Elastic anisotropy of experimental analogues of perovskite and post-perovskite help to interpret D'' diversity

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Recent studies show that the D'' layer, just above the Earth’s core-mantle boundary, is composed of MgSiO3 post-perovskite and has significant lateral inhomogeneity. Here we consider the D'' diversity as related to the single-crystal elasticity of the post-perovskite phase. We measure the single-crystal elasticity of the perovskite Pbnm-CaIrO3 and post-perovskite Cmcm-CaIrO3 using inelastic X-ray scattering. These materials are structural analogues to same phases of MgSiO3. Our results show that Cmcm-CaIrO3 is much more elastically anisotropic than Pbnm-CaIrO3, which offers an explanation for the enigmatic seismic wave velocity jump at the D'' discontinuity. Considering the relation between lattice preferred orientation and seismic anisotropy in the D'' layer, we suggest that the c axis of post-perovskite MgSiO3 aligns vertically beneath the Circum-Pacific rim, and the b axis vertically beneath the Central Pacific.
Since the discovery of the transition from perovskite (Pv) to post-perovskite (pPv) in MgSiO₃ (refs 1,2), pPv-MgSiO₃ has been considered to be a major component of the D′ layer just above the core–mantle boundary (CMB). Seismological observations of the D′ layer are difficult to interpret as they suggest both non-uniform response at the layer boundary and non-uniform anisotropy inside the layer3–5. With this background, elastic anisotropy or single-crystal elasticity of pPv-MgSiO₃ has been the focus of theoretical calculations6–10. However, no experimental data are available on the single-crystal elasticity of pPv-MgSiO₃, because it is unstable at ambient pressure. Cmcm-CaIrO₃ or pPv-CaIrO₃ has been frequently selected as a representative analogue of pPv-MgSiO₃ (refs 11–13). According to the phase diagram14, pPv-CaIrO₃ is the stable phase at ambient conditions and Pv-CaIrO₃ is the high-temperature and low-pressure phase. Here we report the first experimental data on single-crystal elasticity of Pv and pPv structures in a homogeneous chemical composition of CaIrO₃.

Results
Characterization of single-crystal specimen. Figure 1a,b shows single crystals of Pv-CaIrO₃ and pPv-CaIrO₃ used in the present study. The Pv-CaIrO₃ sample was synthesized at 2 GPa and 1,450 °C for 15 h in a piston cylinder apparatus at Okayama University. We produced many grains with size of a few tens of microns, although most of them turned out to have multiple domains. We examined more than 20 grains using a four-circle X-ray diffractometer at Yamaguchi University and selected a quasi-single-domain crystal with size of ~20 μm. Its lattice constants were determined as \(a = 5.3527(1), \ b = 5.5969(5)\) and \(c = 7.6804(6)\) Å, which yields density (\(\rho\)) of 8,091 kg m\(^{-3}\). The pPv-CaIrO₃ sample was synthesized by slow cooling from 1,000 °C in CaCl₂ flux\(^{15}\). It had a pine-needle shape with length of \(\sim 100\) microns and width of a few 10s of microns. Its lattice constant and densities were respectively \(a = 3.145(2), \ b = 9.861(6), \) and \(c = 7.297(5)\) Å, and \(\rho = 8,211\) kg m\(^{-3}\), which are consistent with literature values\(^{16}\).

Inelastic X-ray scattering measurement and results. Figure 1c,d compare experimental velocities obtained from the inelastic X-ray scattering (IXS) spectra with velocities calculated from the optimized elastic constants. The details of the IXS measurement are described in the Methods section for the measurement of IXS at SPring-8 BL35XU; one of the important advantages of IXS compared with Brillouin scattering is its geometrical freedom (see Supplementary Methods). The analytical procedures for determining single-crystal elastic constants are described in the Methods section for data analysis for elastic constants. The measurement data are summarized in Supplementary Tables 1 and 2.

Table 1 summarizes the results of elastic constants and gives isotropic averages. The values are reasonably consistent with the theoretical results\(^{17}\) except for a few discrepancies. The high-pressure phase of pPv-CaIrO₃ is less compressible than Pv-CaIrO₃, whereas the theoretical prediction suggests the opposite relation. In addition, isotropic averages of \(V_p\) and \(V_s\) are significantly larger for pPv-CaIrO₃ than for Pv-CaIrO₃ according to the present results. Figure 2 compares the present results and theoretical calculations for the velocity surfaces of Pv-CaIrO₃ and pPv-CaIrO₃. We see fairly good agreement between the two results for pPv-CaIrO₃, although we recognize significant differences between them for Pv-CaIrO₃. Figure 3 shows the compatibility of the elastic constants with compression data\(^{18}\). The present data are consistent with the compression data for Pv- and pPv-CaIrO₃; the theoretical result for Pv-CaIrO₃, however, yields a different slope from the others.

