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Structural stabilities, robust half-metallicity, magnetic anisotropy, and thermoelectric performance of the pristine/Ir-doped $\text{Sr}_2\text{CaOsO}_6$: strain modulations†

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Half-metallic (HM) ferromagnetic (FM)/ferrimagnetic (FIM) materials with a large energy-gap (E_g) and high magnetocrystalline anisotropy energy (MAE) are receiving consideration for their potential usage in solid-state electronic devices. This study explores various traits of the pristine (prs.)/Ir-doped (dop.) $\text{Sr}_2\text{CaOsO}_6$ structure using *ab initio* calculations, where Ir is doped at the Os-site. To determine the synthesis feasibility of the structures under ambient conditions, the formation energy, elastic constants, and phonon curves are determined. The prs. structure manifests a FM semiconducting nature with an E_g of 0.048 eV. Strikingly, the Ir-dop. structure becomes HM FIM because additional electrons provided by the dopant (Ir) cause a repulsion in the Os t_{2g}^2 spin-minority channel, resulting in conductivity. Conversely, an E_g of 1.15 eV in the spin-majority channel exists, which is high enough to keep the HM state stable. The computed partial spin-moment on the Os in the prs. system is 1.19 μ_B . In the Ir-dop. system it is 1.09/–1.39 μ_B on the Os/Ir ion holding an $\text{Os}^{+6}/\text{Ir}^{+4}$ state with electronic distributions of $5d^2(t_{2g}^{\uparrow}t_{2g}^{\downarrow}e_g^0\uparrow e_g^0)/5d^5(t_{2g}^{\uparrow}t_{2g}^{\downarrow}e_g^0\uparrow e_g^0)$ with $S = 1/(\frac{1}{2})$. Further, the spin-magnetization density isosurfaces assist in determining the m_s values and FM/FIM state of the prs./Ir-dop. system holding a Curie temperature (T_C) of 185/171 K. Besides this, we computed the thermoelectric properties of the prs./Ir-dop. motifs; the figure of merit (0.33/0.02), Seebeck coefficient (147/30 $\mu\text{V K}^{-1}$), and low thermal conductivity ($0.21/0.71 \times 10^{-19} \Omega\text{m}^{-1} \text{s}^{-1}$) at 300 K highlight their potential for conversion devices. Interestingly, a semiconducting-to-HM transition is predicted at a crucial compressive strain of –3% in the prs. structure. Conversely, the HM state in the dop. structure displays robustness against strain. Additionally, it is shown that an applied tensile strain can significantly improve ZT , while compressive strains illustrate a positive impact on the T_C value.

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1 Introduction

Depending on the B and B' cations, double perovskite oxides (DPOs) display a wide range of ferromagnetic (FM), ferrimagnetic (FIM), and anti-ferromagnetic (AFM) spin ordering (SO) in addition to half-metal (HM)^{1,2} and semiconducting (SC)/insulating phases,^{3–5} and an insulator-to-HM transition.⁶ Materials research continues to focus on the insulator-to-metal transitions (IMTs) because of the conceptual gaps that lead to an inability to regulate the electronic conduction in prospective

magnetic memory devices.⁷ Likewise, structure modifications in the crystals allow the material to transform from a band insulator to Mott insulator (MI).⁸ In particular, DPOs based on the 5d transition metals (TMs) provide an ideal environment for orbital, charge, and lattice degrees of freedom. In this respect, osmium-based materials have attracted a lot of interest due to the successful synthesis process of the oxides, which generates a variety of unusual phases. For example, a MI FM state in $\text{Ba}_2\text{NaOsO}_6$ (ref. 9) and a magnetic insulating state in Sr_2MOsO_6 (where M = Cu/Ni)¹⁰ are predicted. Interestingly, the HM FIM state has been discovered with an energy gap (E_g) of 0.40 eV in the spin-minority channel (N^{\downarrow}) of $\text{Sr}_2\text{CuOsO}_6$,¹¹ where the spin-majority channel (N^{\uparrow}) is metallic. The combined effect of electron correlation and spin–orbit coupling (SOC) results in a FIM MI character of the $\text{Ca}_2\text{FeOsO}_6$ structure.¹² Similarly, a half semi-metallic FIM phase in $\text{Sr}_2\text{CrOsO}_6$ is theoretically predicted within the GGA+U+SOC method¹³ owing to a FIM MI state with a massive T_C (Curie temperature (temp.)) of 725 K.¹⁴ Moreover, the $\text{Sr}_2\text{CrOsO}_6$ system displays a compensatory HM behavior

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without the influence of SOC. However, when SOC is included, a net magnetic moment of $0.54 \mu_B$ is obtained,¹³ which exposes that the dominating influence of SOC cannot be properly attributed to the magnetic moment correction.

Systems with high T_C and HM FM/FIM stable ground states are in huge demand for the fields of spintronics,^{15–17} magnetoelectronics,¹⁸ magnetodielectric capacitors,^{19,20} and data storage devices.²¹ It has been discovered that enhancing the antisite disorder deficiencies and grain boundaries may improve the physical aspects of DPOs.^{22,23} Hence, the d-orbital occupancy can be effectively regulated by doping the B/B' site with various TMs to significantly improve magnetic exchange interactions between them, which often results in high- T_C HM systems.^{24,25} The T_C for $\text{Sr}_2\text{CrOsO}_6$ jumps dramatically from 490 to 660 K when the larger Sr cation is substituted at the Ca-site.^{26–28} Furthermore, the inclusion of disordered BO_6 and $\text{B}'\text{O}_6$ octahedra accumulating irregularly leads to a crystal field splitting and enhances the unusual electronic and magnetic features.²⁹ The structural distortions are reduced when Ba is substituted at the Sr-site in AlaNiOsO_6 ($A = \text{Sr}, \text{Ba}$) and the Weiss temp. (θ_w) changes from negative to positive.³⁰ Experimental observations reveal that doping of Ni^{2+} at the Fe-site in $\text{Sr}_2\text{Fe}_{1.5}\text{Mo}_{0.5}\text{O}_{6-\delta}$ leads to competition between $\text{Fe}^{3+}/\text{Mo}^{5+}$ and $\text{Fe}^{2+}/\text{Mo}^{4+}$ along with induced conductivity.³¹ By adopting a similar strategy, Bhandari *et al.*,³² utilizing density functional theory (DFT) calculations, predicted that the system enters into an HM state when Ni^{2+} is doped (dop.) at the Cr^{3+} site (electron doping) in the FIM MI $\text{Ca}_2\text{CrOsO}_6$ system. In addition to doping of various elements at the A or B site in $\text{A}_2\text{BB}'\text{O}_6$, the doping of osmium at the B' site in $\text{Sr}_2\text{CrReO}_6$ significantly increases the T_C and causes an HM-to-insulator transition.³³ However, partial magnetizations in $\text{Sr}_2\text{CrReO}_6$ at the Re-site dramatically decrease at a 40% W-amount and a drop in T_C is also observed with an increase in dopant (W) concentration.³⁴

