

High-pressure structural study of the scheelite tungstates CaWO_4 and SrWO_4 D. Errandonea,^{1,*} J. Pellicer-Porres,¹ F. J. Manjón,² A. Segura,¹ Ch. Ferrer-Roca,¹ R. S. Kumar,³ O. Tschauer,³ P. Rodríguez-Hernández,⁴ J. López-Solano,⁴ S. Radescu,⁴ A. Mujica,⁴ A. Muñoz,⁴ and G. Aquilanti⁵¹*Departamento de Física Aplicada-ICMUV, Universitat de València, Edificio de Investigación, c/Dr. Moliner 50, 46100 Burjassot (Valencia), Spain*²*Departamento de Física Aplicada, Universitat Politècnica de València, Cno. de Vera s/n, 46022 València, Spain*³*High Pressure Science and Engineering Center, Department of Physics, University of Nevada, 4505 Maryland Parkway, Las Vegas, Nevada 89154-4002, USA*⁴*Departamento de Física Fundamental II, Universidad de La Laguna, La Laguna 38205, Tenerife, Spain*⁵*European Synchrotron Radiation Facility, BP 220, Grenoble, F-38043 France*

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Angle-dispersive x-ray-diffraction and x-ray-absorption near-edge structure measurements have been performed on CaWO_4 and SrWO_4 up to pressures of approximately 20 GPa. Both materials display similar behavior in the range of pressures investigated in our experiments. As in the previously reported case of CaWO_4 , under hydrostatic conditions SrWO_4 undergoes a pressure-induced scheelite-to-fergusonite transition around 10 GPa. Our experimental results are compared to those found in the literature and are further supported by *ab initio* total-energy calculations, from which we also predict the instability at larger pressures of the fergusonite phases against an orthorhombic structure with space group *Cmca*. Finally, a linear relationship between the charge density in the AO_8 polyhedra of ABO_4 scheelite-related structures and their bulk modulus is discussed and used to predict the bulk modulus of other materials, like hafnion.

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I. INTRODUCTION

Scheelite ABX_4 compounds are important materials from both a theoretical and a technological point of view. Scheelite fluorides (ABF_4) like YLiF_4 and GdLiF_4 are used in rare-earth solid-state lasers,¹ scheelite oxides (ABO_4) like CaWO_4 and PbWO_4 are used as solid-state scintillators,^{2,3} and there is much interest in the use of scheelite compounds in optoelectronic devices.^{4–6} Moreover, a family of superhard materials has been found in ABO_4 compounds with A and B atoms having valence +4 (Ref. 7).

In the last years there has arisen renewed interest in ABX_4 compounds and their evolution under pressure. Many of these compounds crystallize in the scheelite structure (space group: $I4_1/a$, No. 88, $Z=4$) or in related structures like zircon (space group: $I4_1/amd$, No. 141, $Z=4$), pseudoscheelite (space group: $Pnma$, No. 62, $Z=4$), wolframite (space group: $P2_1/c$, No. 13, $Z=2$), *M*-fergusonite (space group: $I2/a$, No. 15, $Z=4$), hereafter called fergusonite, and *M'*-fergusonite (space group: $P2_1/c$, No. 14, $Z=2$). In particular, the ambient conditions scheelite structure of CaWO_4 and SrWO_4 has eight symmetry elements and a body-centered tetragonal primitive cell that includes two formula units; see Fig. 1(a). Each W site is surrounded by four equivalent O sites in tetrahedral symmetry about that site. Each Ca (Sr) cation shares corners with eight adjacent WO_4 tetrahedra.

Several experimental and theoretical works have been reported in the last decade on the pressure behavior of scheelite oxides and fluorides.^{8–33} Upon compression most of these compounds undergo structural transitions to monoclinic structures. However, several of these low-symmetry structures are difficult to characterize in high-pressure x-ray diffraction experiments and it has been further suggested that their formation could depend on the stress conditions in the

pressure chamber. In particular, a discussion regarding the high-pressure phase of CaWO_4 was open in recent years.^{8,9}

The occurrence of pressure-driven phase transitions in CaWO_4 and CaMoO_4 was first reported by Nicol and Durana,¹⁰ who postulated that the high-pressure phases had the wolframite structure. Other monoclinic structures that were considered during decades as candidate structures for the ABO_4 compounds at high pressure were those of α - MnMoO_4 type (space group: $C2/m$, No. 12, $Z=8$),¹¹ $\text{BaWO}_4(\text{II})$ type (space group: $P2_1/n$, No. 14, $Z=8$),¹² and HgWO_4 type (space group: $C2/c$, No. 15, $Z=4$).¹³ Errandonea and co-workers⁸ performed for the first time energy-dispersive x-ray powder diffraction (EDXRD) experiments on CaWO_4 up to pressures where the high-pressure phase was observed. They observed the occurrence of the pressure-driven phase transition at 10 GPa. These authors considered the four monoclinic structures previously postulated for the high-pressure phase of CaWO_4 to index their EDXRD patterns. Based on the quality of the unit-cell fit, they concluded that the high-pressure phase of CaWO_4 was most likely of the wolframite type⁸ [see Fig. 1(b)]. The same was also concluded by Shieh *et al.* from a high-pressure x-ray diffraction study of CdMoO_4 (Ref. 14). However, more recently Grzechnik *et al.*⁹ performed high-resolution angle-dispersive x-ray powder diffraction (ADXRD) on CaWO_4 and reported the high-pressure structure to be fergusonite like [see Fig. 1(c)]. Later measurements on BaWO_4 (Ref. 13), BaMoO_4 (Ref. 33), and CaMoO_4 (Ref. 29) also reported a scheelite-to-fergusonite phase transition, but in the case of SrWO_4 , a recent study combining x-ray diffraction and -absorption observed a phase transition at 11.7 GPa and characterized the high-pressure phase as wolframite.³² From the theoretical side, support to the scheelite-to-fergusonite transition with increasing pressure in ABX_4 scheelite compounds has been

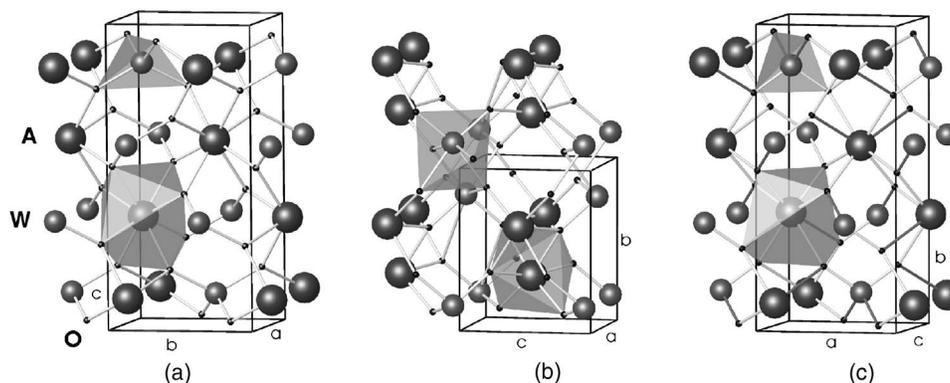


FIG. 1. The (a) scheelite, (b) wolframite, and (c) fergusonite structures of AWO_4 compounds. Large circles represent the A (Ca, Sr) atoms, middle-size circles correspond to the W atoms, and the small circles are the O atoms. The unit cell, A-O bonds, and W-O bonds are also shown. As a consequence of the scheelite-to-fergusonite transition two A-O and W-O bonds are enlarged (see text and Fig. 5); these bonds are shown as dark lines in (c). The AO_8 and WO_4 polyhedra are also shown. By comparing (a) and (c) it can be seen the polyhedra distortion caused by the scheelite-to-fergusonite transition.

given by the works of Sen *et al.*,^{15,16} while support to the scheelite-to-wolframite transition was reported in the work of Li *et al.*¹⁷

In this work we report new high-pressure ADXRD experiments up to nearly 18 GPa and x-ray-absorption near-edge structure (XANES) measurements up to nearly 20 GPa on $CaWO_4$ and $SrWO_4$ along with *ab initio* total-energy calculations in both compounds. From our ADXRD data we find that under hydrostatic conditions both compounds undergo a scheelite-to-fergusonite phase transition with increasing pressure, which is supported by the high-pressure XANES measurements and the *ab initio* total energy calculations.

II. EXPERIMENTAL DETAILS

$CaWO_4$ and $SrWO_4$ crystals were grown with the Czochralski method starting from raw powders having 99.999% purity.⁴ Samples were prepared as finely ground powders from the single crystals of $CaWO_4$ and $SrWO_4$. High-pressure ADXRD measurements were carried out in a 450- μm culet Merrill-Basset diamond-anvil cell (DAC) for $CaWO_4$ and in a 400- μm culet Mao-Bell DAC for $SrWO_4$. In the first case, powder samples were loaded together with a ruby chip into a 180- μm -diam hole drilled on a 200- μm -thick rhenium (Re) gasket preindented to 60 μm . In the second case, the Re gaskets were preindented to 40 μm and the diameter of the gasket hole was 100 μm . Silicone oil was used as pressure-transmitting medium in both cases. For XANES measurements under pressure, fine powder samples were loaded together with a ruby chip into a 200- μm -diam hole drilled on a 200- μm -thick Inconel gasket preindented to 50 μm and inserted between the diamonds of a 400- μm culet membrane-type DAC with silicone oil as pressure-transmitting medium. The pressure was measured by the shift of the $R1$ photoluminescence line of ruby.³⁴

ADXRD experiments were performed at the 16-IDB beamline of the HPCAT facility at the Advanced Photon Source (APS) using monochromatic radiation with $\lambda = 0.3679 \text{ \AA}$ [a Si (311) double-crystal monochromator was used]. The monochromatic x-ray beam was focused down to

$10 \times 10 \mu\text{m}^2$ using multilayer bimorph mirrors in a Kickpatrick-Baez configuration.³⁵ Diffraction images were recorded with a Mar345 image plate detector, 230 mm away from the sample, and were integrated and corrected for distortions using the FIT2D software.³⁶ The indexing, structure solution, and refinements were performed using the GSAS (Ref. 37) and POWDERCELL (Ref. 38) program packages.