![Figure 1](image_url)  
**Figure 1 | Specimen and measurement data.** Left and right columns refer to Pbnm- and Cmcm-CaIrO₃, respectively. (a,b) Photographs of single crystals used. The horizontal edge lengths are ~5 mm and ~800 μm, respectively, for a and b. (c,d) Overall picture for the experimental data and analytical fitting. Black lines indicate the direction of wave propagation in the crystal, and blue triangles and red circles are experimental velocities and recalculated values after least-squares analysis. The vertical direction is positive z direction, while horizontal leftward and rightward directions are positive x and y directions, respectively. Note that the difference between experimental and recalculated velocities is less than 5%. The misfit uniformly distributed indicates untwinned crystal specimens.
Discussion

Figure 4 presents stereo projections of elastic wave velocities for Pv and pPv structures of CaIrO$_3$. It shows that the ranges of variation of $V_P$, the average of $V_S$ values, and the difference in $V_S$ values are $\pm 600$, $\pm 300$, and $\pm 500$ m s$^{-1}$ for Pv-CaIrO$_3$ and $\pm 1,000$, $\pm 800$ and $\pm 1,100$ m s$^{-1}$, for pPv-CaIrO$_3$, respectively. We see that pPv-CaIrO$_3$ has approximately double the anisotropy of Pv-CaIrO$_3$ in terms of the elastic wave velocity. For example, anisotropy in the polarization shear wave velocity is $\pm 30\%$ in pPv-CaIrO$_3$ while it is only $\pm 15\%$ in Pv-CaIrO$_3$.

From Fig. 4d,e for pPv phase, the $b$ axis is found to be the direction of the lowest $V_P$, while the highest $V_S$ is between the $a$ and $c$ axes. These pronounced features of the pPv elasticity are understandable as alternate stacking of compliant and stiff layers perpendicular to the $b$ axis; the stiff layer has the largest shear wave constant and lowest longitudinal wave constant between $a$ and $c$ axes. We recognize similarity between $a$ and $c$ axes in pPv-CaIrO$_3$ elasticity, while the $a$ axis is slightly stiffer than the $c$ axis in terms of both longitudinal and shear wave velocities.

Table 1 | Single-crystal elastic moduli of Pbnm- and Cmcm-CaIrO$_3$.

|       | $C_{11}$ (GPa) | $C_{22}$ (GPa) | $C_{33}$ (GPa) | $C_{12}$ (GPa) | $C_{23}$ (GPa) | $C_{31}$ (GPa) | $C_{44}$ (GPa) | $C_{55}$ (GPa) | $C_{66}$ (GPa) | $K_R$ (GPa) | $K_H$ (GPa) | $G_H$ (GPa) | $V_P$ (m s$^{-1}$) | $V_S$ (m s$^{-1}$) |
|-------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|-------------|-------------|-------------|----------------|----------------|
| Pbnm  | 235 (6)        | 278 (6)        | 286 (11)       | 132 (6)        | 138 (11)       | 120 (10)       | 87 (4)         | 60 (2)         | 79 (2)         | 173 (6)     | 174 (6)     | 72 (2)      | 5,776 (54)    | 2,976 (42)     |
|       | 248            | 359            | 319            | 204            | 162            | 115            | 79             | 42             | 97             | 195         | 202         | 68          | 6,001         | 2,892          |
| Cmcm  | 378 (4)        | 255 (11)       | 360 (5)        | 104 (7)        | 137 (9)        | 73 (6)         | 76 (5)         | 56 (2)         | 85 (3)         | 178 (4)     | 179 (4)     | 84 (1)      | 5,955 (46)    | 3,201 (27)     |
|       | 388            | 241            | 386            | 121            | 149            | 85             | 67             | 46             | 64             | 186         | 189         | 98          | 5,913         | 2,990          |