In the same way, the strain approach is one of the most effective means of adjusting or controlling a material's traits by altering the lattice parameters under various growth circumstances.^{35,36} For example, when a $\text{Sr}_2\text{FeMoO}_6$ film is formed on a SrTiO_3 substrate, lattice mismatch causes compressive (comp.) strain of -1.2% , leading to an HM FM state.^{36,37} Furthermore, at -8% comp. and $+1\%$ to $+5\%$ tensile (tens.) strains, a FM SC to metal and HM transition is verified in $\text{La}_2\text{-FeMnO}_6$, respectively.³⁸ Likewise, in the $\text{Lu}_2\text{NiIrO}_6$ DPO, a FIM SC to HM transition is noted under hydrostatic stress of -6% and biaxial strain of -8% .³⁹ Besides this, experimental observations reveal that lattice mismatch occurs when $\text{Re}_2\text{NiMnO}_6$ thin films (where $\text{Re} = \text{La}, \text{Pr}, \text{Nd}, \text{Sm}, \text{and Y}$) grow on a LaAlO_3 substrate; the B(B')O_6 octahedra are further deformed by induced strain, which improves the films' magnetic qualities and serves as an effective approach to get the maximum T_C in DPOs.⁴⁰ Interestingly, a magnetic transition from the FM SC to the HM FM state in $\text{La}_2\text{FeMnO}_6$ is demonstrated by DFT investigations with the application of a biaxial ($[[110]]$) strain ranging from 0% to $+10\%$.³⁸ Moreover, under volume compression, a HM state with an extremely high T_C of 624 K is predicted in the $\text{Ca}_2\text{MoOsO}_6$ structure.⁴¹

The above-mentioned considerations make it abundantly evident that B'-site doping in DPOs is also a suitable method for altering physical aspects. Thus, we theoretically study the effect of Ir-doping at the Os-site on the distinct traits of FM SC $\text{Sr}_2\text{-CaOsO}_6$ (SCOO). Because of the partial filling of the 5d orbitals of the dopant (Ir) and the host (Os), a SC-to-metal transition (MT) is observed in the Ir-doped (dop.) structure. Each system shows an appropriate amplitude of magnetic anisotropy energy (MAE), which improves the system's functionality for the data storage devices.²¹ Along with this, we do a thorough examination of the material's thermoelectric (TE) qualities for its use in renewable energy devices. Additionally, detailed investigations of the biaxial ($[[110]]$) strain effect on the physical behavior of the systems were conducted, which give interesting outputs.

2 Computational and structural details

A full-potential linearized augmented plane-wave approach based on spin-polarized (SP) DFT, as implemented in the WIEN2K code,⁴² was employed for the current calculations. The exchange–correlation functional utilized in this study, which is derived from the generalized gradient approximation (GGA) combined with on-site Coulomb interaction (GGA+U), is used, keeping U as 2.8/2.6 eV on the 5d states of the Ir/Os ion.⁴³ Additionally, because of the heavy Os/Ir element, SOC effects are also employed in the scalar relativistic form. In the wavefunction expansion within the atomic spheres, $l_{\text{max}} = 12$, $R_{\text{mt}} \times K_{\text{max}} = 7$, and $G_{\text{max}} = 24$ within the irreducible wedge of the Brillouin zone is taken. A $6 \times 6 \times 4$ k -mesh with 76 points is found to be highly converged. Also, the atomic positions of the ions are fully relaxed by lowering the overall forces below 5 mRy a.u.⁻¹. Self-consistency of the system is assumed for a total energy (E_t)/charge convergence to 10^{-5} Ry/ 10^{-5} C. Moreover, Boltzmann's theory is used to compute the TE parameters using the relaxation-time approximation, as defined in the BoltzTrap code.⁴⁴

The monoclinic SCOO keeps CaO_6 and OsO_6 octahedra alternately organized in a rock-salt pattern, providing a full 1 : 1 ordering with space group no. 14 ($P2_1/n$). The experimental lattice constants are $a = 5.7643$, $b = 5.8191$, and $c = 8.1796$ Å

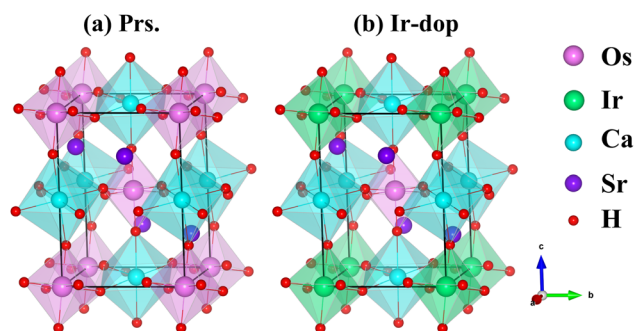


Fig. 1 Crystal diagram of the (a) prs. and (b) Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures.



with $\beta = 90.22^\circ$.⁴⁵ In the primitive unit cell of SCO, 4/2/2/12 Sr/Ca/Os/O ions exist. The atomic coordinates for the Sr, Ca, Os, O₁, O₂, and O₃ are (0.0104, 0.0334, 0.2490), (0.5, 0, 0), (0.5, 0, 0.5), (0.265, 0.312, 0.028), (−0.186, 0.234, −0.038), and (−0.088, 0.479, 0.227), correspondingly. The Ir-dop. SCO structure is designed by replacing one of the Os ions with one Ir in its primitive unit, as reported in the previous experimental^{46,47} and theoretical^{5,48} works, where it changes the $P2_1/n$ to $P1$ structural symmetry. The crystal structure of the prs./Ir-dop. motif is presented in Fig. 1(a) and (b).

3 Results and discussion

3.1 Unstrained systems

First, the thermodynamic stability of the structures is examined by determining the formation enthalpy (ΔH_f) as follows:

$$\Delta H_f^{\text{prs.}} = E_{\text{t}}^{\text{Sr}_4\text{Ca}_2\text{Os}_2\text{O}_{12}} - 4E_{\text{t}}^{\text{Sr-fcc}} - 2E_{\text{t}}^{\text{Ca-fcc}} - 2E_{\text{t}}^{\text{Os-hcp}} - \frac{12}{2}E_{\text{t}}^{\text{O}_2\text{-monoclinic}} \quad (1)$$

$$\Delta H_f^{\text{Ir-dop.}} = E_{\text{t}}^{\text{Sr}_4\text{Ca}_2\text{OsIrO}_{12}} - 4E_{\text{t}}^{\text{Sr-fcc}} - 2E_{\text{t}}^{\text{Ca-fcc}} - E_{\text{t}}^{\text{Os-hcp}} - E_{\text{t}}^{\text{Ir-fcc}} - \frac{12}{2}E_{\text{t}}^{\text{O}_2\text{-monoclinic}} \quad (2)$$