XANES experiments were conducted at the ID24 energy-dispersive x-ray absorption station of the European Synchrotron Radiation Facility (ESRF).^{39,40} The key component of the dispersive setup is a curved monochromator that selects an energy span around the absorption edge and focuses the beam in the horizontal direction. All the energies contained in the diffracted beam are detected simultaneously by means of a position-sensitive detector. In order to establish the energy-pixel correlation, the spectrum of a reference standard is measured and compared with an equivalent spectrum acquired with the classical setup, where the knowledge of the Bragg angle allows for a determination of the energy. A more detailed description of the principles of energy-dispersive x-ray-absorption data collection is given in Ref. 41.

All XANES experiments were performed at the W L_3 edge (10.207 keV). At ID24, the combination of a profiled curved Si (111) monochromator⁴² and a vertically focusing mirror defined a focus spot of approximately $30 \times 20 \mu\text{m}^2$. The membrane DAC was situated at the focus position. The incident and transmitted beams were alternatively measured. In our experiments, the incident intensity was measured outside the pressure chamber. An essential experimental aspect of x-ray absorption spectroscopy (XAS) experiments in a DAC is the presence of diffraction peaks originating from diffraction from the diamond single crystals. The pressure cell is oriented with respect to the polychromatic x-ray beam so as to remove these glitches from the widest spectral range around the x-ray-absorption edge. This operation takes advantage of the real-time visualization of the XAS spectra, characteristic of the energy-dispersive setup. The presence of harmonics was avoided thanks to the grazing incidence mirrors situated between the undulator source and the monochromator. The reference standard for the energy calibration was metallic W.

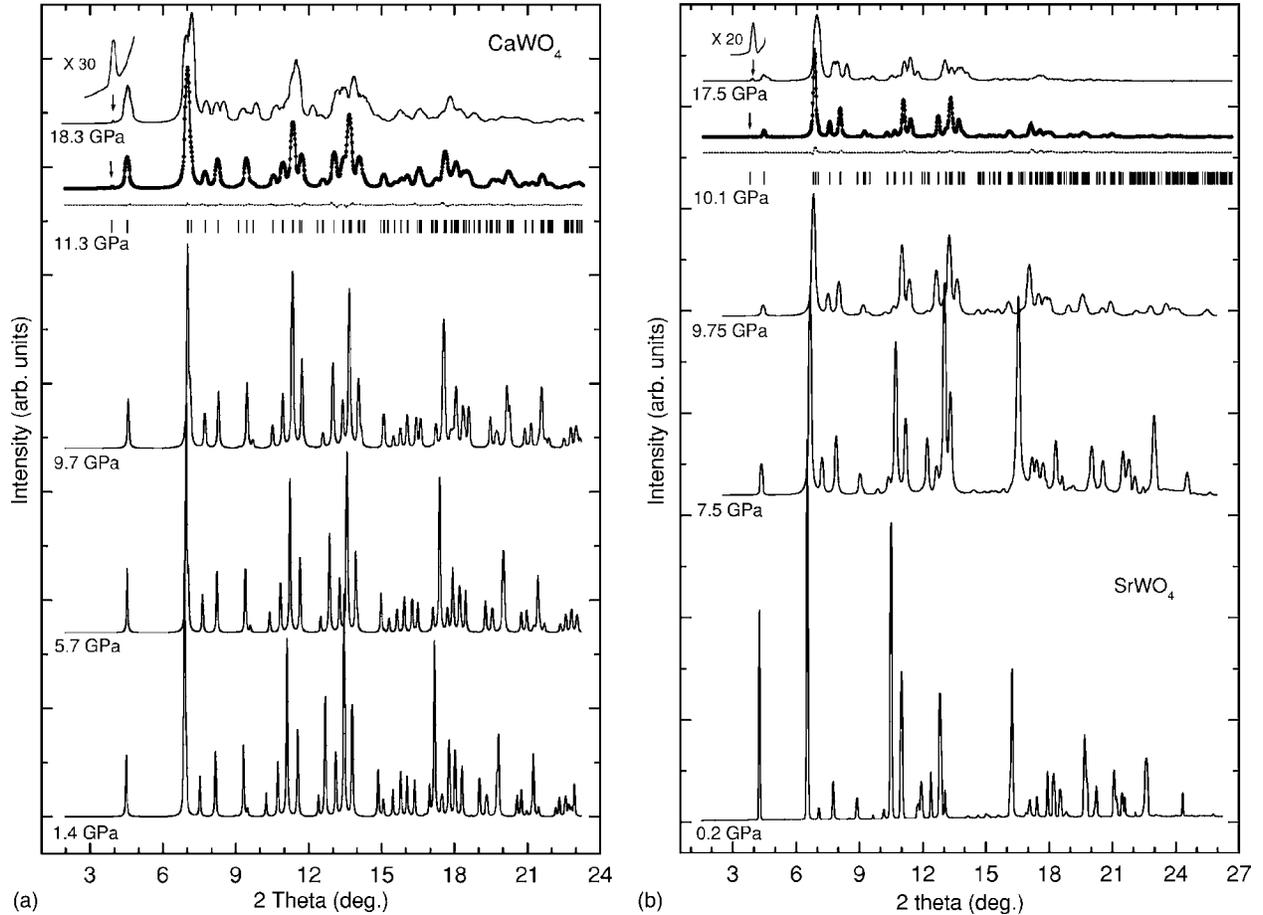


FIG. 2. Room-temperature ADXRD data of (a) CaWO_4 and (b) SrWO_4 at different pressures up to 18 GPa. In all diagrams the background was subtracted. To better illustrate the appearance of the (020) Bragg reflection of the fergusonite structure around $2\theta \approx 4^\circ$ a section of the upper trace is enlarged. In the ADXRD pattern of CaWO_4 collected at 11.3 GPa and of SrWO_4 at 10.1 GPa (which are representative of the high-pressure fergusonite structure) we also show the refined profile (symbols) and the difference between the measured data and the calculated profile (dotted line). The bars indicate the calculated positions of the reflections.

III. OVERVIEW OF THE CALCULATIONS

The structural stability of the phases of CaWO_4 and SrWO_4 was further investigated theoretically by means of total-energy calculations performed within the framework of the density functional theory (DFT) with the Vienna *ab initio* simulation package (VASP).⁴³ A review of DFT-based total-energy methods as applied to the theoretical study of phase stability can be found in Ref. 44. The exchange and correlation energy was evaluated within the generalized gradient approximation (GGA).⁴⁵ We used ultrasoft Vanderbilt-type pseudopotentials⁴⁶ and basis sets including plane waves up to a kinetic-energy cutoff of 850 eV for CaWO_4 and 495 eV for SrWO_4 . The tetrahedron method combined with Blöchl corrections was used for the Brillouin-zone integrations. The total energies were converged to below 1 meV per formula unit. The structural relaxation of the phases at each volume was conducted through the calculation of the forces on the atoms and the components of the stress tensor.

IV. RESULTS AND DISCUSSION

A. ADXRD measurements at high pressures

1. Low-pressure phase

Figure 2 shows our ADXRD data for CaWO_4 and SrWO_4 at several selected pressures up to 18 GPa. The evolution with pressure of the volume, lattice parameters, and axial ratios is plotted in Figs. 3 and 4 where we also compare our results with previously reported data for CaWO_4 (Refs. 8, 9, 19, 47, and 48) and SrWO_4 (Refs. 47 and 49) (in this case only for ambient pressure).

