>
</tr>
</tbody>
</table>

Figure 6 TG curve of SrMoO\textsubscript{4} under ammonia atmosphere from room temperature to 800 °C.
3.3 Ammonolysis of SrMoO₄ vs. SrMoO₃

In order to investigate the influence of the oxide precursor on the final oxynitride, we converted the scheelite-type SrMoO₄ into SrMoO₂.77(3)N₀.23(3) via a two-step process as well. In a first step, the scheelite-type oxide SrMoO₄ was easily reduced to the 5 perovskite-type SrMoO₃ (Figure S7) upon thermal annealing at 900 °C for 6 h under forming gas (mixture of 5 vol% H₂ and 95 vol% N₂). In a subsequent step, SrMoO₃ underwent ammonolysis at 700 °C for 4 h (same conditions as for SrMoO₄) to achieve SrMo(O,N)₃. Interestingly, the nitrogen content of the phase-pure 10 perovskite-type oxynitride (empirical formula SrMoO₂.77(3)N₀.23(3), see the Rietveld refinement data of the neutron diffraction pattern shown in Figure 7a) obtained from perovskite-type SrMoO₃ was significantly lower than that of the oxynitride obtained under the same conditions from SrMoO₄ (SrMoO₂.19(2)O₀.81(2)). This obviously relates to the oxidation state of Mo in SrMoO₃ and SrMoO₄ and its evolution in ammonia atmosphere which will be discussed later.

Moreover, the attempt to prepare perovskite-type SrWO₃ upon reducing SrWO₄ at high temperatures was unsuccessful.

Figure 7 Rietveld patterns of the neutron powder diffraction data of the sample obtained upon ammonolysis of the (a) SrMoO₃ at 700 °C for 4 h (FIREPOD, E9); (b) SrMoO₄ at 700 °C for 4 h (HRPT, SINQ) and (c) SrWO₄ at 900 °C for 4 h (FIREPOD, E9). Blue tick marks are Bragg peak positions of related phase as (a) SrMoO₂.77(3)N₀.23(3); (b) SrMoO₂.19(2)O₀.81(2) and (c) SrWO₁.50(6)N₁.50(6). Green line at the bottom denotes the difference intensities between the observed and calculated profiles.

3.4 Structure verification of perovskite oxynitrides

The neutron powder diffraction data measured at room temperature for SrMo(O,N)₃ and SrW(O,N)₃ were refined by the Rietveld method on the basis of the cubic Pm-3m perovskite-type structure (Figure 7, Table 2). The refined O/N content of SrMoO₂.19(2)N₀.81(2) (700 °C for 4 h) and SrWO₁.50(6)N₁.50(6) (900 °C for 4 h) are consistent with the results of elemental analysis (Supporting Information, see Table S6).

<table>
<thead>
<tr>
<th>Specimens and parameters</th>
<th>SrMoO₂₋₇₋₃₋₁₋₂₋₃</th>
<th>SrMoO₂₋₈₋₂₋₂₋₁₋₂</th>
<th>SrWO₁₋₅₋₀₋₁₋₆</th>
</tr>
</thead>
<tbody>
<tr>
<td>S.G.</td>
<td>Pm-3m, Nr. 221</td>
<td>Pm-3m, Nr. 221</td>
<td>Pm-3m, Nr. 221</td>
</tr>
<tr>
<td>Z</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>a,b,c, Å</td>
<td>3.9744(3)</td>
<td>3.9756(1)</td>
<td>3.9856(2)</td>
</tr>
<tr>
<td>Sr x,y,z, Å²</td>
<td>0.5, 0.5, 0.5</td>
<td>0.5, 0.5, 0.5</td>
<td>0.5, 0.5, 0.5</td>
</tr>
<tr>
<td>Mo/W x,y,z, Å²</td>
<td>0.298(23)</td>
<td>0.693(18)</td>
<td>0.880(57)</td>
</tr>
<tr>
<td>O/N x,y,z, Å²</td>
<td>0.5, 0.0, 0.0</td>
<td>0.0, 0.0, 0.0</td>
<td>0.5, 0.0, 0.0</td>
</tr>
<tr>
<td>Biso, Å²</td>
<td>0.748(18)</td>
<td>0.799(12)</td>
<td>0.798(32)</td>
</tr>
<tr>
<td>Occ.</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>O/cubic</td>
<td>1.50(6)</td>
<td>1.50(6)</td>
<td>1.50(6)</td>
</tr>
</tbody>
</table>

