Relationships between elastic anisotropy and thermal expansion in A₂M₀₃O₁₂ materials†

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We report calculated elastic tensors, axial Grüneisen parameters, and thermal stress distributions in Al₂Mo₃O₁₂, ZrMgMo₃O₁₂, Sc₂Mo₃O₁₂, and Y₂Mo₃O₁₂, a series of isomorphic materials for which the coefficients of thermal expansion range from low-positive to negative. Thermal stress in polycrystalline materials arises from interactions between thermal expansion and mechanical properties, and both can be highly anisotropic. Thermal expansion anisotropy was found to be correlated with elastic anisotropy: axes with negative thermal expansion were less compliant. Calculations of axial Grüneisen parameters revealed that the thermal expansion anisotropy in these materials is in part due to the Poisson effect. Models of thermal stress due to thermal expansion anisotropy in polycrystals following cooling showed thermal stresses of sufficient magnitude to cause microcracking in all cases. The thermal expansion anisotropy was found to couple to elastic anisotropy, decreasing the bulk coefficient of thermal expansion and leading to lognormal extremes of the thermal stress distributions.

Introduction

Materials with the general formula A₂M₀₃O₁₂, where A is a trivalent cation and M is W or Mo, have the unusual property of linear coefficients of thermal expansion spanning the range from low positive (2.4 × 10⁻⁶ K⁻¹) to negative (−9 × 10⁻⁶ K⁻¹), including zero. Their structures have considerable chemical flexibility, as solid solutions can be made incorporating cations with large differences in size (e.g., Al³⁺ and Y³⁺). Aliovalent substitutions of Zr⁴⁺ or Hf⁴⁺ with Mg²⁺ for A³⁺ can be performed, offering additional possibilities for the tailoring of physical properties within this family. The structures of these materials are composed of vertex-linked networks of AO₆ octahedra and MO₄ tetrahedra; negative contributions to thermal expansion arise from transverse motions of bridging oxygen atoms accompanied by distortion of the AO₆ octahedra. Negative thermal expansion (NTE) has been correlated with the compliance of the A-O ionic bond, and the subsequent distortability of the AO₆ octahedra.

The open-framework structure allows for unusual thermal expansion and chemical flexibility, and it typically leads to low stiffness in comparison to other oxide ceramics. The thermal expansion of materials from the A₂M₀₃O₁₂ family is known to be highly anisotropic. Such anisotropy can cause large thermal stresses and subsequent microcracks in polycrystalline materials with randomly oriented grains. However, the elastic anisotropy of these materials is generally unknown; furthermore, it is more difficult to determine experimentally than the anisotropy of the coefficient of thermal expansion (CTE). Critically, the mechanical properties determine the ability of thermomiotic (negative thermal expansion) materials to counteract positive thermal expansion and affect the stress distributions in polycrystalline A₂M₀₃O₁₂ materials. Therefore, the elastic tensors of Al₂Mo₃O₁₂, Sc₂Mo₃O₁₂, and ZrMgMo₃O₁₂ have been calculated and reported herein. Together with that of Y₂Mo₃O₁₂ (reported previously), we now have elastic constants of an isomorphic family of materials ranging from low positive to large negative thermal expansion.

The study of the anisotropy of the elastic properties of these materials generally has previously been limited to determination of bulk moduli by variable-pressure XRD; the exception is Y₂Mo₃O₁₂. The variable-pressure XRD experiment can yield information about directional compressibilities; for example in Sc₂W₂O₁₂ and Sc₂Mo₂O₁₂ the compressibility along an axis was shown to be positively correlated with the thermal expansion along that axis. However, this technique has limitations; for example the compressibilities in orthorhombic Al₃W₀₃O₁₂ could not be accurately determined due to a pressure-induced phase transition, and no information about shear elasticity was obtained. The use of ultrasonic measurement to obtain

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shear moduli of bulk polycrystalline samples\(^5,16,21\) contributes some additional information, but experimental determination of the full elastic tensor would require single crystals for Brillouin spectroscopy or resonant ultrasound spectroscopy.