The pressure-volume (P - V) curves shown in Fig. 3(a) were analyzed in the standard way using a Birch-Murnaghan equation of state (EOS),⁵⁰

$$P = \frac{3}{2}B_0(x^{7/3} - x^{5/3}) \left[1 + \frac{3}{4}(B'_0 - 4)(x^{2/3} - 1) \right], \quad (1)$$

with $x = V_0/V$, where the parameters V_0 , B_0 , and B'_0 are the zero-pressure volume, bulk modulus, and pressure derivative of the bulk modulus, respectively. For scheelite CaWO_4 we

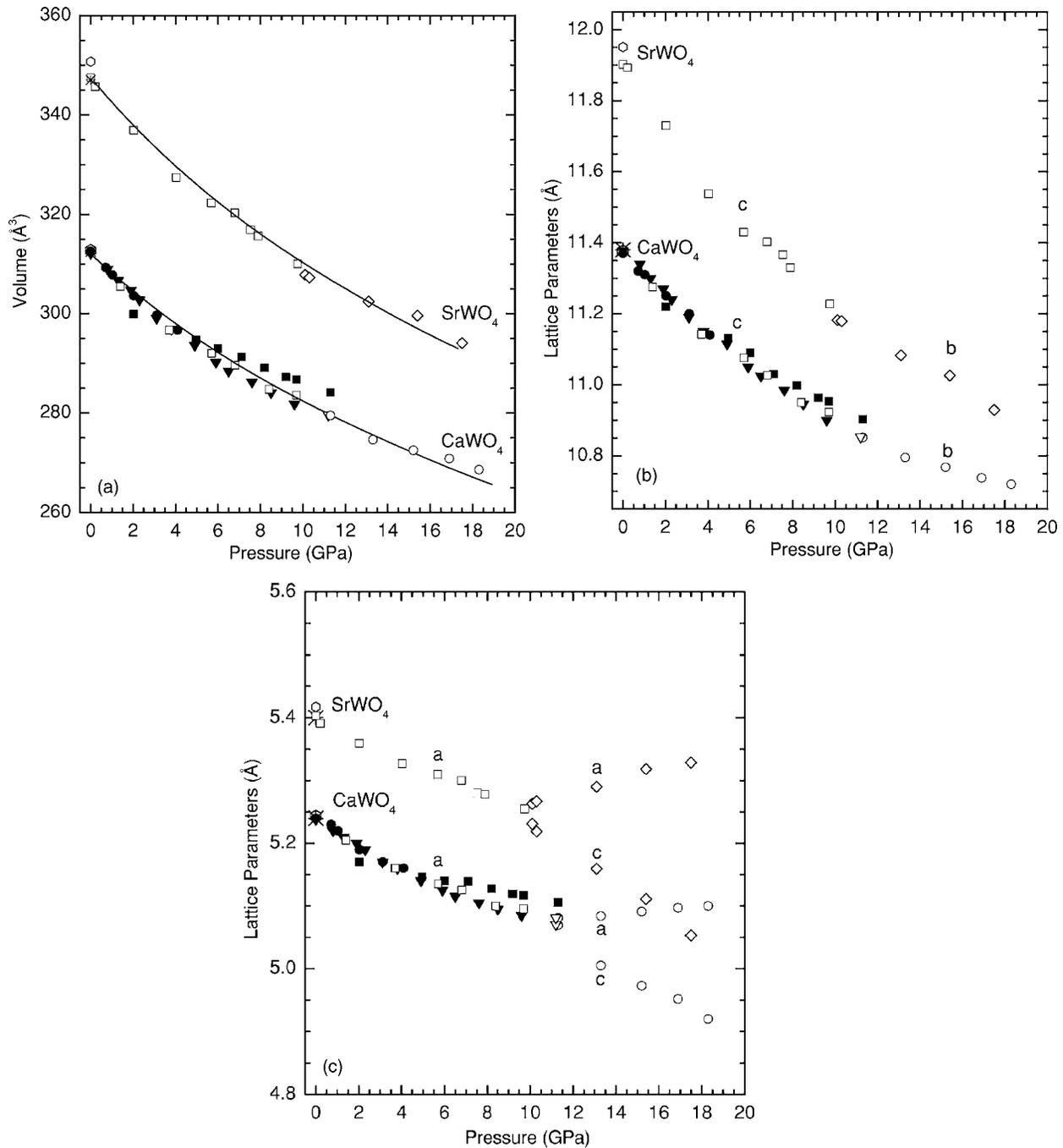


FIG. 3. Evolution of the (a) volume and (b),(c) lattice parameters of CaWO₄ and SrWO₄ with pressure. Open squares correspond to our data for the scheelite phase and open circles and diamonds to those for the fergusonite phase. Solid squares (Ref. 8), solid triangles (Ref. 9), solid circles (Ref. 19), stars (Ref. 47), and open hexagons (Refs. 48 and 49) are other data for the scheelite phase obtained from the literature. Open triangles are the fergusonite data reported in Ref. 9. In (a) the solid lines represent the EOS of the scheelite phase described in the text.

find $V_0=312(1) \text{ \AA}^3$, $B_0=74(7) \text{ GPa}$, and $B'_0=5.6(9)$ [$V_0=347.4(9) \text{ \AA}^3$, $B_0=63(7) \text{ GPa}$, and $B'_0=5.2(9)$ for scheelite SrWO₄]. These parameters are in good agreement with previous reported results^{9,19} and indicate that SrWO₄ is more compressible than CaWO₄, which is a direct consequence of the different compressibility of the *c* axis in the two compounds; see below. It is worth mentioning that the evolution of the volume of CaWO₄ with pressure reported in Ref. 8, and plotted as solid squares in Fig. 3(a) for the sake of com-

parison, underestimates the decrease of the volume above 7 GPa. This result gives support to the idea that a nonhydrostatic pressure environment may affect the structural pressure behavior of scheelite tungstates, as we will comment later on.

Figure 3(b) shows that the compressibility of the *c* axis of the scheelite structure is larger for SrWO₄ than for CaWO₄, while the *a* axis compresses in the same way in the two compounds [see Fig. 3(c)]. The larger compressibility of the

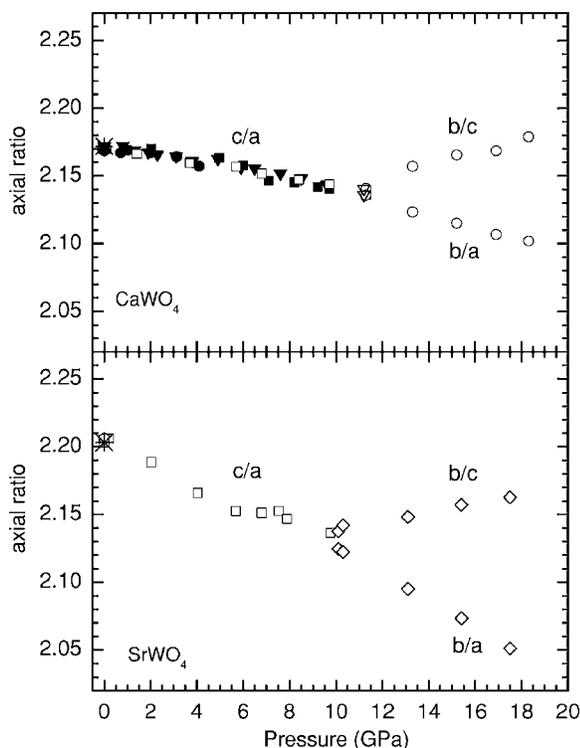


FIG. 4. Pressure dependence of the axial ratios of CaWO_4 and SrWO_4 . For a description of the symbols see Fig. 3.

c axis in SrWO_4 compared to that of CaWO_4 can be related to the difference in size of the Ca^{2+} and Sr^{2+} cations, which implies a larger charge density in the Ca environment with respect to that around Sr, as we will discuss later. The larger compressibility along the c axis as compared to that along the a axis is evident in Fig. 4.

We have also investigated the evolution of cation-anion distances in both compounds. According to the single-crystal high-pressure investigation carried out by Hazen *et al.*¹⁹ up to 4.1 GPa, the relative positions of the atoms in the CaWO_4 unit cell do not vary under pressure within the experimental error. In our experiment we have determined the internal parameters at the lowest pressure by means of a Rietveld refinement and then maintained them constant at higher pressures (see Table I). Figure 5 shows the evolution of the atomic distances between nearest neighbors with increasing pressure. The interatomic distances in CaWO_4 evolve in a similar way as previously reported,^{19,27} but the present results systematically differ by less than $\sim 2\%$ from those reported in Ref. 27. This difference was observed before by Hazen *et al.*¹⁹ between experiments performed inside and outside a DAC and can be attributed to the limited access to the reciprocal space of the used DAC (Ref. 19) and to the presence of impurities in the studied samples.⁵¹ The good agreement between our results and previous ambient pressure results^{19,52} suggests that the pressure evolution of the interatomic distances reported here is more reliable than previously published data. The decrease of Ca-O and Sr-O distances can be compared with the rigidity of the W-O bond distance in both compounds. In Fig. 5 it can be seen that there are two Ca-O and Sr-O distances, the largest distances

TABLE I. Structural parameters of the scheelite and fergusonite structure of CaWO_4 and SrWO_4 . These parameters were obtained from the present Rietveld refinements (see text).

(a) Structural parameters of scheelite CaWO_4 at 1.4 GPa: $I4_1/a$, $Z=4$, $a=5.205(5)$ Å, $c=11.275(7)$ Å

	Site	x	y	z
Ca	$4b$	0	0.25	0.625
W	$4a$	0	0.25	0.125
O	$16f$	0.2289(3)	0.0910(4)	0.0421(5)

(b) Structural parameters of fergusonite CaWO_4 at 11.3 GPa: $I2/a$, $Z=4$, $a=5.069(2)$ Å, $b=10.851(5)$ Å, $c=5.081(7)$ Å, $\beta=90.091(9)^\circ$

	Site	x	y	z
Ca	$4e$	0.25	0.6100(8)	0
W	$4e$	0.25	0.1325(3)	0
O_1	$8f$	0.9309(39)	0.9684(23)	0.2421(24)
O_2	$8f$	0.4850(35)	0.2193(31)	0.8637(37)

(c) Structural parameters of scheelite SrWO_4 at 0.2 GPa: $I4_1/a$, $Z=4$, $a=5.391(8)$ Å, $c=11.893(7)$ Å

	Site	x	y	z
Sr	$4b$	0	0.25	0.625
W	$4a$	0	0.25	0.125
O	$16f$	0.2497(9)	0.0925(9)	0.0421(6)

(d) Structural parameters of fergusonite SrWO_4 at 10.1 GPa: $I2/a$, $Z=4$, $a=5.263(9)$ Å, $b=11.182(6)$ Å, $c=5.231(6)$ Å, $\beta=90.35(1)^\circ$

	Site	x	y	z
Sr	$4e$	0.25	0.6027(9)	0
W	$4e$	0.25	0.1243(8)	0
O_1	$8f$	0.9309(49)	0.9598(53)	0.2619(42)
O_2	$8f$	0.4903(39)	0.2278(35)	0.8779(32)

being more compressible than the shorter ones.