The enthalpies of dissolution (ΔHds) and formation (ΔHf) of scheelite-type SrMoO₄ and SrWO₄ and the corresponding perovskite-type oxynitride samples measured by high temperature oxide melt solution calorimetry are listed in Table 3.

<table>
<thead>
<tr>
<th>Composition</th>
<th>Crystal structure</th>
<th>ΔHds (kJ/mol)</th>
<th>ΔHf (kJ/mol)</th>
<th>ΔHf (kJ/g-atom)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SrMoO₄</td>
<td>Tetragonal/Scheelite</td>
<td>161.8 ± 1.5</td>
<td>-1561.3 ± 3.1</td>
<td>-260.2 ± 0.5</td>
</tr>
<tr>
<td>SrMoO₂₋₉₋₂₋₁₋₂</td>
<td>Cubic/Perovskite</td>
<td>-291.9 ± 2.3</td>
<td>-1119.1 ± 3.6</td>
<td>-223.8 ± 0.7</td>
</tr>
<tr>
<td>SrWO₄</td>
<td>Tetragonal/Scheelite</td>
<td>-162.8 ± 1.5</td>
<td>-1641.2 ± 3.1</td>
<td>-273.4 ± 0.5</td>
</tr>
<tr>
<td>SrWO₂₋₅₋₀₋₁</td>
<td>Cubic/Perovskite</td>
<td>-537.2 ± 1.9</td>
<td>-952.9 ± 3.6</td>
<td>-190.4 ± 0.7</td>
</tr>
</tbody>
</table>

The enthalpies of formation of the oxides and oxynitrides from the elements were calculated through the thermodynamic cycles shown in Tables S 7 and S 8 and are given in Table 3. The enthalpy of formation of SrMoO₄ (-260.2 ± 0.5 kJ/g-atom) is ~ 36 kJ/g-atom more exothermic than that of SrMoO₂₋₉₋₂₋₁₋₂ (-223.8 ± 0.7 kJ/g-atom). Likewise, the enthalpy of formation of SrWO₄ (-273.4 ± 0.5 kJ/g-atom) is ~ 83 kJ/g-atom more exothermic than that of SrWO₂₋₅₋₀₋₁ (-190.4 ± 0.7 kJ/g-atom). Thus, perovskite-type oxynitrides show less favorable enthalpies of formation than their corresponding scheelite-type oxides. Furthermore, the difference of the enthalpy of formation for Sr-W is larger than that of Sr-Mo. This suggests that forming the Sr-W oxynitride is
less favorable and requires higher temperature (as observed), probably for both thermodynamic and kinetic reasons.

In order to attain further insights related to the energetics of the conversion of SrMoO$_4$ into SrM(O,N)$_3$ (M=Mo, W) in ammonia atmosphere, the Gibbs free energy ($\Delta G$) of the reaction (8) and (9) was calculated (Table S9 and S10). Since the entropies of SrMoO$_2$N and SrWO$_1.5$N$_{1.5}$ are not available, we estimated them as 5/6 of the entropy of corresponding scheelite-type oxide. Equation (9) and (11) describe the temperature evolution of the Gibbs free energy of the reaction of SrMO$_4$ with 10NH$_3$ to deliver SrM(O,N)$_3$, indicating that the reaction is spontaneous at temperatures exceeding 992 K (i.e., 719.⁰C) for SrWO$_4$; whereas for SrMoO$_4$ the reaction seems to be thermodynamically favorable at any of the temperatures used for its ammonolysis (Figure 8). It is worth to point out that only a thermodynamic consideration might not be enough to describe the ammonolysis processes of the scheelite oxides. The kinetics (e.g., activation energy) of the ammonolysis probably play also an important role and thus might explain why the conversion of SrMoO$_4$ to the perovskite oxynitride needs temperatures exceeding 600 ⁰C and proceeds through an intermediate phase.