Therefore, for practical reasons, computational methods are the best choice to study the full elastic anisotropy of A\(_2\)M\(_3\)O\(_{12}\) materials, and this is the approach taken herein. There is an associated drawback: the DFT calculations are performed for \(T = 0\) K. Since A\(_2\)M\(_3\)O\(_{12}\) materials (except Y\(_2\)M\(_3\)O\(_{12}\)) transition to a monoclinic phase upon cooling, the calculations were performed on a phase which is not stable at absolute zero. Therefore the phononic structures contain vibrational modes with negative energy that destabilize the orthorhombic structure.\(^22–24\)

However, since the orthorhombic–monoclinic phase transition involves rotation of the coordination polyhedra,\(^7\) the unstable mode is located away from the \(\Gamma\) point of the Brillouin zone\(^25,26\) and the instability is not caused by negative terms in the elastic stiffness tensor.\(^27\) Additionally, because many thermomiotic materials have highly temperature-dependent bulk moduli\(^18,28–30\) DFT results will not necessarily match room-temperature elastic constants, but our approach is validated by comparison with experimental behaviour at room temperature.

The energies of the optic phonons at the \(\Gamma\) point were also calculated using DFT. Calculation of the full dispersion relationship would be computationally expensive, however, with the large unit cell of the orthorhombic A\(_2\)M\(_3\)O\(_{12}\) materials (68 atoms)\(^1,5\) there is little dispersion.\(^16,18\) Low-energy optic phonon modes in A\(_2\)M\(_3\)O\(_{12}\) materials corresponding to polyhedral distortions can have large negative Grüneisen parameters,\(^31,32\) and therefore their energies play an important role in the thermal expansion behaviour. Determination of optic phonon frequencies also allows further validation of the calculations by comparison to experimental Raman spectra.

### Methods

DFT calculations were carried out using the ABINIT\(^43\) code in order to determine the elastic tensors and \(\Gamma\)-point phonon energy of A\(_2\)M\(_3\)O\(_{12}\) and AMgM\(_3\)O\(_{12}\) materials. Computation of elastic tensors and \(\Gamma\)-point phonon energies were performed using the ACENET\(^14\) and WestGrid\(^35\) clusters, taking advantage of the massive parallelization functionality of ABINIT.\(^36\) ABINIT composes wavefunctions as linear combinations of a plane wave basis set and performs calculations in reciprocal space.\(^37,38\) Calculations of elastic tensors and phonon frequencies were performed using the response-function capability of ABINIT.\(^39–41\) Phonon frequencies were corrected for errors due to finite sampling on the effective charges by imposition of the acoustic sum rule (i.e., requiring that the frequencies of the acoustic branches are zero at \(\Gamma\)).\(^39\)

The CIF2Cell program\(^42,43\) was used to create input files from published crystal structure data.\(^5,44,45\) PBE GGA exchange–correlation functionals were used in all cases.\(^46\) In the cases of Al\(_2\)Mo\(_3\)O\(_{12}\) and Sc\(_2\)Mo\(_3\)O\(_{12}\), norm-conserving\(^47\) pseudopotentials generated using the OPIUM code\(^47–50\) from the Bennet and Rappe pseudopotential library\(^51\) were used. For ZrMgMo\(_3\)O\(_{12}\), two-projector optimized norm-conserving Vanderbilt pseudopotentials\(^52\) were used. These pseudopotentials were tested by comparison of calculated bulk moduli to experimental values for MgO (\(-7.2\%\) deviation),\(^53\) ZrO\(_2\) (\(-1.6\%\) deviation),\(^54\) and Mo (\(-3.4\%\) deviation),\(^54\) yielding reasonable results.\(^56\) The plane wave cut-off energy and \(k\)-point grid spacing were determined by convergence studies (Table 1); the criterion used was convergence of the internal pressure to within 1%. A \(k\)-point grid spacing smaller than 0.04 Å\(^{-1}\) was used in all cases. The structures were relaxed until the remaining pressure was less than 1 MPa. Changes in the unit cell axis lengths following relaxation are shown in Table 1.