Our results support the description of AWO_4 tungstate scheelites in terms of hard anionlike WO_4 tetrahedra surrounded by charge compensating cations. When pressure is applied the WO_4 units remain essentially undistorted and the reduction of the unit-cell size is mainly associated to the compression of the A-cation polyhedral environment.¹⁹ Along the a axis the WO_4 units are directly aligned, whereas along the c axis there is an A cation between two WO_4 tetrahedra. Therefore, the different arrangement of hard WO_4 tetrahedra along the c and a axes accounts for the different compressibility of the two cell axes. The different pressure behavior of the two A-O distances (Fig. 5) is associated with the different compressibility of the cell parameters. Effectively, the longest A-O distance has the largest projection along the c axis. It is important to point out that the asymmetric behavior of the c and a axes is also revealed in their different thermal expansion,⁵³ as well as in the evolution of the c/a ratio along a cationic A series.⁴⁷

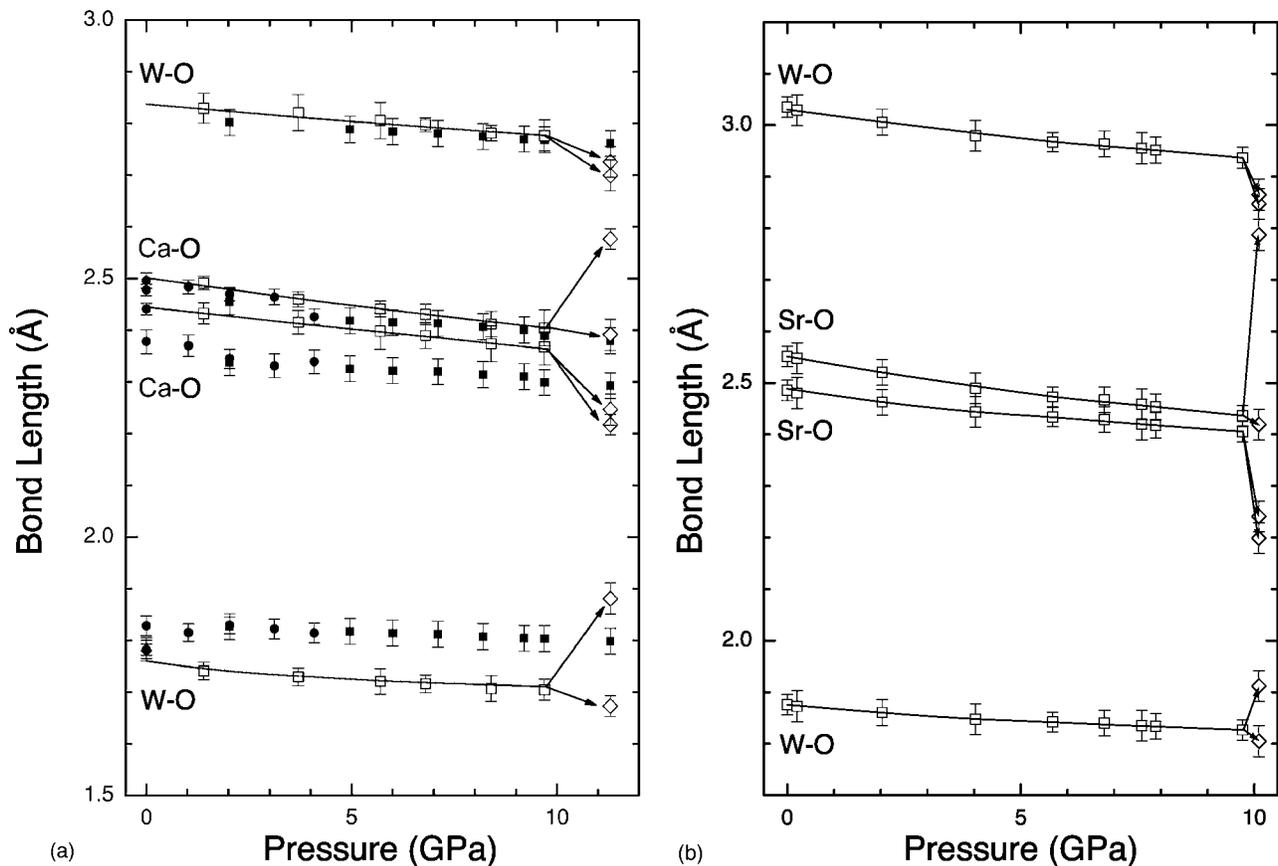


FIG. 5. Pressure dependence of the interatomic bond distances in the scheelite phase of (a) CaWO_4 and (b) SrWO_4 . Open squares represent the distances in the scheelite phase here reported. Solid circles (Ref. 19), solid squares (Ref. 27), and solid diamonds (Ref. 52) represent the distances in the scheelite phase reported in the literature. Open diamonds represent the new bond distances in the fergusonite phase after the phase transition.

2. High-pressure phases

The ADXRD spectra of CaWO_4 exhibit a change around 11.3 GPa, while in SrWO_4 the change occurs near 10.1 GPa (see Fig. 2). These changes are completely reversible upon pressure release. Below those pressures the observed diffraction peaks shift smoothly with compression and all the reflections observed in the diffraction patterns can be indexed within the scheelite structure whereas above those pressures some of the diffraction peaks split and additional diffraction peaks emerge. In particular, the appearance of a new peak around $2\theta \approx 3.8^\circ$ (depicted by an arrow in Fig. 2) is clearly distinguishable. The observed splitting of peaks and the appearance of new reflections suggests the occurrence of a second-order phase transition. The measured ADXRD patterns of the high-pressure phase can be indexed on the basis of the fergusonite structure but not on the basis of the wolframite structure, confirming the results of Grzechnik *et al.* for CaWO_4 (Ref. 9). The new Bragg peaks observed at $2\theta \approx 3.8^\circ$ in the high-pressure phase of both compounds correspond to the (020) reflection of the fergusonite structure of CaWO_4 and SrWO_4 . Two further facts support the assignment of the fergusonite structure to the high-pressure phase of both compounds and rule out the wolframite structure: The first one is that two of the stronger Bragg peaks of the wolframite structure—viz., the (011) and (110) expected at

$2\theta \approx 5.7^\circ$ —are absent in the measured diffraction patterns. The second one is that the (100) reflection of the wolframite structure is not present at $2\theta \approx 4.15^\circ$.

Figure 2 also shows the Rietveld refinements to the experimental spectra of CaWO_4 at 11.3 GPa and of SrWO_4 at 10.1 GPa obtained assuming the fergusonite structure. In order to perform the Rietveld refinement the starting Ca (Sr), W, and O positions were taken from Ref. 9. For both tungstates, we obtained good agreement with the experimental diffraction patterns. The residuals are $R_{\text{WP}}=1.75\%$, $R_{\text{p}}=1.1\%$, and $R(F^2)=1.5\%$ for CaWO_4 (197 reflections) and $R_{\text{WP}}=2.07\%$, $R_{\text{p}}=1.4\%$, and $R(F^2)=1.9\%$ for SrWO_4 (324 reflections). Similar refinement quality was obtained for scheelite CaWO_4 at 1.4 GPa and scheelite SrWO_4 at 0.2 GPa. Table I summarizes the lattice parameters and atomic positions of CaWO_4 at 1.4 and 11.3 GPa and of SrWO_4 at 0.2 and 10.1 GPa. Our structural parameters for fergusonite CaWO_4 agree with those reported by Grzechnik *et al.*⁹