$E_{\text{t}}^{\text{Sr}_4\text{Ca}_2\text{Os}_2\text{O}_{12}}/E_{\text{t}}^{\text{Sr}_4\text{Ca}_2\text{OsIrO}_{12}}$, $E_{\text{t}}^{\text{Sr-fcc}}$, $E_{\text{t}}^{\text{Ca-fcc}}$, $E_{\text{t}}^{\text{Os-hcp}}$, $E_{\text{t}}^{\text{Ir-fcc}}$ and $E_{\text{t}}^{\text{O}_2\text{-monoclinic}}$ represent the E_{t} of the prs./Ir-dop. SCO, Sr ($Fm\bar{3}m-225$), Ca ($Fm\bar{3}m-225$), Os ($P6_3/mmc-194$) and Ir ($Fm\bar{3}m-225$) atoms and oxygen ($C2/m-12$) molecule in their respective ground states, correspondingly. To prevent the overestimation of the ΔH_f value, the suggested adjustments by Wang *et al.*,⁴⁹ are also considered. The estimated ΔH_f of the prs./Ir-dop. structure is −29.25/−25.85 eV, where the “−” sign ensures the thermodynamic stability of the system. Next, to determine the mechanical stability of both structures, we calculated the elastic tensors (C_{ij}) by generating six finite constants using conventional strain and stress relations.^{50,51} The computed 13 independent elastic stiffness tensors for the monoclinic systems are listed in Table 1, which fulfilled the basic requirements and Born stability criteria⁵² for mechanical stability. Additionally, Fig. 1S of the ESI† displays the bulk modulus (B), shear modulus (G), and Young modulus (Y) of both systems. B measures a compound's ability to deform in response to the pressure surrounding its surface, commonly known as hardness. Along with this, a higher value of G suggests that the material is stiffer

and less likely to deform under shear forces, whereas lower values indicate that it is more flexible or ductile. Y is also called the modulus of elasticity and is a measure of a material's stiffness. A high value means that the materials are stiff and resistant to deformation under stress, while lower denotes a more elastic and easily deformed material. Hence, the B , G , and Y values of the prs./Ir-dop. system are 129/128, 145/142, and 55/54, respectively. It shows that the prs. system is stiffer than the doped one. Furthermore, we computed Pugh's ratio (B/G), Cauchy's pressure (C_P), and Poisson's ratio (ν), which illustrate whether a compound is ductile or brittle,⁵³ as displayed in Fig. 2. If the B/G value is less than 1.75, ν is less than 0.25, and the C_P value is less than 0, this reveals a brittle nature. In contrast, if the B/G value is greater than 1.75, ν is greater than 0.25, and the C_P value is greater than 0, this results in a ductile nature.⁵⁴ Hence, our computed values of the B/G (see Fig. 2(a)), C_P (see Fig. 2(b)) and ν (see Fig. 2(c)) for both structures demonstrate that they are ductile. Finally, phonon calculations were performed for both structures to confirm their dynamical stability. The determined phonon spectra for the prs./Ir-dop.

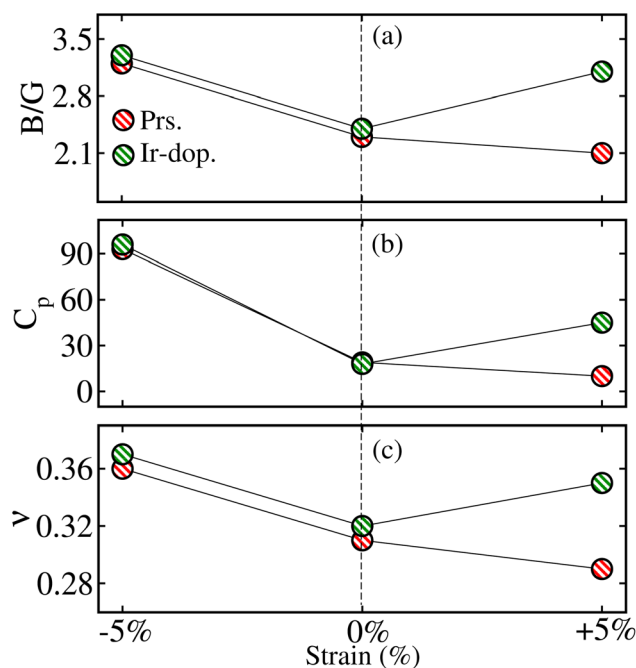


Fig. 2 Computed (a) Pugh's ratio (B/G), (b) Cauchy's pressure (C_P), and (c) Poisson's ratio (ν) in the prs./Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structure for −5% (compressive)/0% (unstrained)/+5% (tensile) strain.

Table 1 Computed 13 independent elastic constants (C_{ij}) of the prs. and Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures under −5%/0% (unstrained)/+5% biaxial ([110]) strain

Systems	Strain	C_{11}	C_{12}	C_{13}	C_{15}	C_{22}	C_{23}	C_{25}	C_{33}	C_{35}	C_{44}	C_{46}	C_{55}	C_{66}
prs.	−5%	203.70	144.29	164.91	6.71	204.85	162.51	16.06	258.55	−4.48	69.14	−4.86	62.03	101.73
	0%	193.76	113.58	90.42	3.94	188.95	78.33	−3.98	222.13	4.44	59.37	−6.03	49.71	67.45
	+5%	144.22	68.45	52.47	−0.95	128.78	56.99	−2.54	183.14	0.98	46.39	−2.03	32.00	45.07
Ir-dop.	−5%	201.35	144.26	165.27	5.13	200.23	162	14.39	252.84	−6.38	66.81	−3.98	60.87	102.21
	0%	190.53	113.98	90.20	4.09	183.40	75.84	−3.81	222.68	3.36	57.46	−5.58	47.99	68.69
	+5%	189.78	94.99	76.84	7.37	149.68	96.63	14.68	187.91	−17.41	51.62	−4.55	25.27	67.67



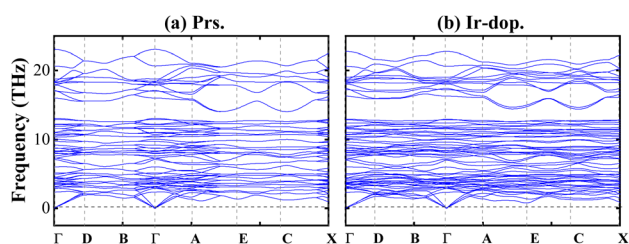


Fig. 3 Computed phonon dispersion curves for the (a) prs. and (b) Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures.

systems are plotted in Fig. 3(a) and (b). In principle, each atom typically contributes three phonon branches with the total number of phonon branches in a unit cell being precisely proportional to the three times number of atoms. There are $3n$ total branches, which include $3n - 3$ optical modes and 3 acoustic modes.⁵⁵ Both structures contain 20 atoms in their basic unit cell and have 60 vibrational modes including 3 acoustic modes along with 57 optical modes. The phonon dispersion curves do not contain any negative frequencies for both structures (see Fig. 3), indicating that they are dynamically stable as well.

To find the magnetic ground state (MGS) of the prs. system, we plotted the computed E_t for the non-magnetic (NM), FM, and AFM MGS spin ordering (SO) in Fig. 2S of the ESI† within the GGA+U/GGA+U+SOC scheme. In the case of the NM phase, both Os ions remain non-SP. In contrast, they are aligned ($\uparrow\uparrow$)/anti-aligned ($\uparrow\downarrow$) for the FM/FIM SO. It is demonstrated that the E_t of the FM SO is lower than that of the NM/AFM state in both methods (see Fig. 2S of the ESI†). However, it is experimentally observed that the system persists in a paramagnetic state for 600 K to 2 K temp. and there is no magnetic phase transition occurring.⁴⁵ Additionally, the computed E_t values of the FM and FIM SO are compared to determine the MGS for the Ir-dop. SCOO DPO. In the case of the FM/FIM SO, Os and Ir ions, the spins remain parallel ($\uparrow\uparrow$)/anti-parallel ($\uparrow\downarrow$) to each other, where the small and large arrow lengths represent their respective spin magnitudes. It is established that the FIM state is more stable than the FM one with an energy difference of $\Delta E = E_{\text{FIM}} - E_{\text{FM}} = -96 \text{ meV}$ (see Table 2). This indicates that Ir and Os ion spins favor anti-alignment in both the in-plane and out-of-plane orientations. Hence, for further investigations only FM/FIM SO is considered for the prs./Ir-dop. motif.