Table 1 Parameters determined by convergence studies and lattice parameter \((a, b, c)\) changes and final pressures following structural relaxation

<table>
<thead>
<tr>
<th>Material</th>
<th>Plane wave cut-off energy/Hartree</th>
<th>(k)-Point grid dimensions</th>
<th>Change in (a) relative to experiment/%</th>
<th>Change in (b) relative to experiment/%</th>
<th>Change in (c) relative to experiment/%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Al(_2)Mo(<em>3)O(</em>{12})</td>
<td>30</td>
<td>(2 \times 2 \times 2)</td>
<td>2.2</td>
<td>1.7</td>
<td>2.0</td>
</tr>
<tr>
<td>ZrMgMo(<em>3)O(</em>{12})</td>
<td>35</td>
<td>(2 \times 3 \times 3)</td>
<td>3.0</td>
<td>2.1</td>
<td>2.0</td>
</tr>
<tr>
<td>Sc(_2)Mo(<em>3)O(</em>{12})</td>
<td>35</td>
<td>(2 \times 3 \times 3)</td>
<td>3.0</td>
<td>2.1</td>
<td>2.1</td>
</tr>
</tbody>
</table>

\(\dagger\) Mo was chosen rather than an oxide as the test material because MoO\(_3\) has a layered structure and therefore is an unsuitable point of comparison,\(^56\) and the bulk modulus of MoO\(_3\) has not been reported in the literature.
Table 2. The thermal expansion tensor and the corresponding directional elastic moduli are shown in Fig. 1. A slice through the microstructure of a modeled polycrystal with randomly oriented grains. The relative orientations of the grains are shown as different colours, with similar colours corresponding to similar orientations. The compliant buffer layer is shown as a grey boundary.

Table 3. The experimental thermal expansion tensor is shown in Fig. 1. A slice through the microstructure of a modeled polycrystal with randomly oriented grains. The relative orientations of the grains are shown as different colours, with similar colours corresponding to similar orientations. The compliant buffer layer is shown as a grey boundary.

Results and discussion

Elasticity

The calculated elastic stiffness tensor (c) of orthorhombic Al₂Mo₃O₁₂ (Pbcn setting) is given by:

\[
c = \begin{bmatrix}
79.1 & 19.3 & 19.3 & 0 & 0 & 0 \\
19.3 & 128 & 51.3 & 0 & 0 & 0 \\
19.3 & 51.3 & 126 & 0 & 0 & 0 \\
0 & 0 & 0 & 45.7 & 0 & 0 \\
0 & 0 & 0 & 0 & 29.8 & 0 \\
0 & 0 & 0 & 0 & 0 & 29.4
\end{bmatrix} \text{ GPa} \tag{1}
\]

and the corresponding directional elastic moduli are shown in Table 2. The thermal expansion tensor (a) from the literature is shown for comparison of its anisotropy to that of c:

\[
a = \begin{bmatrix}
7 & 0 & 0 \\
0 & 0.5 & 0 \\
0 & 0 & -1
\end{bmatrix} \times 10^{-6} \text{K}^{-1}. \tag{2}
\]

Al₂Mo₃O₁₂ showed elastic anisotropy similar to that of its thermal expansion, with only a small difference between the b- and c-axes. The overall elastic anisotropy of Al₂Mo₃O₁₂, expressed as the universal anisotropy parameter, is equal to 1.9. By comparison, Y₂Mo₃O₁₂ has Aₜ = 3.9. Unlike in the case of Y₂Mo₃O₁₂, the axial Young’s moduli and directional compressibilities followed the expected trends and were inversely proportional to each other. The Voight–Reuss–Hill (VRH) values for the bulk elastic constants of Al₂Mo₃O₁₂ were determined to be K = 35.3 GPa and G = 36.2 GPa. These stiffnesses are close to those obtained experimentally for monoclinic Al₂Mo₃O₁₂ (K = 13.5 ± 1.3 GPa, G = 18.2 ± 0.7 GPa). A similar discrepancy was seen in the case of Y₂Mo₃O₁₂, obscuring whether the difference is due to temperature dependence of the elastic constants or the difference between the monoclinic and orthorhombic phases.