It is worthwhile to discuss here the differences between the present results and those of Grzechnik *et al.*⁹ with previous structural studies of CaWO_4 and SrWO_4 . As we mentioned above, in a previous EDXRD study Errandonea *et al.*⁸ characterized the high-pressure phase of CaWO_4 as wolframite type. This conclusion was a result of an structural analysis performed using the extraction technique of LeBail *et al.*⁵⁴ and considering four candidate structures, among

which the fergusonite structure was not included. The exclusion of this structure was not accidental but a consequence of the fact that the (020) Bragg peak and other characteristic reflections of the fergusonite structure were not present in the EDXRD patterns of the high-pressure phase reported in Ref. 8. Furthermore, in these patterns there are also two reflections around 23 keV which were assigned to the (011) and (110) Bragg peaks of the wolframite structure and which cannot be indexed with the fergusonite structure—that is, the experimental situation was quite different from what we observe in the present experiments. We think that in the previous EDXRD experiments the presence of large nonhydrostatic stresses inside the DAC (Ref. 8) may have favored a transition to the wolframite structure instead of the fergusonite structure. In the study by Grzechnik *et al.*, both helium and a 4:1 methanol-ethanol mixture were used as pressure-transmitting medium.⁹ In the present study silicone oil was used as pressure-transmitting medium. In contrast, in Ref. 8 no pressure-transmitting medium was used. Using a nonhydrostatic pressure medium as NaCl, Nicol and Durana assigned the wolframite structure to the high-pressure phase of CaWO_4 (Ref. 10). The bulk modulus of CaWO_4 is 3 times larger than that of NaCl, and therefore the absence of a pressure-transmitting medium could create highly nonhydrostatic conditions at the onset of the transition.⁵⁵ It is well known that phase transitions can be greatly affected by nonhydrostatic conditions,⁵⁵ and therefore the fact that the less hydrostatic media was used in Ref. 8 could then have affected the characterization of the high-pressure phase of CaWO_4 . The observation of a scheelite-to-wolframite transition in CdMoO_4 in experiments performed by Shieh *et al.*¹⁴ using CdMoO_4 without pressure-transmitting medium, as well as the differences between the compressibility observed for the scheelite phase in these experiments and the one observed when a 4:1 methanol-ethanol mixture was used as pressure-transmitting medium,¹⁹ gives additional support to this hypothesis. Regarding SrWO_4 , Kuzmin *et al.*³² concluded recently from their x-ray-diffraction and -absorption measurements that the high-pressure phase of this compound is of the wolframite type. There are two principal facts that may explain the differences between the results reported in Ref. 32 and the present results. The first one is the lower quality of the EDXRD patterns reported in Ref. 32 in comparison with the ADXRD patterns reported here. The x-ray patterns reported in Ref. 32 do not allow the authors to perform a structural refinement, and all they can conclude is that there is a phase transition at 11.7 GPa, a pressure that is in fairly good agreement with our own results. The second one is that the extended x-ray-absorption fine-structure (EXAFS) measurements reported in Ref. 32 show that the local structure around the W atoms is compatible with an octahedral coordination at 30 GPa. However, from the EXAFS analysis alone, it is not possible to identify the structure of the high-pressure phase. Then, the possible existence of a post-fergusonite phase with the tungsten atoms in an octahedral coordination will resolve apparent controversies between our results and those reported by Kuzmin *et al.*³² Another fact to be taken into consideration is the possible metastability of two different monoclinic structures, a scenario that is supported by the polytypism observed in other tungstates (e.g., PbWO_4) even at ambient conditions.⁵⁶

Figures 3(b) and 3(c) show the lattice parameters of the fergusonite phases of CaWO_4 and SrWO_4 as a function of pressure up to ~ 18 GPa. Above 15 GPa the quality of the ADXRD patterns deteriorated, but it was still possible to obtain the lattice parameters at different pressures using the extraction technique of LeBail *et al.*⁵⁴ The degradation of the x-ray-diffraction patterns was observed previously in CaWO_4 (Ref. 9) and in similar compounds,^{13,57} and is independent of the pressure-transmitting medium employed in the experiments. This observation may be related to precursor effects either of a martensitic transition⁵⁸ or of the amorphization observed in alkaline-earth tungstates⁸ and other scheelite-structured compounds³¹ at higher pressures. The β angle was found to increase slightly from 90.09° at 11.3 GPa to 93° at 18.3 GPa in CaWO_4 and from 90.35° at 10.1 GPa to 92° at 17.5 GPa in SrWO_4 . The difference between the b/a and b/c axial ratios of the fergusonite phases of CaWO_4 and SrWO_4 also increases upon compression; see Fig. 4. These two facts imply an increase of the monoclinic distortion with pressure. A volume discontinuity is not apparent at the transition pressure, consistent with a second-order phase transition. The Birch-Murnaghan fit to both the scheelite and the fergusonite pressure-volume data gives EOS parameters (V_0 , B_0 , and B'_0) that differ by less than one standard deviation from those obtained for the scheelite data only. Hence, the EOS reported above can be assumed as a valid EOS for CaWO_4 and SrWO_4 up to 18 GPa, as illustrated in Fig. 3(a). A Birch-Murnaghan fit to only the high-pressure fergusonite data gives slightly larger values for B_0 and B'_0 [e.g., for CaWO_4 , we obtained $V_0=312(2) \text{ \AA}^3$, $B_0=78(9) \text{ GPa}$, and $B'_0=5.7(12)$ and, for SrWO_4 , $V_0=347(2) \text{ \AA}^3$, $B_0=64(8) \text{ GPa}$, and $B'_0=5.4(11)$]. A similar conclusion can be drawn from our *ab initio* calculations; see Sec. IV C.

In order to close the discussion on the ADXRD results we would like to comment that in both compounds the phase transition implies a distortion of the WO_4 tetrahedra accompanied by a small shear distortion of alternate (100) cation planes in the [001] direction. The scheelite-to-fergusonite transition occurs together with a slight decrease of two W-O bonds and the increase of the other two W-O bonds inside the WO_4 tetrahedra; however, as a consequence of this deformation, the volume of the WO_4 tetrahedra is enlarged by less than 10%. On the other hand, at the transition six of the A-O bonds in the AO_8 polyhedra are compressed and the remaining two are enormously expanded; see Fig. 5. The consequence of these changes is a decrease of the volume of the AO_8 polyhedra. In this way, as a result of the phase transition the WO_4 tetrahedra in the fergusonite phase are only slightly distorted, while the AO_8 polyhedra are quite distorted (see Fig. 1).

B. XANES measurements at high pressures

1. Low-pressure phase

The XANES part of the absorption spectrum is very sensitive to modifications in the neighborhood of the absorbing atom, and thus it can be used as a tool to detect structural changes. We have performed XANES experiments on CaWO_4 and SrWO_4 under compression with the aim of in-

TABLE II. Atomic positions used to perform the XANES simulations for the wolframite structure ($P2/c$, $Z=2$) (Ref. 60).

	Site	x	y	z
A	$2f$	0.5	0.3027	0.75
W	$2e$	0	0.1785	0.25
O ₁	$4g$	0.242	0.372	0.384
O ₂	$4g$	0.202	0.096	0.951

investigating changes in W coordination after the phase transition. In the scheelite structure the W environment is formed by four O atoms in tetrahedral configuration. If the high-pressure phase were fergusonite, the tetrahedron would become distorted which results in two slightly different near-neighbor distances but the main characteristics of the W environment would be maintained. In this situation we would expect small changes in the XANES spectra. If however the high-pressure phase would be of the wolframite type, the W coordination would change to 6 (2+4) and one would expect significant changes in the XANES spectra.

In order to confirm these ideas and as a guide to interpret changes in the experimental spectra, we have performed XANES simulations of the scheelite, fergusonite, and wolframite phases. The XANES simulations were carried out using the real-space multiple-scattering code implemented in the FEFF8 package.⁵⁹ We employed a self-consistent potential calculated using 120 atoms clusters (6.9 Å or 14 shells) and the Hedin-Lundqvist energy-dependent self-energy. Full multiple-scattering XANES calculations were performed using 87-atom clusters (6.5 Å or 11 shells). No pseudo-Debye-Waller factor has been considered in our simulations. The structural data used are given in Table I for the scheelite and fergusonite structures and in Table II for the wolframite structure. The description of wolframite is based on that of CdWO₄ (Ref. 60). For this structure, the lattice parameters have been scaled to give the same volume per formula unit as in the fergusonite structure. In Fig. 6 we present the results of the XANES spectra simulated in the three structures for CaWO₄ and SrWO₄. The spectra corresponding to both compounds are similar, with five resonances. The most dramatic change observed when passing from fourfold coordination to sixfold coordination affects the resonance denoted *B* in Fig. 6. In the scheelite and fergusonite structures the *B* resonance is clearly observable, but it disappears in the wolframite simulation. Other noticeable changes concern the intensity and width of the white line (*A* resonance).

Figure 7 shows the experimental XANES spectra at different pressures for CaWO₄ and SrWO₄. The spectra of both compounds at atmospheric pressure show the five resonances predicted by our simulations for the scheelite structure. The position and intensity of each feature agree qualitatively with those of the simulation, except for the resonances *D* and *E* in CaWO₄ whose relative intensities are inverted. In the theoretical spectra the resonances are more pronounced as a consequence of not considering the pseudo-Debye-Waller factor.

