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Pressure induced octahedral tilting distortion in Ba$_2$YTaO$_6$†

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Herein we communicate the first example of a pressure induced octahedral tilting distortion in a double perovskite phase, which was observed during the structural characterization of Ba$_2$YTaO$_6$ using high-pressure synchrotron X-ray powder diffraction.

Perovskites and structurally related oxides have been one of the most widely studied structure types because of their interesting physical properties and extensive structural diversity. The perovskite oxide has the general stoichiometry ABO$_3$ and is composed of corner sharing BO$_6$ octahedra with the A site cation occupying the void created by the three dimensional octahedral framework. Although many compounds adopt the ideal cubic perovskite aristotype, most perovskites undergo a distortion away from the high symmetry cubic structure. It is these subtle structural distortions, such as a cooperative octahedral tilting distortion that occurs when the A site cation is undersized for the corner sharing network, that often greatly influence the physical properties. Research on ordered double perovskites has been pursued by a large number of researchers, in part, because of the diverse physical properties and remarkable structural chemistry observed in these materials. Octahedral tilting in cation ordered double perovskites has been examined in detail by several groups using a large number of techniques, including group theoretical analysis. While temperature is the most common intensive variable utilized in the study of phase transitions in these materials, pressure also may induce a structural phase transition; an example being the first order rhombohedral (R$\bar{3}$) to monoclinic (I$\bar{2}$m) transition in Ba$_3$BiTiO$_6$ that can be induced by either lowering the temperature or through the application of pressure.

Here we describe the first example of a pressure induced octahedral tilting distortion in Ba$_2$YTaO$_6$, observed by variable pressure synchrotron X-ray powder diffraction experiments on Ba$_2$YTaO$_6$ were performed using a diamond anvil cell at the X7A beam line at the National Synchrotron Light Source at Brookhaven National Laboratory. The experimental setup and detailed procedure are described elsewhere. Diffraction patterns were collected using a wavelength of 0.62245 Å. Diffraction data analysis was performed using the EXPGUI interface of GSAS. A pseudo Voigt function was used to model the peak shape. An angular range employed in the refinements was from 6° to 35° 20, and excluded the regions that contained strong Bragg peaks from the pressure cell gasket.

Inspection of the variable pressure data above 4.3 GPa indicated that the diffraction peaks of certain reflections began to split. An example of this peak splitting is the 620 peak in space group Fm$\bar{3}$m near 27.2° 20. Fig. 1 clearly shows the splitting of the peak at pressures of 5.6 GPa and 6.3 GPa, compared to the single reflection at 4.3 GPa. The splitting of the reflections is well described by a cubic to tetragonal phase transition and is consistent with a phase transition to either space group P4/mncc or I$\bar{4}$m. Reflections that violate the body centering conditions, such as the 212 reflection shown to be indicative of primitive symmetry in Ba$_3$PrIr$_x$Ru$_{1-x}$O$_6$ were absent. Refinements were attempted using structure models generated by the SPrDS software in space groups I$\bar{4}$m and P4/mncc. The fit was superior in space group I$\bar{4}$m compared to P4/mncc. A phase transition from cubic (Fm$\bar{3}$m) to tetragonal (I$\bar{4}$m) was also reported to occur at 253(1) K; therefore the pressure induced phase transition from Fm$\bar{3}$m to I$\bar{4}$m is the same as that which occurs on lowering of the temperature. Crystal structures shown in Fig. 2 were refined from the 4.3 GPa and 6.3 GPa data in space groups Fm$\bar{3}$m and I$\bar{4}$m, respectively.

![Fig. 1 Splitting of the 620 peak in Fm$\bar{3}$m to the 116/332/240 peaks in I$\bar{4}$m at P = 4.3, 5.6 and 6.3 GPa. The bottom pattern was fitted with space group Fm$\bar{3}$m, the top and middle patterns with I$\bar{4}$m.](Image)
This pressure induced phase transition is most clearly under
stood in terms of an octahedral tilting distortion. The untitled ideal
cubic ordered double perovskite crystallizes in space group \textit{Fm\textit{3}m}. In space group \textit{I\textit{4}lm}, an antiphase octahedral tilting distortion
occurring about the \(c\) axis is clearly shown in Fig. 2. The
octahedral tilting phase transition allows for an additional means
of compression of the structure. In perovskites, the two
dominating processes during compression are bond shortening
and octahedral tilting. In the cubic form of Ba\textsubscript{2}YTaO\textsubscript{6}, compression with retention of the cubic symmetry is possible only \textit{via} the
simultaneous shortening of the Y O, Ta O and Ba O bonds. In
its tetragonal form, on the other hand, both bond shortening and
octahedral tilting can take place. However, while a decrease in the
\(c\) lattice parameter is the result of only the compressive shortening of the
Y O, Ta O and Ba O bond distances, a reduction of the \(a\) lattice parameter can have contributions from both bond compression and octahedral tilting.

In an ideal \textit{I\textit{4}lm} perovskite crystal structure with symmetric
octahedra and \(0^\circ\) octahedral tilting, the \(2a\) and \(c\) lattice
parameters are equal and the \(c/(2a)\) ratio is unity. The observation
of a larger relative compression of the \(a\) lattice parameter compared to the \(c\) lattice parameter, that is when \(c/(2a)\) becomes larger than unity, is indicative of an additional compression mechanism, namely octahedral tilting.