The calculated elastic stiffness tensor of orthorhombic ZrMgMo₃O₁₂ (P2₁nb setting) is given by:

\[
c = \begin{bmatrix}
56.3 & 16.8 & 14.2 & 0 & 0 & 0 \\
16.8 & 85.2 & 43.6 & 0 & 0 & 0 \\
14.2 & 43.6 & 88.4 & 0 & 0 & 0 \\
0 & 0 & 0 & 33.9 & 0 & 0 \\
0 & 0 & 0 & 0 & 20.1 & 0 \\
0 & 0 & 0 & 0 & 0 & 18.0
\end{bmatrix} \text{ GPa} \tag{3}
\]

and the corresponding directional elastic moduli are shown in Table 3. The experimental thermal expansion tensor is shown for comparison of its anisotropy to that of c:

\[
a = \begin{bmatrix}
7.6 & 0 & 0 \\
0 & 2.5 & 0 \\
0 & 0 & -4.7
\end{bmatrix} \times 10^{-6} \text{K}^{-1}. \tag{4}
\]

As for Al₂Mo₃O₁₂, ZrMgMo₃O₁₂ shows increased stiffness along its thermomotic axes both in terms of Young’s modulus and compressibility. The stiffnesses of the b- and c-axes were similar even though their CTEs are quite different in magnitude. The overall elastic anisotropy of ZrMgMo₃O₁₂ was very similar to that of Al₂Mo₃O₁₂; both have Aₜ = 1.9. The VRH bulk and shear moduli were 26.5 GPa and 23.8 GPa, respectively. These values are close to those obtained experimentally (K = 31 ± 3 GPa, G = 22 ± 1 GPa).

The calculated elastic stiffness tensor of orthorhombic Sc₂Mo₃O₁₂ (Pbcn setting) is given by:

\[
c = \begin{bmatrix}
53.1 & 14.9 & 16.7 & 0 & 0 & 0 \\
14.9 & 88.1 & 49.3 & 0 & 0 & 0 \\
16.7 & 49.3 & 92.2 & 0 & 0 & 0 \\
0 & 0 & 0 & 31.8 & 0 & 0 \\
0 & 0 & 0 & 0 & 15.8 & 0 \\
0 & 0 & 0 & 0 & 0 & 18.5
\end{bmatrix} \text{ GPa}. \tag{5}
\]
and the corresponding directional elastic moduli are shown in Table 4. The experimental thermal expansion tensor \( \beta \) is shown for comparison of its anisotropy to that of \( c \):

\[
\alpha = \begin{bmatrix}
11 & 0 & 0 \\
0 & -8.7 & 0 \\
0 & 0 & -8.4
\end{bmatrix} \times 10^{-6} \text{K}^{-1}.
\]  

(6)

The elastic anisotropy of \( \text{Sc}_2\text{Mo}_3\text{O}_{12} (A_1 = 2.9) \) was larger than that of \( \text{Al}_2\text{Mo}_3\text{O}_{12} \) and \( \text{ZrMgMo}_3\text{O}_{12} \), which could be expected given its larger CTE anisotropy. The VRH values for the bulk and shear moduli were \( K = 24.0 \text{ GPa} \) and \( G = 22.3 \text{ GPa} \); unlike in the cases of \( \text{Al}_2\text{Mo}_3\text{O}_{12} \) and \( \text{Y}_2\text{Mo}_3\text{O}_{12} \), the calculated bulk modulus of \( \text{Sc}_2\text{Mo}_3\text{O}_{12} \) is significantly less than the experimental value \( (K = 32 \pm 2 \text{ GPa} \text{ from variable-pressure XRD}) \).\(^\text{14}\)