The negative temperature dependence of the free energy reflects a positive entropy of reaction because 1.5 moles of gas are produced.

\[
\begin{align*}
\text{SrMoO}_4 + 2\text{NH}_3 & = \text{SrMoO}_2\text{N} + 2\text{H}_2\text{O} + \frac{1}{2}\text{N}_2 + \text{H}_2 \quad (8) \\
\Delta G_{\text{Sr-Mo}}(kJ/mol) & = 50.404 - 0.197T \\
\text{SrWO}_4 + 2\text{NH}_3 & = \text{SrWO}_1.5\text{N}_{1.5} + \frac{5}{2}\text{H}_2\text{O} + \frac{1}{2}\text{H}_2 + \frac{1}{4}\text{N}_2 \quad (10) \\
\Delta G_{\text{Sr-W}}(kJ/mol) & = 175.615 - 0.177T
\end{align*}
\]

The negative temperature dependence of the free energy reflects a positive entropy of reaction because 1.5 moles of gas are produced.

**Figure 8** Gibbs free energy ($\Delta G$) for the ammonolysis of SrMoO$_4$ and SrWO$_4$ (Eqs. (8) and (10), respectively) as function of the temperature.

### 3.5 Factors affecting the formation of perovskite-type oxynitrides

As addressed above, the experimental results related to the conversion of BaMoO$_4$, BaWO$_4$, SrMoO$_4$ and SrWO$_4$ into perovskite-type oxynitrides are consistent with our prediction. However, CaMoO$_4$ and CaWO$_4$ appear to not be converted to oxynitrides.

Scheelite-type ABO$_4$ oxides are rather common precursors for the synthesis of perovskite oxynitrides, e.g. Nd$^{3+}$Y$^4+$O$_2$N$^{41}$, Eu$^{3+}$Nb$^5+$O$_2$N$^{31}$, La$^{3+}$Nb$^5+$O$_2$N$^{32}$, Ca$^{2+}$Sr$^{2+}$W$_{5^+}$O$_2$N$^{43}$ and so on. The formation of hydrogen due to the dissociation of ammonia under high temperature is beneficial for the reduction of the B-site cation in scheelite-type oxide (e.g. from A$^{2+}$B$^{6+}$O$_4$ to A$^{2+}$B$^{5+}$O$_2$N or from A$^{3+}$B$^{5+}$O$_4$ to A$^{3+}$B$^{4+}$O$_2$N). In the case of using perovskite oxides as precursors for perovskite-type oxynitrides, the B-site cation has to be oxidized in order to compensate the increasing of the negative charge resulted from nitrogen incorporation (e.g. from Sr$^{2+}$Mo$^{4+}$O$_3$ to Sr$^{2+}$Mo$^{5+}$O$_2$N). Thus, it seems that scheelite-type oxide precursors are more favorable for the synthesis of perovskite-type oxynitrides.

Moreover, parameters such as the tolerance factor (describing the distortion of the cubic perovskite structure) were shown to be crucial for the formability of perovskite-type oxynitrides. As defined by Goldschmidt, the tolerance factor ($t$) in ABX$_3$ is expressed as:

\[
t = \frac{(r_A - r_B)}{\sqrt{r_A - r_X}} \quad [10]
\]

In our previous work, the formability of perovskite-type oxynitrides was also rationalized upon assessing the values of the tolerance factor, defined as in Eq. [11] (see Table 4, for O/N ratio 2; i.e., ABO$_2$N):

\[
t_{\text{ox}} = \frac{[(r_A + r_B)^{12} + 2(r_A + r_B)^{11} + (r_A + r_B)^{10} - (r_A + r_B)^9 - r_A^9 - r_B^9]^{1/12}}{2[(r_A + r_B)^{12} + 2(r_A + r_B)^{11} + (r_A + r_B)^{10} - (r_A + r_B)^9 - r_A^9 - r_B^9]^{1/12}} \quad [11]
\]