Next, to examine the electronic structure of the prs./Ir-dop. SCOO system, we computed the total density of states (TDOS) in the stable FM/FIM SO within the GGA+U method. Fig. 4(a) shows that the prs. motif is SC with an E_g of 0.048 eV (as listed in Table 2). The remarkable finding of the present work is that a transition from the SC to HM state occurs in the Ir-dop. structure (see Fig. 4(b)), where a few states cross the Fermi level (E_F) in the N^\downarrow , while it contains a definite large E_g of 1.15 eV in the N^\uparrow .

Additionally, we presented the orbital resolved partial density of states (PDOS) on the 5d-states of the Os and Ir/Os ions in the prs. and Ir-dop. system in Fig. 3S of the ESI† to better examine the states close to E_F . The Os 5d states are dominant at the valence and the conduction band edges (VBE and CBE) as

Table 2 Computed total energy difference (ΔE), energy gap in the spin majority/minority channel (E_g^N/E_g^{\downarrow}), total/partial spin/orbital moment ($m_t/m_s/m_{\text{orb}}$) in μ_B , magnetocrystalline anisotropy energy (MAE) in meV, Curie temperature (T_C) and MAE constant (K) per unit volume ($\times 10^7 \text{ erg cm}^{-3}$) within the GGA+U/GGA+U+SOC method for the prs. and Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures. The symbol M symbol represents the metal

Property	GGA+U		GGA+U+SOC	
	prs.	Ir-dop.	prs.	Ir-dop.
ΔE	−51	−131.8	—	−96.4
$E_g^{N^\uparrow}$	0.048	1.15	0.004	M
$E_g^{N^\downarrow}$	1.86	M	—	—
MAE	—	—	2.95	0.79
K	—	—	1.72	0.46
T_C	185	171	—	—
m_t	4.0	1.0	3.84	0.91
	Os1	Os2	Os	Ir
m_s	1.19	1.19	1.09	1.09
m_{orb}	—	—	−0.5	−0.5
	Os1	Os2	Os	Ir
	1.19	1.01	−1.39	0.89
	—	—	—	1.25
	—	—	—	0.44
	—	—	—	0.083

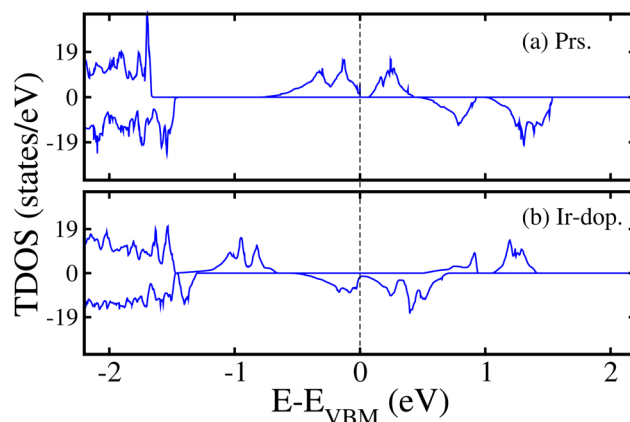


Fig. 4 GGA+U computed non-degenerate total density of states (TDOS) in the (a) prs. and (b) Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures.

demonstrated in Fig. 3S(a) of the ESI†. As Os is in a $+6(5d^2)$ oxidation state, this results in the filling of two t_{2g} states in the N^\uparrow and the e_g states are empty. Hence, the filled states are in the VB, while the two empty states shift towards the CB. For the Ir-dop. structure, metallicity in the N^\downarrow mainly arises from the Os-5d states along with substantial contributions from Ir-5d states (see Fig. 3S(b) of the ESI†). It is found that the Os-5d states become partially occupied and significantly shift towards lower energies in the CB, which leads the system into the HM phase. This happens because the Ir ion is in a $+4(5d^5)$ oxidation state and it contributes 3 additional electrons to the system; one of the additional electrons occupies the t_{2g}^\downarrow and two occupy the t_{2g}^\uparrow state. Hence, they produce a repulsive force in the Os t_{2g} states, causing the Os states to move towards the CB from the VB in the N^\downarrow . Along with this, we plotted the computed non-degenerate band structures for both systems within the GGA+U scheme in Fig. 5 to provide further confirmation of their electronic states. The corresponding MI/HM behavior in the



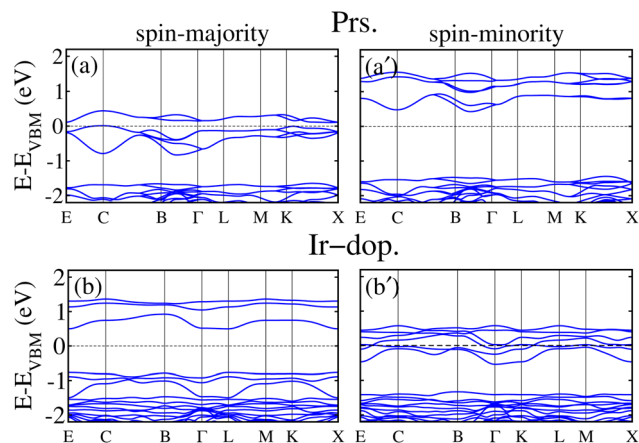


Fig. 5 GGA+U computed non-degenerate spin-majority (left column)/spin-minority (right column) band structures for the (a and a') prs. and (b and b') Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures.

prs./Ir-dop. structure is well illustrated by Fig. 5(a, a') and (b, b'), which also support the estimated TDOS in Fig. 4(a) and (b). Moreover, we plotted the GGA+U+SOC computed TDOS in Fig. 4S of the ESI† for the prs./Ir-dop. SCOO system in the most stable FM/FIM SO. A small E_g of 0.0045 eV exists in the prs. system (see Fig. 4S(a) of the ESI†), while the Ir-dop. motif turns metallic (see Fig. 4S(b) of the ESI†). It is predicted that the E_g computed within the GGA+U+SOC scheme for the prs. motif decreases (see Fig. 4S(a) of the ESI†) as compared to the GGA+U one (see Fig. 4(a)). This is because relativistic effects and SOC interactions often lower the E_g by mixing the orbitals in a way that reduces the energy splitting of the electronic states close to the E_F .