Correlations of axial elastic properties with axial thermal expansion, axial Young’s moduli \( (\gamma_\alpha) \), axial compressibilities \( (\beta_\alpha) \), and directional shear moduli \( (G_\alpha\beta) \) are visualized in Fig. 2 as functions of the axial CTEs \( (\gamma_\alpha)^{2,5,6}\) for the three materials described above and \( \text{Y}_2\text{Mo}_3\text{O}_{12} \).\(^\text{3,16}\) The elastic behaviours of the thermomiotic \( b \)- and \( c \)-axes are correlated with their axial CTEs, while such a correlation is not seen for the \( a \)-axis. The \( b \)- and \( c \)-axes are stiffer than the \( a \)-axis except in the case of \( \text{Y}_2\text{Mo}_3\text{O}_{12} \), where the \( a \)-axis has a higher Young’s modulus but also higher compressibility. Stiffer thermomiotic axes could decrease the bulk CTE of a polycrystal, as the increased stiffness would yield a larger contribution to the bulk expansion. As described below, this effect can be predicted in some cases. The trends in the VRH-averaged isotropic moduli (Fig. 2(d)) are more pronounced than those in the axial stiffnesses. A decrease in overall stiffness with increasingly negative thermal expansion can be seen clearly. When a thermomiotic material is used to counteract positive thermal expansion, its stiffness can be as important as its CTE in determining the amount of CTE reduction achieved,\(^\text{15,20,21,63}\) and therefore the negative correlation between stiffness and CTE might act to discourage the use of materials such as \( \text{Y}_2\text{Mo}_3\text{O}_{12} \) in applications. The trend relating the elastic moduli to the CTE is very similar to the previously reported correlation between the A site ionic force and the CTE,\(^\text{5}\) suggesting that polyhedral distortability is a common factor between thermal expansion and elastic properties in this group of materials.

The materials studied show significantly different behaviour when subjected to different stress conditions. The \( b \)- and \( c \)-axes

### Table 4 Directional elastic moduli of \( \text{Sc}_2\text{Mo}_3\text{O}_{12} \) (Pbcn setting)

<table>
<thead>
<tr>
<th></th>
<th>Young’s modulus/GPa</th>
<th>Shear modulus/GPa</th>
<th>Compressibility/ GPa(^{-1})</th>
<th>Poisson ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>( Y_{\alpha\alpha} )</td>
<td>49.5</td>
<td>31.8</td>
<td>( \beta_{\alpha\alpha} = 1.57 \times 10^{-2} )</td>
<td>0.51</td>
</tr>
<tr>
<td>( Y_{\beta\beta} )</td>
<td>60.9</td>
<td>15.8</td>
<td>( \beta_{\beta\beta} = 6.00 \times 10^{-3} )</td>
<td>0.13</td>
</tr>
<tr>
<td>( Y_{\gamma\gamma} )</td>
<td>63.1</td>
<td>18.5</td>
<td>( \beta_{\gamma\gamma} = 4.80 \times 10^{-3} )</td>
<td>0.10</td>
</tr>
</tbody>
</table>

Fig. 2 Calculated directional elastic properties of (a) Young’s moduli, (b) compressibilities, and (c) shear moduli, as functions of axial thermal expansion along the \( a \)- ( ), \( b \)- ( ), and \( c \)- ( ) axes for orthorhombic \( \text{Al}_2\text{Mo}_3\text{O}_{12} \), \( \text{ZrMgMo}_3\text{O}_{12} \), \( \text{Sc}_2\text{Mo}_3\text{O}_{12} \), and \( \text{Y}_2\text{Mo}_3\text{O}_{12} \) in the \( \text{Pbcn} \) or \( \text{P}2_1\text{nb} \) setting. (d) Volumetric thermal expansion as a function of calculated VRH bulk and shear moduli for orthorhombic \( \text{Al}_2\text{Mo}_3\text{O}_{12} \), \( \text{ZrMgMo}_3\text{O}_{12} \), \( \text{Sc}_2\text{Mo}_3\text{O}_{12} \), and \( \text{Y}_2\text{Mo}_3\text{O}_{12} \). Axial CTEs and the elastic tensor of \( \text{Y}_2\text{Mo}_3\text{O}_{12} \) were taken from the literature; see the text for references.
show a decrease in uniaxial stiffness (Fig. 2(a)) with increasingly negative thermal expansion. However, when subjected to isotropic pressure (Fig. 2(b)), less variation in the stiffness is seen. These differences illustrate the importance of computational studies of the full elastic tensor, as only $b_{ii}$ can be obtained from experimental variable-pressure XRD measurements on polycrystalline samples. The directional shear moduli generally show increased stiffness for $G_{23}$ by comparison with $G_{13}$ and $G_{12}$, which is consistent with the increased stiffness of the $b$- and $c$-axes compared to the $a$-axis. This trend leads to the shear modulus in a plane ($G_{ij}$) increasing with the thermal expansion of the axis perpendicular to the plane ($a_{ii}$).