2. High-pressure phases

The high-pressure XANES spectra of CaWO₄ show no significant changes up to 11.3 GPa; see Fig. 7(a). At this

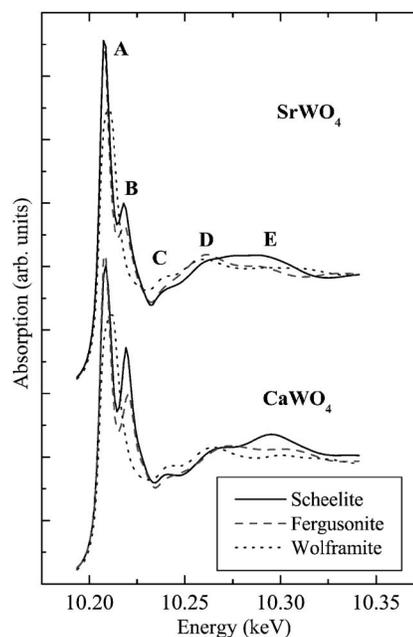


FIG. 6. *Ab initio* simulation of XANES spectra of CaWO₄ and SrWO₄ in the three phases of scheelite, fergusonite, and wolframite. The main difference between the fergusonite and the wolframite phases affects the intensity of the *B* resonance and the intensity and width of the white line (labeled *A* in the figure). There are also minor intensity changes in the *C*, *D*, and *E* resonances.

pressure the *B* resonance loses intensity and the ratio of intensities between the *D* and *E* resonances also decreases. Meanwhile, the intensity and width of the white line remain unaffected. The changes described indicate a transition to the fergusonite phase at 11.3(10) GPa, in agreement with ADXRD results. It is interesting to note that XANES spectra continue to evolve up to the maximum pressure attained of 20.2 GPa, suggesting, as we observed in our ADXRD measurements, that the structural distortions leading to the fergusonite structure become more pronounced when applying pressure. The phase transition is reversible, as the spectrum of the recovered phase is identical to the initial one except for a diminution in the white line intensity which we interpret as due to a decrease in sample thickness.

As regards to SrWO₄ the XANES spectra up to 12.4 GPa show only a small reduction of the intensity of the *B* resonance; see Fig. 7(b). At 15.0 GPa an acceleration in the decrease of the *B* resonance is accompanied by the progressive disappearance of the *C* resonance and an increase of the *D* resonance, while the white line remains unchanged. These changes continue up to the maximum pressure attained of 22.2 GPa. At this pressure the *B* resonance is still visible in the spectrum. Once again, the evolution of the spectra is reversible and suggests a transition towards the fergusonite phase. However, the onset of the phase transition is not as clear as in CaWO₄ and the distortion of the W tetrahedral environment is not evident up to 13.7(17) GPa.

C. *Ab initio* calculations

We compare now the experimental body of data presented in the previous sections with the results from our total-

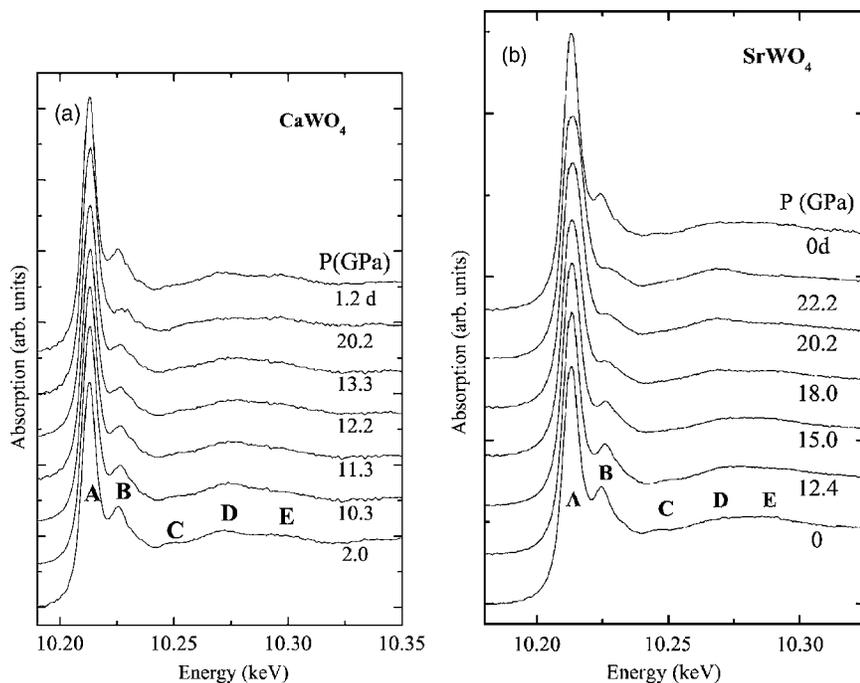


FIG. 7. Experimental XANES spectra of (a) CaWO_4 and (b) SrWO_4 measured at different pressures. The spectra collected on pressure release are marked with *d*. The analysis of the spectra reveals a transition to the fergusonite phase in both compounds. At the transition we observed intensity changes in the resonances. In CaWO_4 , *B* decreases by 8% and the ratio between the intensities of *D* and *E* decreases by 7%.

energy theoretical study of several structural phases of CaWO_4 and SrWO_4 . Along with the observed scheelite and fergusonite phases we have also considered the wolframite structure previously proposed for the high-pressure phase of CaWO_4 (Ref. 8) as well as other candidate structures on account of their observation or postulation in previous high-pressure work for related compounds: M' -fergusonite,¹⁶ LaTaO_4 ,⁶¹ $\text{BaWO}_4\text{-II}$,¹² and $\text{YLiF}_4\text{-Sen}$ (as we call the very-high-pressure structure found in the molecular dynamics study reported by Sen *et al.*¹⁶). Several of these phases are structurally related and can be represented within the monoclinic space group $P2_1/c$ (No. 14), which has thus also received our special attention.

Figure 8 shows the energy-volume curves for the different structures of CaWO_4 and SrWO_4 , from which the relative stability and coexistence pressures of the phases can be extracted by the common-tangent construction.⁴⁴ At all the pressures investigated and for both compounds the M' -fergusonite structure reduced upon full relaxation to fergusonite—it is thus not shown in Fig. 8. This figure shows the scheelite phases as being stable at zero and low pressure, with $V_0=318.3 \text{ \AA}^3$, $B_0=72 \text{ GPa}$, and $B'_0=4.3$ for CaWO_4 and $V_0=362.2 \text{ \AA}^3$, $B_0=62 \text{ GPa}$, and $B'_0=4.9$ for SrWO_4 . These values compare well with the experimental results, with differences within the typical reported systematic errors in DFT-GGA calculations. A similar degree of agreement exists for the calculated values of the internal parameters of the scheelite phases [O(16*f*) at (0.244, 0.097, 0.039) and $c/a=2.16$ for CaWO_4 ; O(16*f*) at (0.237, 0.111, 0.042) and $c/a=2.20$ for SrWO_4 ; cf. Table I].

As pressure increases, the scheelite structure becomes unstable against fergusonite. The fergusonite structure, a distortion of scheelite, only emerges as a structurally different and thermodynamically stable phase above a compression threshold of about 10–11 GPa in both compounds; at the lower pressures investigated, the relaxation of the fergusonite struc-

ture resulted in the scheelite structure. This is consistent with a continuous or quasicontinuous scheelite-to-fergusonite transition with none or very little volume collapse. The calculated structural parameters of the fergusonite phases are also in good agreement with the experimental results [$y(\text{Ca})=0.624$, $y(\text{W})=0.132$, $\text{O}_1(8f)$ at (0.912, 0.963, 0.242), $\text{O}_2(8f)$ at (0.492, 0.217, 0.822), $b/a=2.104$, $c/a=0.977$, and $\beta=91.6^\circ$ for CaWO_4 at 11 GPa; $y(\text{Sr})=0.624$, $y(\text{W})=0.128$, $\text{O}_1(8f)$ at (0.905, 0.961, 0.235), $\text{O}_2(8f)$ at (0.485, 0.213, 0.840), $b/a=2.145$, $c/a=0.990$, and $\beta=90.3^\circ$ for SrWO_4 at 11 GPa; cf. Table I].

The $\text{BaWO}_4\text{-II}$ and $\text{YLiF}_4\text{-Sen}$ structures are very high in enthalpy and nowhere close to stability in either compound. The LaTaO_4 -type structure is similarly high in enthalpy in CaWO_4 though in SrWO_4 it is placed considerably lower and is in fact a competitive candidate for stability in a post-fergusonite regime around 20 GPa. The wolframite structure is not thermodynamically stable in any interval of pressures though it is close in energy (20–40 meV) to fergusonite in CaWO_4 in the relevant range around 10–20 GPa which might have a bearing on its observation in previous experimental work in which nonhydrostatic conditions were used.⁸

A difficulty found in the relaxation of the monoclinic phases belonging to space group $P2_1/c$ is the existence of a number of local minima. For a significant interval of medium and high pressures these structurally different minima are located very close in energy, sometimes separated by shallow barriers, which make the precise determination of the absolute minimum within this set of low-symmetry crystal structures a rather tedious and difficult task. Nevertheless, we have carried out such minimization, ensuring great care in the relaxation procedure, which requires in particular repeating relaxation starting from different initial conditions and checking for local stability. In the course of this minimum-trapping quest we have arrived at a well-defined minimum in the compressed region for a structure which after refinement