Now, the system's magnetism is described by calculating the total/partial spin magnetic moment (m_t/m_s) in each case along with the three-dimensional (3D) spin magnetization density isosurfaces. The estimated m_t for the prs./Ir-dop. system is 4.0/1.0 μ_B f.u.⁻¹ (see Table 2). The computed m_s for the Os1/Os2 ion in the prs. system is 1.19 μ_B , whereas the m_s on the Os/Ir is 1.01/−1.39 μ_B for the Ir-dop. structure, as displayed in Table 2. The “−” sign shows that Ir and Os ions' m_s are aligned antiparallel ($\uparrow \downarrow$) to each other, which indicates that an AFM coupling is dominant, and turns out to be a FIM SO in the Ir-dop. structure. Further, the computed m_s on the Os/Ir assures that they are in a $+6(t_{2g}^2 \uparrow t_{2g}^0 \downarrow e_g^0 \uparrow e_g^0 \downarrow)/+4(t_{2g}^3 \uparrow t_{2g}^2 \downarrow e_g^0 \uparrow e_g^0 \downarrow)$ state. Moreover, the computed m_{orb} on the Os is −0.5 μ_B in the prs. and 0.44/0.08 μ_B on the Os/Ir in the Ir-dop. structure. Also, the 3D spin magnetization density isosurfaces, keeping an iso-value of $\pm 0.05\% \text{ e } \text{\AA}^{-3}$, are plotted in Fig. 6 for the direct observation of m_s as well as to further affirm the SO in both structures. An appropriate density arises around the Os ions in the prs. system (see Fig. 6(a)), where the same density colors confirm that their spins' parallel ($\uparrow \uparrow$) alignment and t_{2g}^2 orbital characterization is visualized due to its +6 state. Meanwhile in the Ir-dop. motif, the magnitude of density on the Ir ion is greater than that on the Os ion, which further verifies the magnitude of the calculated m_s on each ion. Besides this, the color contrast of the densities also

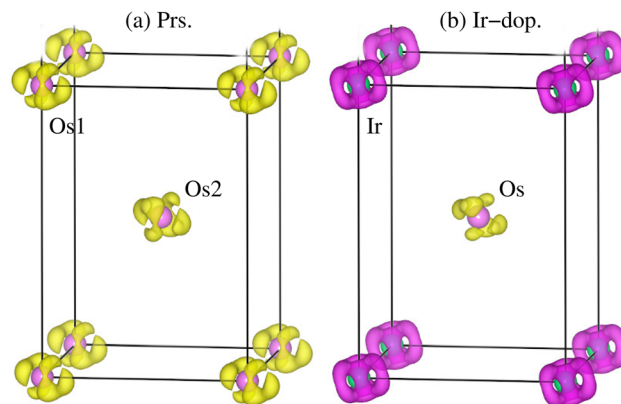


Fig. 6 Computed spin magnetization density iso-surfaces for the (a) prs. and (b) Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures with an iso-value of $\pm 0.05\% \text{ e } \text{\AA}^{-3}$.

assures that the spins of both ions align antiparallel ($\uparrow \downarrow$) to each other.

Now, the FM SO in the prs. structure is explained by virtual hopping between less-than-half-filled d-orbitals *via* oxygen ($\text{Os}^{+6}-\text{O}^{2-}-\text{Os}^{+6}$), as depicted in Fig. 7(a). Since the B-site atom is a non-magnetic cation, the magnetic aspects are determined by exchange coupling within the Os^{+6} ion, and a FM superexchange interaction occurs at 180° . But in the Ir-dop. system, the electron hopping takes place between the half-filled d-orbitals of Os^{+6} and half-filled d-orbitals of the Ir^{+4} ion, which occurs 180° between Os and Ir ions as $\text{Os}^{+6}-\text{O}^{2-}-\text{Ir}^{+4}$ (see Fig. 7(b)). This supports the strong AFM superexchange

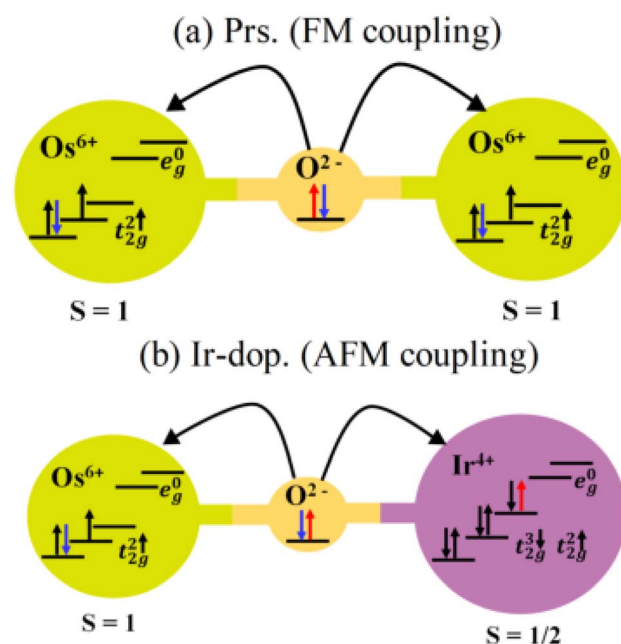


Fig. 7 The superexchange interaction process between (a) Os^{+6} 5d and Os^{+6} 5d *via* oxygen results in a ferromagnetic (FM) coupling in the prs. and that between (b) Ir^{+4} 5d and Os^{+6} 5d *via* oxygen results in a strong antiferromagnetic (AFM) coupling in the Ir-doped $\text{Sr}_2\text{CaOsO}_6$ structure, which leads to ferromagnetic ordering.



mechanism in this structure, resulting in the FIM SO. Now, it is a well-established fact that MAE plays an important part in defining coercivity, which is required for permanent magnets with long-term magnetization.^{56,57} Therefore, we investigated the MAE and MAE constant $\left(K = \frac{\text{MAE}}{V}\right)$ as well as its relation to the structural distortions. To do this, we set the magnetization direction along the [001]/[010]/[100]-axis and compare E_t . Moreover, a dense k -mesh of $12 \times 12 \times 9$ is utilized to verify that the E_t and MAE values converged correctly. Our results reveal that the prs. system has a MAE/ K of 2.95 meV/ 1.72×10^7 erg per cm^3 with an easy axis of [100] and average $\angle \text{Os-O-Os}$ of 153.2° . In comparison, the Ir-dop. system has a lower MAE/ K of 0.79 meV/ 0.42×10^7 erg per cm^3 , as listed in Table 2, with an easy axis of [010] and an average $\angle \text{Os-O-Ir}$ of 153.6° . This clearly illustrates that the MAE value is highly dependent on the structural distortions as reported in previous works.^{5,21}

Next, to evaluate the T_C of the prs./Ir-dop. SCOO motif, the exchange constants (J) are determined using the Heisenberg model, as follows:⁵⁸

$$H = -\sum_{i \neq j} J_{ij} \vec{S}_i \cdot \vec{S}_j \quad (3)$$

where \vec{S}_i/\vec{S}_j specifies the spin vectors and can be calculated as $S(S+1)^{1/2}$. The S indicates the m_s on each magnetic ion, where J represents the interactions between ions. The E_t of FM, FIM, and AFM SO can be written as:

$$E_t^{\text{FM}} = E_0 + (S_1(S_1 + 1))^{1/2}(S_2(S_2 + 1))^{1/2}(2J_1 + 4J_2) \quad (4)$$

$$E_t^{\text{FIM}} = E_0 + (S_1(S_1 + 1))^{1/2}(S_2(S_2 + 1))^{1/2}(-2J_1 + 4J_2) \quad (5)$$

$$E_t^{\text{AFM}} = E_0 + (S_1(S_1 + 1))^{1/2}(S_2(S_2 + 1))^{1/2}(-2J_1 - 4J_2) \quad (6)$$