The anisotropic Young’s moduli of Al$_2$Mo$_3$O$_{12}$, ZrMgMo$_3$O$_{12}$, Sc$_2$Mo$_3$O$_{12}$, and Y$_2$Mo$_3$O$_{12}$ are compared in all directions in Fig. 3. The overall trend of decreased overall stiffness with decreased CTE can be seen clearly, as can variations in the elastic anisotropy. In the case of Al$_2$Mo$_3$O$_{12}$, the predominant feature is the increased stiffness of the thermomiotic $bc$-plane relative to the $a$-axis. However, as the thermal expansion of the material becomes more negative, the stiffness of the $bc$-plane decreases more dramatically than that along the $a$-axis. Additionally, the stiffness within the $bc$-plane becomes more anisotropic, with the [011] and [011] directions becoming considerably stiffer than [010] and [001]. This results in the relatively large values of $G_{23}$ and discrepancies between $Y_{ii}$ and $b_{ii}$ seen in Fig. 2 and the indentations along [010] and [001] seen in Fig. 3.

Phonon frequencies

The $\Gamma$-point optic phonon frequencies calculated in ABINIT are presented in tabular form in the ESI,† and visually in Fig. 4 (along with the previously reported phonon spectrum of Y$_2$Mo$_3$O$_{12}$). Fig. 4 shows the common features of the phonon spectra of the four materials studied: a band of low-energy librational, translational, and bending modes separated from stretching modes with higher energy, with asymmetric stretches at lower energies than symmetric stretches. As discussed in ref. 5, ZrMgMo$_3$O$_{12}$ has a larger spread of stretching mode energies due to the splitting of the A site into a Zr site and Mg site, but otherwise its $\Gamma$-point phonon spectrum is similar to that of the Al$_2$Mo$_3$O$_{12}$ materials. The energies of the low-energy phonons can be compared more easily in Fig. 4, which shows the cumulative distribution function of the modes below 500 cm$^{-1}$. The phonons below ca. 200 cm$^{-1}$ are expected to have negative mode Grüneisen parameters and contribute to NTE, while those of higher energies are expected to have positive or near-zero Grüneisen parameters.

Fig. 4 shows that Y$_2$Mo$_3$O$_{12}$ has significantly more modes with energies below 200 cm$^{-1}$ and has generally lower phonon energies in this region, which is consistent with its observed NTE over a wide temperature range. Conversely, Al$_2$Mo$_3$O$_{12}$ has considerably fewer modes below 200 cm$^{-1}$, with a much broader distribution of its phonon frequencies in the region below 500 cm$^{-1}$. Of particular interest is a mode at 26 cm$^{-1}$ in Al$_2$Mo$_3$O$_{12}$ which is anomalously low in energy by comparison to the other translational and librational modes in this material. At some point in the Brillouin zone, this mode could be the soft mode with negative frequency that causes the orthorhombic–monoclinic phase transition. The mode distributions for Sc$_2$Mo$_3$O$_{12}$ and ZrMgMo$_3$O$_{12}$ are very similar, especially in the region below 200 cm$^{-1}$. Sc$_2$Mo$_3$O$_{12}$ has a negative CTE while ZrMgMo$_3$O$_{12}$ displays ca. zero thermal expansion, implying that the mode Grüneisen parameters of Sc$_2$Mo$_3$O$_{12}$ are more negative.
The calculated $\Gamma$-point phonon frequencies of Al$_2$Mo$_3$O$_{12}$ and ZrMgMo$_3$O$_{12}$ are compared to experimental Raman spectra (measured at room temperature) in the ESI.$^{\dagger}$ Overall, the results of this comparison, in addition to the comparison of experimental elastic moduli to calculated values presented above, show that the calculated elastic and phononic properties of these materials, while not quantitatively accurate, show reasonable agreement with the experimental data.