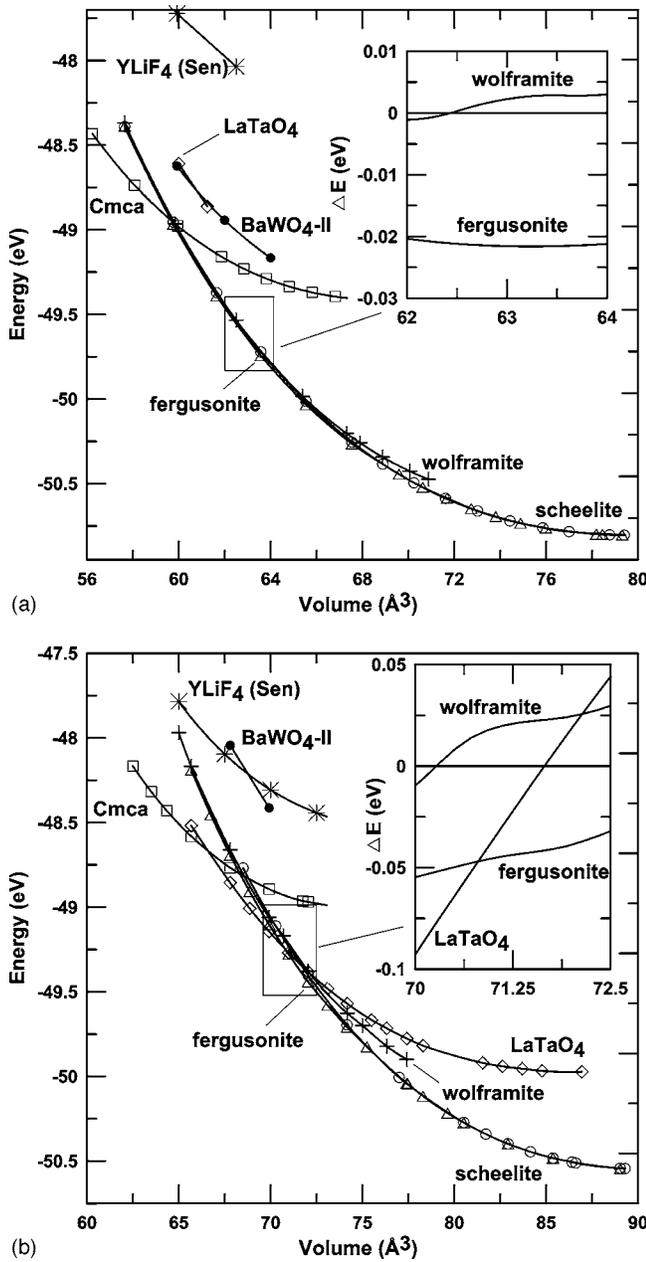


FIG. 8. Energy-volume curves (both per formula unit) calculated for (a) CaWO_4 and (b) SrWO_4 . The structures shown are scheelite (circles), fergusonite (triangles), wolframite (crosses), LaTaO_4 (diamonds), Cmca (squares), $\text{BaWO}_4\text{-II}$ (dots), and $\text{YLiF}_4\text{-Sen}$ (stars). The insets show differences in energy with respect to the scheelite phase in the marked areas.

and further analysis turned out to have *increased* orthorhombic symmetry, with space group Cmca (No. 64). This totally unexpected structural phase⁶² has lower enthalpy than any other of the phases considered above ~ 29 GPa in CaWO_4 and ~ 21 GPa in SrWO_4 [in this case in close competition with the LaTaO_4 -type structure—see Fig. 8(b)]. It has $Z=8$ with Ca atoms in $8e$ positions at (0.25, 0.164, 0.25), W at $8f$ (0, 0.409, 0.226), and O at $8d$ (0.157, 0.5, 0), $8e$ (0.25, 0.348, 0.25), $8f$ (0, 0.288, 0.005), and $8f$ (0, 0.084, 0.094) for CaWO_4 at 30 GPa [for SrWO_4 at 23 GPa: Sr($8e$) (0.25, 0.167, 0.25), W($8f$) (0, 0.413, 0.223), $\text{O}_1(8d)$ (0.149, 0.5, 0),

$\text{O}_2(8e)$ (0.25, 0.359, 0.25), $\text{O}_3(8f)$ (0, 0.292, 0.034), and $\text{O}_4(8f)$ (0, 0.084, 0.077)]. In both materials $b/a \sim 1.65\text{--}1.68$ and $c/a \sim 0.68$. In this structure the Ca (Sr) and W cations are surrounded by ten and six O atoms, respectively. It is worth noting that this new structure is *strongly* energetically favored over fergusonite in the high-pressure regime and thus the figures of ~ 29 GPa in CaWO_4 and ~ 21 GPa in SrWO_4 constitute neat upper bounds for the thermodynamical stability of the respective fergusonite phases. Such high pressures are just above those reached in x-ray-diffraction experiments.

D. Bulk modulus in scheelite ABO_4 compounds

Hazen *et al.* found that the bulk modulus of certain binary oxides and silicates can be directly correlated to the compressibility of the A-cation coordination polyhedra.⁶³ In particular, they proposed that the bulk compressibility in these compounds is proportional to the average volume of the cation polyhedron divided by the cation formal charge; i.e., B_0 is proportional to the cation charge density per unit volume inside the cation polyhedron. They also found that $\text{A}^{2+}\text{B}^{6+}\text{O}_4$ scheelite tungstates and molybdates under pressure compressed in an anisotropic way with the WO_4 and MoO_4 tetrahedra behaving as rigid units.¹⁹ Furthermore, they ordered the compressibility of scheelite compounds according to the A-cation formal charge and, on this basis, suggested that the compressibility of ABO_4 scheelites could be given by the compressibility of the softer AO_8 polyhedron and that the $\text{A}^{4+}\text{B}^{4+}\text{O}_4$ scheelites could be a family of ultrahard materials.

These last conclusions have been confirmed in two recent works, where the bulk moduli of scheelites have been plotted as a function of the bulk volume.^{7,64} Further insight can be obtained with the present data by plotting the bulk modulus of scheelite and scheelite-related compounds as a function of the A-cation charge density per unit volume in the AO_8 polyhedra, given by the A-cation formal charge divided by the cubic average A-O distance (see Fig. 9). All data plotted in Fig. 9, summarized in Table III, correspond to approximately 25% of the ABO_4 compounds with the scheelite and scheelite-related structures that can be found in the Inorganic Crystal Structure Database. The bulk modulus of all the plotted compounds obeys a linear relationship according to the equation

$$B_0 = 610(110) \frac{Z_A}{d_{A-O}^3}, \quad (2)$$

where B_0 is the bulk modulus (in GPa), Z_A is the A-cation formal charge (being $4 \geq Z_A \geq 1$), and d_{A-O} is the average A-O distance (in \AA) inside the AO_8 polyhedron. This simple rule serves as an effective and simple empirical criterion for predicting the bulk modulus of any scheelite or scheelite-related ABO_4 compound. The linear relationship between B_0 and the A-cation charge density of the AO_8 polyhedra is consistent with the fact that AO_8 polyhedra exhibiting a large A-cation charge density result in a larger electronic cloud inside the polyhedra than those AO_8 polyhedra with a low A-cation charge density. In the AO_8 polyhedra with a high Z_A the electrons around the cation are highly localized and the

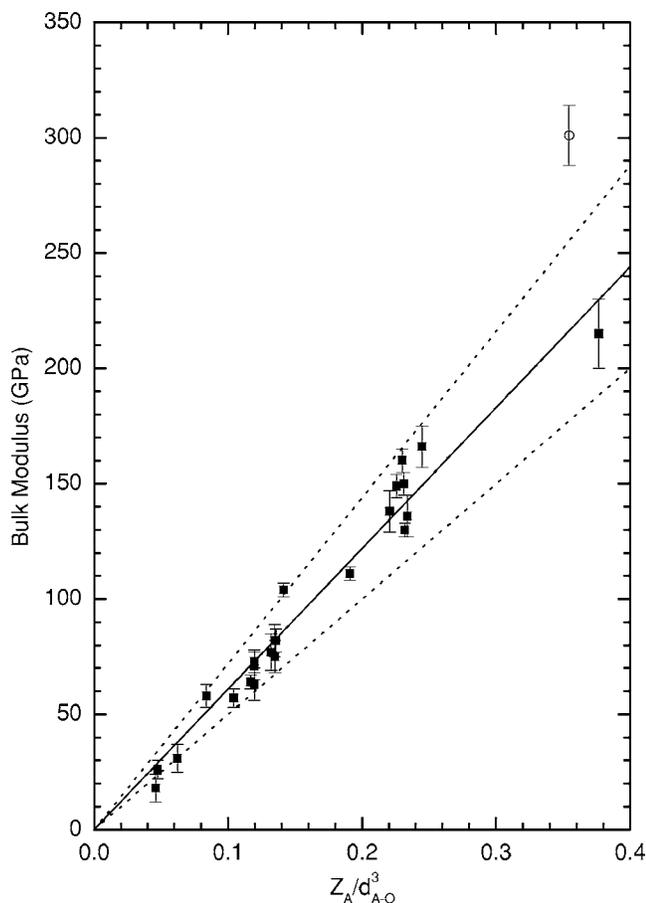


FIG. 9. Values of the ambient-pressure bulk modulus of ABO_4 scheelite and scheelite-related compounds plotted against the value of the cation charge density of the AO_8 polyhedra. $A-O$ distances and ambient-pressure bulk moduli were taken from different references (Refs. 7–9, 13, 16, 19, 27, 51, and 65–80) and are summarized in Table III. The white circle represents the bulk modulus reported by Scott *et al.* (Ref. 7) for scheelite $ZrSiO_4$. The solid line corresponds to the relation given in Eq. (2), and the dashed lines indicate its lower and higher deviations.

bond distances cannot be highly deformed under pressure. On the contrary, in AO_8 polyhedra with a low Z_A the density electrons around the cation are highly delocalized and the bond distances can be considerably deformed under pressure. Then, since the compressibility of ABO_4 compounds is mainly given by the compression of the AO_8 polyhedra, the above-described facts explain why B_0 is proportional to Z_A . In addition to that, they also explain why AO_8 polyhedra with A valence (+1, +2, and +3) are highly deformed as compared to BO_4 polyhedra with B valence (+7, +6, and +5) in ABO_4 scheelites and scheelite-related structures, being the compounds with A - and B -cation valence equal to 4+ the hardest ABO_4 materials. In fact, the linear relationship stated above should not be applicable to $A^{4+}B^{4+}O_4$ scheelites if AO_8 and BO_4 tetrahedra have similar compressibilities. However, despite both A and B cations having equal valence, $B-O$ bonds in tetrahedral configuration are shorter and stronger than $A-O$ bonds and the bulk modulus is again dictated by the AO_8 polyhedra. Therefore, Eq. (2) can also be effectively applied to $A^{4+}B^{4+}O_4$ scheelites as clearly shown.