**Table 4** The tolerance factors for ABO$_3$ and ABO$_2$N calculated with the Equations [11] and [12], respectively.

<table>
<thead>
<tr>
<th>Oxide</th>
<th>BaMoO$_4$</th>
<th>BaWO$_4$</th>
<th>SrMoO$_4$</th>
<th>SrWO$_4$</th>
<th>CaMoO$_4$</th>
<th>CaWO$_4$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$t_o$</td>
<td>1.035</td>
<td>1.027</td>
<td>0.98</td>
<td>0.975</td>
<td>0.954</td>
<td>0.941</td>
</tr>
<tr>
<td>$t_{\text{ox}}$</td>
<td>1.053</td>
<td>1.048</td>
<td>0.995</td>
<td>0.989</td>
<td>0.959</td>
<td>0.955</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Oxynitride</th>
<th>BaMoO$_2$N</th>
<th>BaWO$_2$N</th>
<th>SrMoO$_2$N</th>
<th>SrWO$_2$N</th>
<th>CaMoO$_2$N</th>
<th>CaWO$_2$N</th>
</tr>
</thead>
<tbody>
<tr>
<td>Predicted</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
</tr>
<tr>
<td>Experiment</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
</tr>
</tbody>
</table>

$P$: perovskite; $N$: non-perovskite

: predicted formability of oxynitrides

: formability of oxynitrides for this work

For a general consideration of the formability of perovskite-type oxynitrides, we compare their tolerance factors with those of the
corresponding perovskite-type oxides. The values of the tolerance factors \( t_c \) and \( t_{oct} \) calculated from the ionic radii are shown in Table 4 and indicate that the formal substitution of \( \text{O}^2- \) with \( \text{N}^3- \) in \( \text{SrMoO}_3, \text{SrWO}_3, \text{CaMoO}_3 \) and \( \text{CaWO}_3 \) reduces the structural distortion (i.e., the tolerance factor become closer to unity), which suggests that the formation of the corresponding perovskite-type oxynitrides is unfavorable. This is in agreement with the experiment for \( \text{Sr-Mo/Sr-W compounds} \) and does not fit the experimental observations for \( \text{Ca-Mo/Ca-W compositions} \).

Large basic cations like \( \text{Ca} \) typically stabilize higher oxidation states of the transition metals (\( \text{Mo, W} \) as in our case), \(^{46, 47}\) thus this may explain why the \( \text{Ca scheelite-type oxides} \) cannot be converted into oxynitrates.

On the contrary, incorporation of nitrogen within \( \text{BaMoO}_3 \) and \( \text{BaWO}_3 \) increases the structural distortion; thus, the formation of \( \text{BaMoO}_3\text{N}_x \) and \( \text{BaWO}_3\text{N}_x \) would be less favorable. This is in agreement with our synthetic observation.

Moreover, the higher covalent character of the \( \text{B-site-N} \) bond than that of the \( \text{B-site-O} \) bond might also induce structural distortion into the perovskite structure of oxynitrides as compared to their analogous perovskite oxides, i.e. \( \text{B(O,N)}_6 \) octahedra are expected to be more distorted than their analogous \( \text{BO}_6 \) octahedra. This structural distortion might be quite pronounced, as for compounds which exhibit so-called second-order Jahn-Teller distortion \(^{48}\) (i.e. \( \text{d}^0 \) B-site octahedra such as in \( \text{LaZrO}_2 \), \( \text{NdTiO}_2 \) or \( \text{LaTiO}_2 \)). \(^{4} \) However, we consider in our compounds \( \text{SrM(O,N)}_3 \), the first-order Jahn-Teller effect being relevant and thus the contribution of the \( \text{B-site-N} \) covalency on the distortion might not be significant.