Axial Grüneisen parameters

The axial coefficients of thermal expansion are determined by the anharmonicity of the interatomic potentials, as expressed through Grüneisen parameters, and by the stiffness tensor, which represents the resistance of the lattice to thermal deformation. The present computational analysis does not allow the axial Grüneisen parameters ($\gamma_{ii}$) to be determined directly, but using the calculated anisotropic elastic constants and phonon energies and the experimental CTE tensor the following expression can be invoked:67

$$\gamma_{ii} = \frac{V_m}{C_V} (c_{ii} \xi_{ii} + c_{ij} \xi_{ij} + c_{ik} \xi_{ik})$$

(7)

where $V_m$ is the molar volume, $C_V$ is the isochoric heat capacity, and $c_{ij}$ and $\xi_{ij}$ are stiffness and thermal expansion tensor elements, respectively. Here, $C_V$ was calculated using the Einstein and Debye models following the procedure described in ref. 68 using the calculated elastic tensors and $\Gamma$-point phonon frequencies.$^{16}$ The axial Grüneisen parameters were determined at $T = 300$ K for ZrMgMo$_3$O$_{12}$, Sc$_2$Mo$_3$O$_{12}$, and Y$_2$Mo$_3$O$_{12}$,$^{9,16}$ and at $T = 550$ K for Al$_2$Mo$_3$O$_{12}$ (temperatures chosen so that the materials are in the orthorhombic phase). The results are shown in Fig. 5 and Table 5.

Thermal expansion along the $b$- and $c$-axes strongly correlates with the Grüneisen parameter along that axis, suggesting that vibrational modes with negative Grüneisen parameters along those axes are responsible for NTE. Essentially, this high degree of correlation is because the $c_{ij} \xi_{ij}$ and $c_{ik} \xi_{ik}$ terms in eqn (7) have opposite signs and nearly cancel. These terms are due to the Poisson effect, where a strain (in this case, thermally induced) along one axis causes a corresponding strain along the axes perpendicular to it.$^{69}$ As shown in Tables 2–4, some of the axial Poisson ratios in these materials are quite large, close to the theoretical limit of 0.5 for stable isotropic materials.$^{70}$ However, NTE along the $b$- and $c$-axes increases the CTE along
Y2Mo3O12 can be seen in Fig. 6. The stress extrema are found at the grain boundaries where there are significant differences in the elastic properties of the grains. An example cross section of the thermal stress distribution in Y2Mo3O12 is shown in Fig. 5. The stress distributions within the polycrystals were fit to statistical distributions as described in the ESI.† The distributions in the different materials are generally similar, and using two lognormal distributions for the stress along each axis provides excellent fits to the data (see Fig. S4–S11, ESI†). As the size of the A site cation increases from Al3+ to Y3+, increasing thermal expansion anisotropy is accompanied by decreasing stiffness (vide supra). These two factors have opposite effects on the magnitude of the thermal stresses, resulting in similar stress distributions and extrema. This similarity indicates that it will prove difficult to use chemical control of the A site cation to reduce thermal stress due to thermal expansion anisotropy. When the thermal stresses are sufficiently large to cause microcracking, a further deviation from the intrinsic CTE can be found at elevated temperatures due to crack growth and healing

Table 6 Mean strain energy densities and deviations from the intrinsic linear CTE as predicted for the elastically isotropic case (where Δσ/σ = 0) by Kreher’s method,71,72 and the present models

<table>
<thead>
<tr>
<th>Material</th>
<th>Elastic isotropy (W/kg m⁻³)</th>
<th>Present models</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(W)</td>
<td>Δσ/10⁻⁷ K⁻¹</td>
</tr>
<tr>
<td>Al₂Mo₃O₁₂</td>
<td>311</td>
<td>-2.5</td>
</tr>
<tr>
<td>ZrMgMo₂O₁₂</td>
<td>493</td>
<td>-3.5</td>
</tr>
<tr>
<td>Sc₂Mo₂O₁₂</td>
<td>1338</td>
<td>-11</td>
</tr>
<tr>
<td>Y₂Mo₂O₁₂</td>
<td>837</td>
<td>-1.3</td>
</tr>
</tbody>
</table>