Octahedral tilting has a primary effect on the Ba O bond
lengths, with a minimal change in the octahedral cation bond
lengths. If the octahedral bond distances remain regular, then with
an increase in the magnitude of octahedral tilting the \(a\) lattice
parameter becomes smaller while the \(c\) lattice parameter remains
unchanged. Therefore, we can conclude that the observed increase in the
\(c/(2a)\) ratio in Ba\textsubscript{2}YTaO\textsubscript{6} from 1.0025 to 1.0029 upon a
change in the applied pressure from 5.6 to 6.3 GPa, represents a
greater degree of octahedral tilting. The average tilt angle of the
[YO\textsubscript{6}] and [TaO\textsubscript{6}] octahedra is 4.1\(^\circ\) at 6.3 GPa.

One cause for such a tilting transition is a difference in the
relative compressibilities of the A X vs. B X bonds in an ABX\textsubscript{3}
perovskite. For example, a recent study of the perovskite NaMgF\textsubscript{3}
demonstrated that the compression mechanism was dominated by
a shortening of the Mg F bonds below 6 GPa and controlled by
an increase in octahedral tilting above 12 GPa; however, in that experiment no change in the type of octahedral tilting was observed. We have obtained similar results in this study of Ba\textsubscript{2}YTaO\textsubscript{6}, where bond compression is the dominant mechanism at lower pressures, while octahedral tilting becomes more important at higher pressures. These results also suggest that the
Ba O bonds are more compressible than the Ta O and Y O bonds, which is evident in the onset of the octahedral tilting transition.

The pressure dependence of the lattice parameters and unit cell
volume is given in Table 1. Additional information on the Rietveld refinements in space group \textit{Fm\textit{3}m} and in space group \textit{I\textit{4}lm} at
6.3 GPa are given in the ESI. Group theoretical analysis has shown the
\textit{Fm\textit{3}m} to \textit{I\textit{4}lm} phase transition can be continuous. The pressure dependence of the lattice parameters is shown in Fig. 3. A
second order Birch Murnaghan equation of state was fit to the pressure dependence of the unit cell volume data between 0.9 and
4.3 GPa, giving a bulk modulus of 157(16) GPa and \(V_o\)
601.0(9) \(\text{Å}^3\). The bulk modulus is comparable to that found in
other double perovskites, for example Ba\textsubscript{2}BiTaO\textsubscript{6} at 126(10) GPa.

![Fig. 2](image)

**Fig. 2** Crystal structures of the \textit{Fm\textit{3}m} (top) and \textit{I\textit{4}lm} (bottom) forms of Ba\textsubscript{2}YTaO\textsubscript{6}. Large orange spheres represent Ba, green spheres Y, red spheres are Ta, and blue spheres O.

<table>
<thead>
<tr>
<th>Pressure/GPa</th>
<th>Space group</th>
<th>(a/\text{Å})</th>
<th>(c/\text{Å})</th>
<th>Volume/(\text{Å}^3)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.9</td>
<td>\textit{Fm\textit{3}m}</td>
<td>8.42811(8)</td>
<td>598.67(2)</td>
<td></td>
</tr>
<tr>
<td>1.7</td>
<td>\textit{Fm\textit{3}m}</td>
<td>8.4077(2)</td>
<td>594.33(5)</td>
<td></td>
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<tr>
<td>2.2</td>
<td>\textit{Fm\textit{3}m}</td>
<td>8.39707(8)</td>
<td>592.08(2)</td>
<td></td>
</tr>
<tr>
<td>3.1</td>
<td>\textit{Fm\textit{3}m}</td>
<td>8.38092(9)</td>
<td>588.67(2)</td>
<td></td>
</tr>
<tr>
<td>4.3</td>
<td>\textit{Fm\textit{3}m}</td>
<td>8.37435(9)</td>
<td>587.29(2)</td>
<td></td>
</tr>
<tr>
<td>5.6</td>
<td>\textit{I\textit{4}lm}</td>
<td>8.5829(2)</td>
<td>8.3545(4)</td>
<td>290.11(2)</td>
</tr>
<tr>
<td>6.3</td>
<td>\textit{I\textit{4}lm}</td>
<td>5.8851(1)</td>
<td>8.3474(2)</td>
<td>289.10(1)</td>
</tr>
</tbody>
</table>

![Fig. 3](image)

**Fig. 3** Pressure variation of the lattice parameters. Circles represent \(a\) in \textit{Fm\textit{3}m}, squares \(c\) in \textit{I\textit{4}lm}, and triangles \(2a\) in \textit{I\textit{4}lm}.
Ba$_2$PrRu$_{0.8}$Ir$_{0.2}$O$_6$ at 139(10) GPa,$^9$ and Sr$_2$TbRu$_{0.3}$Ir$_{0.7}$O$_6$ at 196(10) GPa.$^{10}$

In conclusion, we have demonstrated that a phase transition from space group $Fm\bar{3}m$ to $I4/m$ occurs upon application of pressure to Ba$_2$YTaO$_6$ with the onset of an octahedral tilting distortion about the c axis. The phase transition to space group $I4/m$ marks an additional compression mechanism consisting of simultaneous octahedral tilting and bond shortening in contrast to bond shortening only in space group $Fm\bar{3}m$.

Notes and references

† Polycrystalline Ba$_2$YTaO$_6$ was prepared using solid state techniques by reacting Y$_2$O$_3$ and Ta$_2$O$_5$ to form a phase pure YTaO$_4$ precursor, which was then appropriately mixed with BaCO$_3$ and reacted until a single phase sample of Ba$_2$YTaO$_6$ was obtained, as established by laboratory powder X ray diffraction. The laboratory X ray powder diffraction pattern could be indexed with a cubic cell, space group $Fm\bar{3}m$, with $a = 8.42$ Å.  
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