E_0 refers to the energy in the spin-degenerate system and $S_1(S_2) = 1(1)$ for $\text{Os}_1^{+6}/\text{Os}_2^{+6}$ in the prs. system and $S_1(S_2) = 1\left(\frac{1}{2}\right)$ for $\text{Os}^{+6}(\text{Ir}^{+4})$ in the Ir-dop. motif. Hence, T_C can be determined as follows:⁵⁹

$$T_C^{\text{prs.}} = \frac{2}{3k_B}(S_1(S_1 + 1))^{1/2}(S_2(S_2 + 1))^{1/2}(2J_1 + 4J_2) \quad (7)$$

$$T_C^{\text{Ir-dop.}} = \frac{2}{3k_B}(S_1(S_1 + 1))^{1/2}(S_2(S_2 + 1))^{1/2}(-2J_1 + 4J_2) \quad (8)$$

Our results revealed that the determined T_C is 185/171 K for the prs./Ir-dop. motif, as listed in Table 2. As structural distortions are reduced in the Ir-dop. system as compared to the prs. one, this lowers the T_C a bit, which is also consistent with the previous reports that T_C decreases with a reduction in structural distortion.⁵

Next, TE parameters are computed to provide insight into these materials for their potential realization in devices that convert heat into electricity. The BoltzTrap code⁴⁴ is used to calculate the TE factors, such as the electrical conductivity (σ) per relaxation time $\left(\frac{\sigma}{\tau}\right)$, Seebeck coefficient (S), electronic thermal conductivity (κ_e) per relaxation time $\left(\frac{\kappa_e}{\tau}\right)$, susceptibility

(χ), power factor $\left(\text{PF} = \frac{S^2\sigma}{\tau}\right)$, and figure of merit $\left(ZT = \frac{\sigma S^2 T}{\kappa_e}\right)$ at temp. ranging from 200 to 600 K, as presented in Fig. 8. As efficient TE materials require a high value of $\frac{\sigma}{\tau}$, the variation in temp.-dependent $\frac{\sigma}{\tau}$ is illustrated in Fig. 8(a); it depends on the concentration of the free carriers (electrons or holes) and increases with temp. due to an increase in kinetic energy. The values of $\frac{\sigma}{\tau}$ are higher for the Ir-dop. motif, because there is a greater concentration of free charge carriers at E_F . Its value is $0.21/0.71 \times 10^{19} \Omega \text{m}^{-1} \text{s}^{-1}$ at 300 K and reaches $0.45/0.9 \times 10^{19}$ at 600 K for the prs./Ir-dop. structures. The S value, which is another important TE factor for determining the electronic transport properties, measures the TE capacity of a material to generate a potential difference across its edges as a result of a gradient. The temp. difference supports the passage of carriers, creating a gradient that can be determined in $\mu\text{V K}^{-1}$. Fig. 8(b) demonstrates that S varies significantly between 30 to $200 \mu\text{V K}^{-1}$ for the considered temp. range. The computed S values for the prs./Ir-dop. structure are 147/30 $\mu\text{V K}^{-1}$ and 85/37 $\mu\text{V K}^{-1}$ at 300 K and 600 K, respectively. Hence, one can conclude from Fig. 8(a) and (b) that the behavior of S against temp. is in contrast to that of $\frac{\sigma}{\tau}$: whereas S decreases with temp., $\frac{\sigma}{\tau}$ increases. In addition to $\frac{\sigma}{\tau}$, Fig. 8(c) displays $\frac{\kappa_e}{\tau}$, which shows an increase with temp., being $0.41/0.84 \times 10^{14}$ at 300 K and $1.6/2.1 \times 10^{14} \text{W m}^{-1} \text{K}^{-1} \text{s}^{-1}$ at 600 K for the prs./Ir-dop. system. Next, the term χ describes the response of the TE materials to variations in external conditions such as electric or magnetic fields, which may have an impact on their TE performance. A higher temp. can enhance the TE performance of a material because the thermal energy pushes more charge carriers into the CB, increasing sensitivity to external forces. Hence, Fig. 8(d) exhibits that χ increases with the rise in temp. for the prs./Ir-dop. structure, from $0.9/1.6 \times 10^{-9} \text{m}^3 \text{mol}^{-1}$ to $1.5/1.8 \times 10^{-9} \text{m}^3 \text{mol}^{-1}$. Similarly, the power factor (PF) is also an important factor to consider when measuring a material's TE efficiency. The trend of the PF is like that of S , which decreases with an increase in temp. The calculated value of PF at 300 K is $0.45/0.064 \times 10^{11} \text{W m}^{-1} \text{K}^{-2} \text{s}^{-1}$ and at 600 K is $0.32/0.13 \text{W m}^{-1} \text{K}^{-2} \text{s}^{-1}$ for the prs./Ir-dop. system (see Fig. 8(e)). Likewise, ZT is a critical component in determining TE efficiency for practical applications. The computed value of ZT at 300 K is 0.33/0.02 and at 600 K is 0.13/0.038 for the prs./Ir-dop. system (see Fig. 8(f)).

The net thermal conductivity (κ) is calculated as follows:

$$\kappa = \kappa_e + \kappa_l \quad (9)$$

Slack's equation⁶⁰ was used to calculate κ_l as follows:

$$\kappa_l = \frac{A M_{\text{av}} \theta^3 \delta}{\gamma n^{2/3} T} \quad (10)$$

M_{av} , δ , n , T , γ , and θ represent the average atomic mass in the crystal, cubic root of the average atomic volume, total number of