The effect of elastic anisotropy on the thermal stress distributions can be quantified by comparison to Kreher’s model for elastically isotropic polycrystals.71,72 Anisotropic models have lower mean strain energy densities, ⟨W⟩, as shown in Table 6, indicating that elastic anisotropy acts to decrease thermal stress. In the elastically isotropic case, the average strain in the material is required to be equal to the thermal strain by the (unconstrained) boundary conditions and the conditions of static equilibrium.71,72 When the coupling of elastic anisotropy and thermal expansion anisotropy is included, the thermal expansion of the stiffer axes is slightly more expressed than that of the more compliant axes, which allows some relaxation of the thermal stress. This effect is seen in consolidated polycrystals as deviations from the intrinsic linear CTE (Δσ/σ), shown in Table 6. While Δσ/σ is small in comparison to the bulk CTEs of Al₂Mo₃O₁₂ and Y₂Mo₂O₁₂, for ZrMgMo₂O₁₂ and Sc₂Mo₂O₁₂ the deviation is of similar magnitude to the intrinsic CTE. Therefore, elastic anisotropy could explain the relatively large differences observed in dilatometric results compared with the intrinsic CTE (e.g. from XRD) for these materials.25
causing expansion and shrinkage, respectively, of the polycrystalline body.19,20

In all cases, the two lognormal distributions in each fit had opposite signs for their prefactors, leading to positive and negative stress extrema each being fit by a single unbounded lognormal tail. The utility of the lognormal distribution in fitting the extreme stresses is likely connected to some underlying physical phenomenon. The lognormal distribution is known to arise as a result of multiplication of a series of random variables, whereas the normal distribution arises from summation of a series of random variables.73 Therefore, the lognormal fit to the stress distribution implies some multiplicative character to their origin. In anisotropic polycrystalline materials, thermal expansion mismatches cause strains in the material, and consequently reaction forces and stresses. In elastically isotropic materials, the interactions of reaction forces throughout the material are isotropic. This isotropy results in the stresses at any point being related to the strains caused by thermal expansion anisotropy throughout the remainder of material additively, and the normal thermal stress distribution is a logical result. However, in a material where the elastic constants are anisotropic, the transfer of stresses through the material is not so simple. The elastic anisotropy leads to the grains reacting differently to the applied strains based on their orientations, with increased stress in stiffer directions and vice versa. Therefore, elastic anisotropy introduces a multiplicative factor into the thermal stress distributions, since the relationship between the stress at a point and the strain at another point is modified by the variations in the stiffness tensor.

Conclusions

Materials in the A2Mo3O12 and AMgMo3O12 families were shown to have significant elastic anisotropy, a finding consistent with their characteristically large CTE anisotropy. The thermomiotic axes of these materials were shown to be stiffer than the PTE axes, which could act to decrease their bulk CTEs. However, the stiffness along the thermomiotic axes decreases with increasingly negative thermal expansion. The stiffness along the PTE axes did not strongly correlate with their CTEs. The stiffness along the thermomiotic axes of these materials were shown to be stiffer than the PTE axes, which could act to decrease their bulk CTEs. The elastic tensors were found to couple with the thermal expansion tensors to produce small deviations in the bulk CTE, which were large by comparison to the intrinsic CTE in the cases of ZrMgMo3O12 and Sc2Mo3O12.

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References

33 ABINIT, http://www.abinit.org, accessed 02/05/16.
35 WestGrid – Western Canada Research Grid, https://www.westgrid.ca, accessed 02/05/16.
42 CIF2Cell, http://sourceforge.net/projects/cif2cell/, accessed 02/05/16.
51 Rappe Group, http://www.sas.upenn.edu/rappegroup/htdocs/Research/psp_gga.html, accessed 02/05/16.