Several isotropic moduli and velocities are shown on the right-hand side. The upper two rows are for Pbnm or Pv structure, while the lower two rows are for Cmcm or pPv structure. In each case, the first row gives the present experimental results with accuracy and the second row the theoretical results obtained by Tsuchiya and Tsuchiya$^{17}$. The unit is ‘GPa’ except for velocities which have a unit of ‘m s$^{-1}$’ in the last two columns. The numbers in parentheses correspond to fitting error in the least square analysis.
The elasticity of Pv-CaIrO$_3$ resembles that of Pv-MgSiO$_3$ under lower-mantle conditions more than under ambient conditions (Fig. 5). Elasticity of Pv- and pPv-CaIrO$_3$ under ambient conditions seems to be a good indicator of that of Pv- and pPv-MgSiO$_3$ under lowermost mantle conditions; a conversion factor of 2.2–2.4 is found between them.

We interpret the D'' diversity observed seismologically in terms of the single-crystal elasticity of Pv and pPv-CaIrO$_3$. Significant anisotropy of the seismic wave velocity has been observed in the D'' layer, while the lower mantle above the D'' layer is nearly isotropic with a high degree of confidence. Therefore, we plotted the velocity difference between anisotropic pPv and isotropic Pv both for CaIrO$_3$ under ambient conditions and MgSiO$_3$ under mantle conditions (Fig. 6). From the plots for $V_P$ and slower $V_S$, we see that any velocity change over the D'' discontinuity is possible both for P and S waves depending on the lattice preferred orientation (LPO) characteristics of the pPv phase.

The seismic wave phases used in the D'' study (S, ScS, Sdiff) travel nearly horizontally in the D'' layer. Therefore, the two polarized S waves are classified as SH and SV, whose polarizations are in the horizontal and vertical directions, respectively. In seismological study, vertically transverse isotropy (VTI) and the tilted transverse isotropy have been assumed to explain S-wave polarization anisotropy in the D'' layer. According to seismological studies, the region of the Circum-Pacific rim is interpreted as having VTI with $V_{SH} > V_{SV}$, while the region of the Central Pacific has complicated VTI without any fixed relation between $V_{SH}$ and $V_{SV}$. Polarization anisotropy is not recognized beneath the Atlantic Ocean, or the region is considered as having VTI with $V_{SH} \approx V_{SV}$. On the other hand, from the polarization pattern in the $a$--$b$ plane of the faster $V_S$ plot (Figs 2b and 7), we can expect that $V_{SH} > V_{SV}$ in the D'' layer if the $c$ axis of pPv aligns vertically. Similarly, if the $b$ axis aligns vertically, the magnitude relation between $V_{SH}$ and $V_{SV}$ can be complicated.

Figure 4 | Velocity surface of Pv- and pPv-CaIrO$_3$. Stereo projections of elastic wave velocities for Pv- and pPv-CaIrO$_3$ based on the present results. The crystallographic directions, $a$, $b$, and, $c$, are identical in each plot as shown in (a). The unit of scale bars is m s$^{-1}$. (a–c) $V_P$, average of two $V_S$ values, and difference between two $V_S$ values, respectively, for Pv-CaIrO$_3$. (d–f) Same as a, b, and c but for pPv-CaIrO$_3$. 

| Average $V_S$ | Average $V_S$ | Difference between two $V_S$ | Difference between two $V_S$ |
|---------------|---------------|-----------------------------|-----------------------------|
| 2,850         | 2,900         | 2,950                       | 3,000                       |
| 3,050         | 3,100         | 3,150                       | 3,200                       |
| 3,300         | 3,400         | 3,500                       | 3,600                       |
| 5,600         | 5,800         | 6,000                       | 6,200                       |
| 6,400         | 6,600         | 5,400                       | 5,500                       |
| 5,600         | 5,800         | 6,000                       | 6,200                       |
| 6,400         | 6,600         | 5,400                       | 5,500                       |
We can summarize the above conclusion in an alternative way: For the c axis vertical case, horizontal wave propagation directions lie in the a–b plane, for which the in-plane shear mode (SH) has consistently higher wave speeds (Figs 2b and 7). If instead the b axis is vertical, horizontal wave propagation directions lie in the a–c plane, for which neither the in-plane (SV) nor the out-of-plane (SV) is consistently faster for all such propagation directions (Figs 2b and 7).

The dominant slip system of pPv-CaIrO₃ was determined as [100]⟨010⟩ from shear deformation experiments. Since the slip system of pPv-MgSiO₃ has not yet been determined, we assumed that it is similar to that of pPv-CaIrO₃. This assumption leads to LPO of pPv-MgSiO₃, in which the b axis aligns vertically along the lateral flow in the D'' layer. The LPO pattern is consistent with the complicated polarization S-wave anisotropy beneath the Central Pacific (Figs 2b and 7).