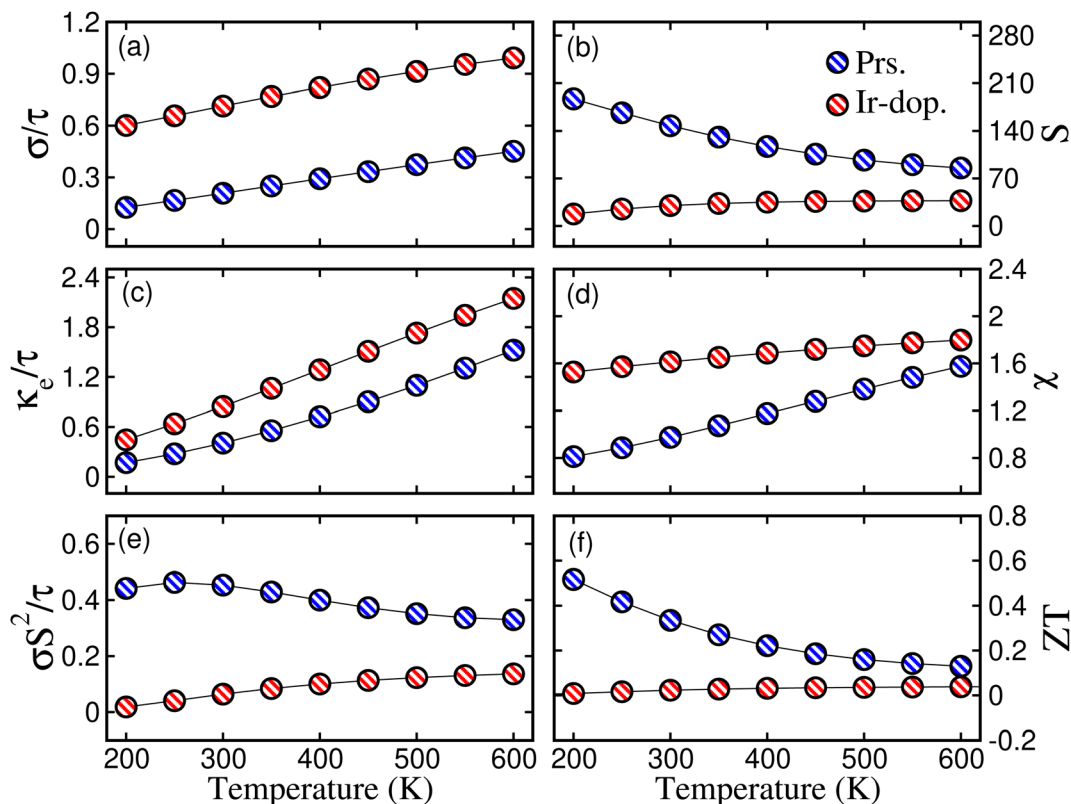


Fig. 8 Computed (a) electrical conductivity per relaxation time ($\frac{\sigma}{\tau}$) in $\times 10^{19} \Omega^{-1} \text{ s}^{-1}$, (b) Seebeck coefficient (S) in $\mu\text{V K}^{-1}$, (c) thermal conductivity per relaxation time ($\frac{\kappa_e}{\tau}$) in $\times 10^{14} \text{ W m}^{-1} \text{ K}^{-1} \text{ s}^{-1}$, (d) susceptibility (χ) in $\times 10^{-9} \text{ m}^3 \text{ mol}^{-1}$, (e) power factor (PF) in $\times 10^{11} \text{ W m}^{-1} \text{ K}^{-2} \text{ s}^{-1}$, and (f) figure of merit (ZT) for the prs./Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structure.

atoms in the unit cell, absolute temp., Grüneisen parameter, and Debye temp., respectively. The Grüneisen parameter is determined *via* Poisson's ratio:

$$\gamma = \frac{3(1 + \nu)}{2(2 - 3\nu)} \quad (11)$$

where A is calculated as follows:⁶¹

$$A = \frac{2.43 \times 10^{-8}}{1 - \frac{0.514}{\gamma} + \frac{0.228}{\gamma^2}} \quad (12)$$

while θ can be determined as follows:⁶²

$$\theta_D = \frac{h}{k_B} \left[\frac{3n}{4\pi} \left(\frac{N_A \rho}{M} \right) \right]^{\frac{1}{3}} \times v_m \quad (13)$$

where h , k_B , n , N_A , ρ , M , and v_m serve as Planck's constant, the Boltzmann constant, the total number of atoms per unit cell, Avogadro's number, density, molecular weight, and sound velocity, respectively. The average sound velocity (v_m) is figured out using

$$v_m = \left[\frac{1}{3} \left(\frac{2}{v_l^3} + \frac{1}{v_t^3} \right) \right]^{-\frac{1}{3}} \quad (14)$$

Here, v_l/v_t is the longitudinal/transverse velocity, which can be measured as follows:

$$v_t = \sqrt{\frac{G}{\rho}} \quad (15)$$

$$v_l = \sqrt{\frac{3B + 4G}{3\rho}} \quad (16)$$

Fig. 9 shows that the estimated $\frac{\kappa_l}{\tau}$ decreases with increasing temp., which is a desired property for TE applications. The decrease in $\frac{\kappa_l}{\tau}$ as temp. rises is due to increased phonon scattering caused by higher lattice vibrations, which decreases the mean free route of phonons. The established values of $\frac{\kappa_l}{\tau}$ for the prs. and Ir-dop. systems are 0.68 and $0.69 \times 10^{14} \text{ W m}^{-1} \text{ K}^{-1}$, respectively.

Here, we would like to mention that the Slack method predominantly considers acoustic phonon contributions, disregarding optical phonon modes, and complicated anharmonic interactions, which may be crucial in materials characterized by poor symmetry or complicated bonding environments. Consequently, the model may generate considerable error in the calculation of κ_l , thereby impacting the precision of resultant TE parameters including ZT . Recent investigations have underscored the ambiguities linked to the Slack model. For example,



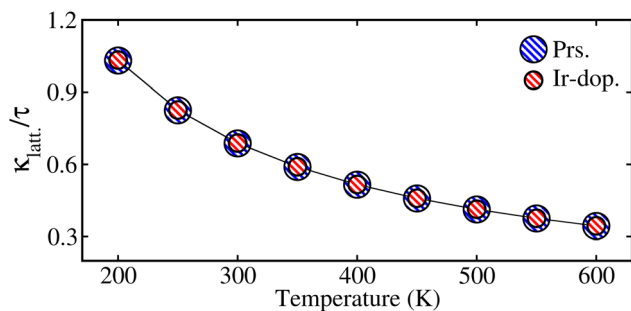


Fig. 9 Computed values of lattice thermal conductivity per relaxation time ($\frac{\kappa_{\text{lat}}}{\tau}$) in $\times 10^{-14} \text{ W m}^{-1} \text{ K}^{-1}$ with temperature, for the prs./Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structure.

Rabin *et al.*⁶³ shown that the Slack formula often overestimates κ in materials characterized by low group velocities and substantial optical phonon contributions, which are particularly influential in materials exhibiting poor symmetry or intricate bonding environments. Consequently, the model may generate significant error in the calculation of κ_{I} , possibly compromising the precision of derived thermoelectric parameters, including the ZT . Likewise, J. Carrete *et al.*⁶⁴ also noted that the empirical coefficient in the Slack equation is not universal as it changes according to the Grüneisen parameter and the bonding properties of the material. Hence, inaccuracies in κ_{I} can influence the assessment of the thermoelectric ZT , either under-representing or overstating a material's potential, thereby impacting the precision of derived thermoelectric parameters such as the ZT . The Slack model provides a valuable first estimate. However, enhanced accuracy in predictions can be attained by resolving the Boltzmann transport equation for phonons using first-principles interatomic force constants. In future endeavors, we intend to integrate sophisticated methodologies to authenticate and enhance the outcomes derived from the empirical methodology.

Usually, the empirical Slack model offers a useful initial estimate of κ , but it fails to adequately represent phonon transport dynamics in intricate materials. Recent developments in computational materials science have resulted in the emergence of first-principles approaches that offer a more precise and physically substantiated framework for forecasting thermal transport parameters. One technique involves solving the Peierls-Boltzmann transport equation (PBTE) alongside interatomic force constants derived from density functional perturbation theory (DFPT). These approaches consider both acoustic and optical phonons and comprehensively capture phonon-phonon scattering processes, allowing more accurate predictions across a diverse array of materials. Ma *et al.*⁶⁵ and Lindsay⁶⁶ examine the theoretical basis and practical application of these first-principles methodologies. Ma *et al.* emphasise the accomplishments and persistent difficulties in modeling heat transport using PBTE + DFPT, particularly in low-dimensional and disordered systems. Lindsay offers a comprehensive examination of the concept and illustrates its efficacy in precisely modeling κ across several crystalline materials.