4 CONCLUSIONS

In the present study, preparative possibilities to access perovskite-type oxynitrides \( \text{AM(O,N)}_3 \) (\( \text{A}=\text{Ba, Sr, Ca; B}=\text{Mo, W} \)) phases upon thermal ammonolysis of scheelite-type \( \text{AMO}_4 \) oxide precursors were investigated. The as-synthesized results of perovskite-oxynitrides are consistent with our previous prediction in general.

The experimental data reveal that both scheelite-type \( \text{SrMoO}_4 \) and \( \text{SrWO}_4 \) transform into a scheelite-type oxynitride intermediate phase \( \text{SrMO}_3\text{N}_x (M=\text{Mo, W}) \), which subsequently converts fast into perovskite-type \( \text{SrM(O,N)}_3 \) at temperatures above \( 600 \) °C and are in agreement with the high temperature oxide melt solution calorimetry experiments which indicate that the conversion of scheelite \( \text{SrMoO}_4 \) into perovskite \( \text{SrM(O,N)}_3 \) is thermodynamically favorable at the used ammonolysis temperatures.

Furthermore, the formability of the perovskite-type oxynitrides depends on the structure of the used oxide precursor (scheelite seems to be favorable, except for large basic \( \text{A} \) cations) and on the structural distortion described by the tolerance factor.

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The authors acknowledge Dr. Samuel Bernard (IEM, University Montpellier 2) for the TGA of \( \text{SrMoO}_4 \) in ammonia atmosphere.

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Notes and references

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\(^{f}\) Electronic Supplementary Information (ESI) available: XRD, FTIR, lattice parameters and phases composition obtained by Rietveld refinement, elemental analysis and the enthalpies of formation results as well as Gibbs free energy calculation. See DOI: 10.1039/b000000x/

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Figure 1 XRD patterns of SrMoO4 after heating at 400, 600 and 700 °C for different times under an ammonia flow in forming gas (mixture of 5 vol% H2 and 95 vol% N2). Arrow indicates the diffraction pattern of the oxynitride obtained upon ammonolysis of SrMoO3 which was synthesized by reducing SrMoO4 under ammonia flow at 700 °C.
Figure 2 XRD patterns of SrWO4 after heating at 400, 600, 700 and 900 °C for different time under an ammonia flow.
Figure 3 FTIR spectrum of the as-synthesized scheelite oxide (a) SrMoO$_4$, (b) SrWO$_4$ and the resulting oxynitrides from ammonolysis at different temperatures (400, 600, 700 and 900 °C) for 6h.
Figure 4 Rietveld patterns of the X-ray powder diffraction data of the sample obtained upon ammonolysis of the SrMoO4 at 600 °C for 4 h. Blue tick marks are Bragg peak positions of related phase as SrMoO3.61(3)N0.39(3) (the ratio of O/N was fixed based on the results of the elemental analysis). The green line at the bottom denotes the difference intensities between the observed and calculated profiles. Table S3 summarizes the results of the structure refinement.
Figure 5 HRTEM micrographs of SrMoO$_4$ after heating at 600 °C for 4 h. 90x180mm (150 x 150 DPI)
Figure 6 TG curve of SrMoO4 under ammonia atmosphere from room temperature to 800 °C.

184x138mm (150 x 150 DPI)
Figure 7 Rietveld patterns of the neutron powder diffraction data of the sample obtained upon ammonolysis of the (a) SrMoO$_3$ at 700 °C for 4 h (FIREPOD, E9); (b) SrMoO$_4$ at 700 °C for 4 h (HRPT, SINQ) and (c) SrWO$_4$ at 900 °C for 4 h (FIREPOD, E9). Blue tick marks are Bragg peak positions of related phase as (a) SrMoO$_{2.77(3)}$N$_{0.23(3)}$; (b) SrMoO$_{2.19(2)}$N$_{0.81(2)}$ and (c) SrWO$_{1.50(6)}$N$_{1.50(6)}$. Green line at the bottom denotes the difference intensities between the observed and calculated profiles.
Figure 8 Gibbs free energy (ΔG) for the ammonolysis of SrMoO4 and SrWO4 (Eqs. (8) and (10), respectively) as function of the temperature.

190x138mm (150 x 150 DPI)