It has been recently reported that both the scheelite and the zircon structure of YVO_4 have a quite similar bulk modulus.⁶⁴ This result is in agreement with our expectations since in both structures the $Y-O$ bond distances differ by less than 2%. A similar behavior should be expected also for $ZrSiO_4$, with similar $Zr-O$ bond distances in the scheelite and zircon structures (see Table III). However, a bulk modulus of 300 GPa has been recently reported for the scheelite phase of $ZrSiO_4$ (Ref. 7). This bulk modulus exceeds by more than 30% the bulk moduli of the zircon structure of $ZrSiO_4$. Therefore, according to the systematic here reported, a bulk modulus of 300 GPa for the scheelite phase of $ZrSiO_4$ is unrealistic and we think that the extremely low compressibility recently reported for this material could be mistaken. Following Eq. (2), we can predict for the scheelite phase of $ZrSiO_4$ a bulk modulus of 220(40) GPa, which is one of the largest bulk moduli of ABO_4 compounds. Theoretical calculations using either the local-density approximation (LDA) or the GGA gave a bulk modulus of 230(25) GPa (Ref. 81), value that agrees well with our estimation. A bulk modulus of 300 GPa can be only expected for a compound with octahedral coordinated silicon atoms, like $\gamma-Si_3N_4$, but not for compounds with tetrahedral coordinated Si atoms,⁸² like scheelite $ZrSiO_4$. We attributed the overestimation of B_0 to (i) the nonhydrostatic conditions of the experiments performed by Scott *et al.*,⁷ who used a 16:3:1 methanol-ethanol-water mixture as pressure-transmitting medium up to 52.5 GPa, and (ii) to the large presence of impurities in the natural zircon samples used by Scott *et al.*, as suggested by van Westrenen *et al.*⁵¹ The first argument leads to large pressure gradients and an inaccurate estimation of the pressure inside the DAC when a 60- μm x-ray beam is used because the pressure-transmitting medium used is not hydrostatic above 15 GPa. In fact, the Pt diffraction peaks used for determining the pressure in Ref. 7 are quite broad. These facts may easily cause an overestimation of the bulk modulus of the scheelite phase of $ZrSiO_4$. The second argument has proved to lead to different transition pressures and different pressure coefficients. New experiments using a microfocus x-ray beam and better hydrostatic conditions are needed to check the pressure behavior of the scheelite phase of $ZrSiO_4$.

To conclude, we would like to mention that attempting to predict the pressure behavior of other scheelite structures and zircon-structured ABO_4 materials we used Eq. (2) to make a back-of-the-envelope estimation of the bulk modulus of several compounds, which have been selected by considering their actual technological interest. Our predictions are summarized in Table IV. In the case of $BaMoO_4$, our estimation of B_0 is in quite good agreement with the recent experimental results of Panchal *et al.*³³ On top of that, according to our estimations, hafnon ($HfSiO_4$) is expected to be one of the least compressible ABO_4 compounds, being therefore a material of interest for potential applications as an interphase component in toughened oxide ceramic composites.⁸³ Our predictions for $NaReO_4$ can be compared with the bulk modulus obtained from DFT calculations by Spitaler *et al.*⁸⁴ These authors reported $B_0=18.3$ GPa. This value is approximately half of the value estimated by us. However, a Birch-Murnaghan fit to the results reported by Spitaler *et al.* gives a negative value for the pressure derivative of B_0 , something

TABLE III. Summary of the data plotted in Fig. 9. The structure, A-O bond distance, cation formal charge, and bulk modulus are given.

ABO_4 compound	Space group	Mean A-O bond distance [Å]	Cation formal charge	B_0 [GPa]	Reference
ZrSiO ₄	$I4_1/a$	2.243	4	301(13)	7
ZrSiO ₄	$I4_1/amd$	2.198	4	215(15)	51, 65, 66
LaNbO ₄	$I4_1/a$	2.505	3	111(3)	67
YVO ₄	$I4_1/a$	2.387	3	138(9)	64
TbVO ₄	$I4_1/amd$	2.369	3	149(5)	68
BiVO ₄	$I4_1/a$	2.350	3	150(5)	69
DyVO ₄	$I4_1/amd$	2.354	3	160(5)	70
YVO ₄	$I4_1/amd$	2.348	3	130(3)	64
ErVO ₄	$I4_1/amd$	2.341	3	136(9)	71
LuPO ₄	$I4_1/amd$	2.306	3	166(9)	72
BaSO ₄	$Pnma$	2.879	2	58(5)	73, 74
BaWO ₄	$I4_1/a$	2.678	2	57(4)	13, 75
PbWO ₄	$I4_1/a$	2.579	2	64(2)	19
PbMoO ₄	$I4_1/a$	2.576	2	64(2)	19
SrWO ₄	$I4_1/a$	2.557	2	63(7)	This work
EuWO ₄	$I4_1/a$	2.557	2	65(6)	75
SrMoO ₄	$I4_1/a$	2.556	2	73(5)	76
NaY(WO ₄) ₂	$I4_1/a$	2.478	2	77(8)	77
CaMoO ₄	$I4_1/a$	2.458	2	82(7)	19, 29
CaWO ₄	$I4_1/a$	2.457	2	75(7)	This work, 8, 9, 19, 74
SrSO ₄	$Pnma$	2.452	2	82(5)	16
CdMoO ₄	$I4_1/a$	2.419	2	104(2)	19
KReO ₄	$I4_1/a$	2.791	1	18(6)	78
TlReO ₄	$Pnma$	2.765	1	26(4)	79
AgReO ₄	$I4_1/a$	2.524	1	31(6)	80

unexpected for a scheelite ABO_4 compound, which suggests the EOS of NaReO₄ may be miscalculated in Ref. 84. This conclusion is also supported by the fact that the value predicted by us for B_0 is very similar to that experimentally observed in other perrhenates (see Table III), as expected.

TABLE IV. Predicted bulk modulus for different scheelite-type and zircon-type compounds.

ABO_4 compound	Space group	Mean A-O bond distance [Å]	Cation formal charge	B_0 [GPa]
HfSiO ₄	$I4_1/amd$	2.186	4	235(40)
YPO ₄	$I4_1/amd$	2.337	3	145(25)
YAsO ₄	$I4_1/amd$	2.383	3	135(25)
EuCrO ₄	$I4_1/amd$	2.410	2	87(15)
ZrGeO ₄	$I4_1/a$	2.203	4	230(40)
BaMoO ₄	$I4_1/a$	2.741	2	59(12)
NaReO ₄	$I4_1/a$	2.446	1	42(8)
KIO ₄	$I4_1/a$	2.816	1	27(5)

V. CONCLUSIONS

We have measured ADXRD and XANES spectra in CaWO₄ and SrWO₄ under pressure up to ~20 GPa. In both cases our results support the existence of a reversible scheelite-to-fergusonite structural transition under hydrostatic conditions. From our ADXRD data we locate the onset of the transition at 10.8(5) GPa in CaWO₄ and at 9.9(2) GPa in SrWO₄. The monoclinic distortion triggered at the phase transition continues up to the maximum pressures attained in our experiment, with no evidence of any further structural transformation. The small changes of the local environment around the absorbing atom make XANES sensitive to the phase transition at slightly higher pressures, around 11.3(10) GPa in CaWO₄ and 13.7(17) GPa in SrWO₄. In the case of SrWO₄ precursor effects of the transition appear at 10 GPa but the transition is not completed up to 15 GPa. The sluggish character of the transition is confirmed not only by the present ADXRD and XANES experiments, but also by the Raman investigation carried out in Ref. 30, where the pressure dependence of some modes related to the internal movement in the WO₄ tetrahedra is found to be strongly nonlinear

up to 3–4 GPa above the transition pressure. Our *ab initio* theoretical study of the energetic of the phases supports the scheelite-to-fergusonite transition and yields structural characteristics for the scheelite and fergusonite phases in very good agreement with the experimental results. In addition, from our *ab initio* study we can place an upper bound (not reached experimentally) on the stability of the fergusonite high-pressure phases, at ~ 29 GPa in CaWO_4 and ~ 21 GPa in SrWO_4 , which calls for experimental structural studies in this higher-pressure region. Finally, we have shown that the ambient-pressure bulk modulus of ABO_4 scheelite and scheelite-related compounds can be easily estimated if the average A-O distance is known.

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