Although the texture development during Pv–pPv transition in MgSiO₃ has not yet been clarified, we expect the c axis alignment of pPv-MgSiO₃ immediately after the phase transition from Pv-MgSiO₃ from analogy with the case for MgGeO₃ (ref. 20). If the c axis aligns vertically, we expect VTI with V₃₈₉ > V₅⁶, which is consistent with the seismic feature beneath the Circum-Pacific rim.

Employing this conceptual scheme, we evaluated the feasibility of the LPO model of pPv in the D'' layer. The volume ratio between (Mg, Fe)SiO₃ Pv and (Mg, Fe)O ferropericlase (Fp) is 0.67:0.33 for the lower mantle. We also used this ratio for (Mg, Fe)SiO₃ pPv and Fp in the D'' layer. This means that the anisotropy resulting from pPv LPO should be weakened by approximately two-thirds in the D'' layer assuming isotropy of Fp. The observed polarization anisotropy of the S-wave velocity is ~3% anisotropy observed in the D'' layer is expected ~4.5% anisotropy to account ~3% anisotropy observed in the D'' layer. This requirement is easily satisfied by partial LPO of pPv (Table 2).

Figure 8 summarizes the present idea for seismic anisotropy and LPO structure in the D'' layer. The VTI of the vertical c axis and that of the vertical b axis can be interpreted as transformation LPO and deformation LPO, respectively. It is noted that isotropic pPv-MgSiO₃ is a reasonable interpretation for the seismic feature beneath the Pacific and Atlantic may be attributed to differences in temperature, chemical composition and strain rate. The proposed LPO behaviour for pPv-MgSiO₃ obtained from the

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**Figure 5 | Stereo projections of elastic wave velocities for Pv- and pPv-MgSiO₃.** The crystallographic directions, a, b, and, c, are identical in each plot as shown in (a). The unit of scale bars is m s⁻¹, although a factor of 10⁴ should be multiplied for Vₚ (a–c). Vₚ average of two Vₚ values, and difference between two Vₖ values, respectively, for Pv-MgSiO₃ under ambient conditions. (d–f) and (g–i) Same as a, b, and c but for Pv-MgSiO₃ and pPv-MgSiO₃ at 125 GPa and 2,500 K, respectively.
present mineral physics perspective is consistent with seismic observation.

To summarize, we presented a new data set of PV and Ppv-CaIrO3 single-crystal elasticity by means of IXS at SPring-8. We used the present results to interpret seismic wave anisotropy in the D00 layer, and proposed a model of Ppv LPO in the D00 layer consistent with seismic observations and mineral physics experiments on analogue materials. The present model may allow sophisticated discussion of global mantle convection, which is triggered by heating at the core–mantle boundary and modulated by lateral flow in the D00 layer.

Methods
Measurement of IXS at SPring-8 BL35XU. The small size and black colour of the present samples limit the techniques that can be used to investigate the elastic properties. In particular, the crystal sizes are too small to employ GHz ultrasonic methods22 or resonant ultrasound spectroscopy23, where both need crystals larger than 100 μm. In addition, the crystal opacity makes it difficult to apply Brillouin scattering. Therefore, IXS with an X-ray beam diameter of ~70 μm was employed.

Each specimen was attached to glass fibres (100 μm in diameter) using manicure resin and mounted on the four-circle goniometer installed at BL35XU of SPring-8 (ref. 24). The incident X-ray beam energy was 17.794 and 21.747 keV for Pbnm-CaIrO3 and Cmcm-CaIrO3 measurements, respectively. The IXS technique25,26 allows us to determine the elastic properties of a material by measuring the energies of acoustic longitudinal and shear phonon modes. When this information is combined with the precise reciprocal space...
position (momentum transfer) for each measurement, we can determine the sound velocity$^{27,28}$. If measurements are made in many different directions, the velocity information can be combined with the density, allowing all the elastic constants of a material to be determined. In the present work, the two-dimensional, $4 \times 3$, analyser array of BL35XU is especially advantageous as it allows access to 12 different directions of momentum transfer simultaneously.

The IXS is applicable on opaque materials; it is the most important advantage of IXS against Brillouin scattering. Further characteristics and procedures of the IXS measurements are summarized in Supplementary Figs 1–5.

Data analysis for elastic constants. A least-squares fit to Christoffel equation was then used to determine the elastic constants$^{29}$. $\text{Pv-CaIrO}_3$ and $\text{pPv-CaIrO}_3$ have orthorhombic symmetry and thus nine independent elastic constants. The six diagonal elastic constants ($C_{11}$, $C_{22}$, $C_{33}$, $C_{44}$, $C_{55}$, $C_{66}$) is related to nearly independent phonon modes propagating along crystallographic axes, while the three off-diagonal constants ($C_{12}$, $C_{23}$, $C_{31}$) can be determined from combination of a few measurements. We carefully planned the measurement to acquire sufficient data to constrain all nine elastic constants. Using the analyser array, we obtained $B_{40}$ and $B_{60}$ mode energies for $\text{Pv-CaIrO}_3$ and $\text{pPv-CaIrO}_3$, respectively.

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Table 2 | Characteristic velocities for crystals of post-perovskite structure.

|                | $V_P$, km s$^{-1}$ | $V_{SH}$, km s$^{-1}$ | $V_{SV}$, km s$^{-1}$ | $V_{SH}/V_{SV}$ | $\Delta V_s/V_s$ |
|----------------|-------------------|----------------------|----------------------|-----------------|-----------------|
| pPv-CaIrO$_3$  |                   |                      |                      |                 |                 |
| $a$–$b$ plane  | 6.1 (5)           | 3.4 (1)              | 2.8 (2)              | 1.19            | 0.11            |
| $b$–$c$ plane  | 6.1 (4)           | 3.1 (1)              | 2.9 (2)              | 1.06            | 0.07            |
| $c$–$a$ plane  | 6.3 (3)           | 3.4 (6)              | 3.1 (1)              | 1.08            | 0.17            |
| pPv-MgSiO$_3$  |                   |                      |                      |                 |                 |
| $a$–$b$ plane  | 14.0 (6)          | 7.6 (3)              | 7.1 (3)              | 1.07            | 0.09            |
| $b$–$c$ plane  | 13.9 (6)          | 7.2 (3)              | 7.4 (5)              | 0.98            | 0.07            |
| $c$–$a$ plane  | 14.2 (3)          | 7.4 (6)              | 7.8 (2)              | 0.94            | 0.09            |

$V_P$, $V_{SH}$ and $V_{SV}$ are the average velocities of the $P$ wave, $SH$ wave and $SV$ wave, respectively, on each plane specified on the left. $\Delta V_s$ is the maximum difference between the $SH$-wave and $SV$-wave velocities, and $V_s$ is the average of $V_{SH}$ and $V_{SV}$ on the corresponding plane. The source data for $\text{pPv-CaIrO}_3$ and $\text{pPv-MgSiO}_3$ are the results of the present IXS study and theoretical computation$^{25}$, respectively. The numbers in parentheses correspond to expected uncertainty.

Figure 8 | Schematic illustration of the LPO of $\text{pPv-MgSiO}_3$ in the D” layer. From a mineral physics viewpoint, the $\text{Pv}$ slab (purple region) transforms to $\text{pPv}$ (light blue region) and the $\text{pPv}$ slab then moves along the path of mantle convection with deformation (light-green region). In contrast, from seismic observation, the region of the Circum-Pacific rim has VTI anisotropy of $V_{SH} > V_{SV}$, while the region of the Central Pacific has complicated VTI with any magnitude relation; that is, $V_{SH} > V_{SV}$, $V_{SH} < V_{SV}$ or $V_{SH} \approx V_{SV}$. The region of the Atlantic Ocean (white circle) has VTI with $V_{SH} \approx V_{SV}$ suggesting isotropy or negligible LPO of $\text{pPv-MgSiO}_3$. The speculated aligned axes of $\text{pPv}$ LPO are shown by arrows for each region; the vertical $c$ axis beneath the Circum-Pacific rim and the vertical $b$ axis beneath the Central Pacific.
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Author contributions

A.Y. and H.F. planned and managed the present study equally. F.X. prepared sample of Pbnm-CaIrO₃. A.N. and A. Yoshiasa characterized single-crystal samples. Y.S. and K.O. joined the IXS measurement at SPring-8. S.T., H.U. and A.B. set up the instruments for the IXS measurement. A.Y. wrote the manuscript with input from H.F. and A.B.

Additional information

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