4 Strained systems

Here, we examine the biaxial ([110]) strain consequences for the structural, mechanical, and dynamical stability of the SCO system by optimizing the two-axis lattice parameters a and b in the range from -5% to $+5\%$. First, the thermodynamical stability of the systems is examined by calculating ΔH_{f} under applied strain, and the results are shown in Fig. 5S of the ESI†. It is clear that ΔH_{f} is negative within all strain ranges; however, it is more negative under tens. strains, which means that these structures are more stable than the comp. strained ones. Next, we determine the mechanical stability of the structures under strain; 13 independent elastic constants are listed in Table 1, which obey the Born criteria and confirm mechanical stability against strain. Further, Fig. 1S of the ESI† displays the $B/Y/G$ values, which show that the prs. system under -5% comp. strain is stiffer and harder than the tens. strain ones. Additionally, we plotted the B/G , C_{p} , and ν values of both systems under -5% and $+5\%$ strain in Fig. 2, which depicted that structures are ductile and show more strength for -5% comp. strain. Moreover, we computed the phonon dispersion curves under two maximum strains of $\pm 5\%$, as shown in Fig. 6S of the ESI†, to ensure the dynamic stability of the motifs. This shows that no negative frequencies exist, indicating that the systems are also dynamically stable for the considered higher strain values. Subsequently, the magnetic ground state stability of the Ir-dop. SCO system is examined by plotting $\Delta E = E_{\text{FIM}} - E_{\text{FM}}$ against strain, as shown in Fig. 7S of the ESI†. It is found that ΔE remains negative within all considered strain ranges, which indicates that FIM is more stable than the FM SO in the whole strain range.

To qualitatively describe how the electronic structure of the systems varies under strain, we plotted the computed E_{g} in the N^{\uparrow} and N^{\downarrow} against strain, as shown in Fig. 10. When tens. strain is varied from 0% to $+5\%$, a slight linear increase in E_{g} from 0.048 to 0.050 eV in the N^{\uparrow} is observed (see Fig. 10(a)) for the prs. system. Conversely, when comp. strain is varied from 0% to -2% , the computed E_{g} decreases from 0.048 to 0.027 eV and ultimately becomes zero at a crucial value of -3% in the N^{\uparrow} (see Fig. 10(a)), while N^{\downarrow} maintains a definite E_{g} for the whole strain range. Thus, the system exhibits a SC-to-HM transition at a crucial -3% strain and for higher comp. strains as well. Likewise, when the Ir-dop. system is subjected to strain from 0% to $+5\%$, E_{g} decreases from 1.15 to 1.12 eV in the N^{\uparrow} and decreases more significantly under comp. strain from 1.15 eV to 0.75 eV (see Fig. 10(b)). In contrast, N^{\downarrow} preserves its metallic character for the whole strain range, which ensures the robustness of the HM phase in a doped system. In addition, we plotted the corresponding SP TDOS from -1% to -5% for the prs. system in Fig. 11, which also assures that no states lie at E_{F} and a small/large E_{g} exists in the $\text{N}^{\uparrow}/\text{N}^{\downarrow}$ for -1 and -2% strains (see Fig. 11(a) and (b)), correspondingly. However, the states overlap at the E_{F} in the N^{\uparrow} and E_{g} becomes zero for a crucial -3% strain (see Fig. 11(c)), while a large E_{g} of 1.86 eV exists in the N^{\downarrow} . Hence, a SC-to-HM transition occurs at a crucial comp. strain of -3% for the prs. system. Moreover, N^{\uparrow} becomes more metallic for the $-4\%/-5\%$ strain (see Fig. 11(d) and (e)) as more electrons reside



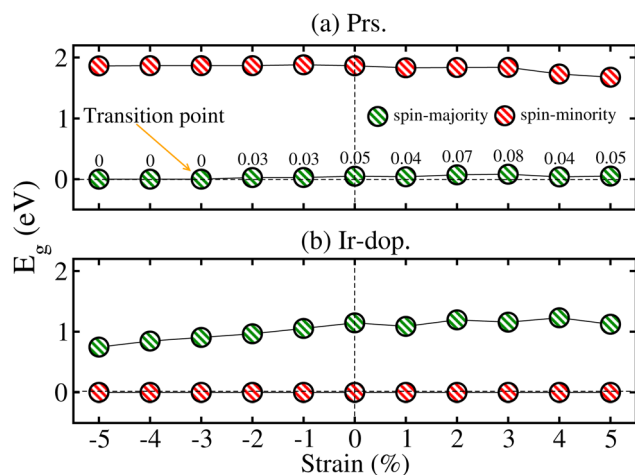


Fig. 10 GGA+U calculated energy gap (E_g) against $\pm 5\%$ biaxial ([110]) strain in the spin majority (N^\uparrow)/spin-minority (N^\downarrow) channel of the (a) prs. and (b) Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures.

at E_F for these strain levels. Along with this, we plotted the SP TDOS for the tens. strain of +1% to +5% in the prs. system in Fig. 8S of the ESI,[†] which shows that no transition occurs at any strain value and a definite E_g exists in both channels. Hence, the system holds its insulating behavior for all tens. strain levels. Likewise, the calculated TDOS for the Ir-dop. under $\pm 5\%$ biaxial ([110]) strains is plotted in Fig. 9S of the ESI,[†] which is consistent with Fig. 10 and no significant changes occur with strain.

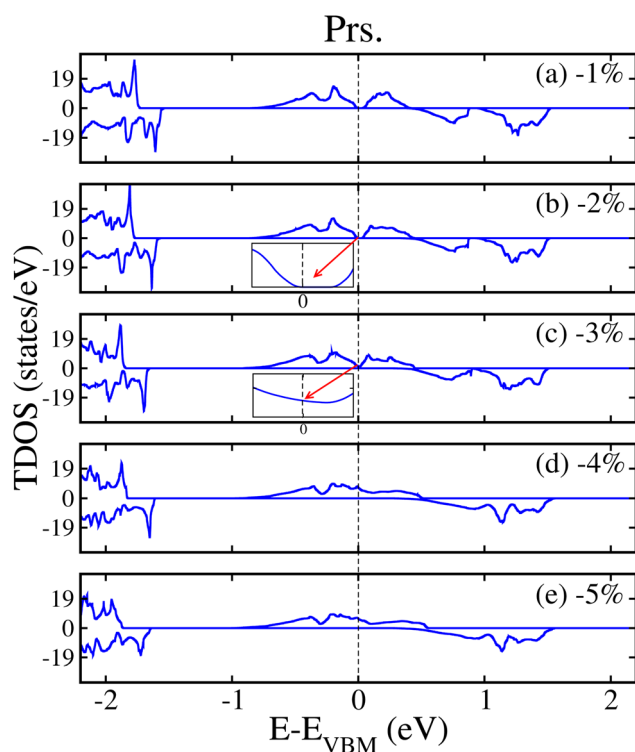


Fig. 11 GGA+U computed non-degenerate total density of states (TDOS) of the prs. $\text{Sr}_2\text{CaOsO}_6$ structure for (a) -1% , (b) -2% , (c) -3% , (d) -4% , and (e) -5% biaxial ([110]) compressive strain.

Next, we investigated the strain-induced changes in the systems' magnetism. To do this, the m_s on the Os/Ir ion in the prs. and Ir-dop. SCO motifs is plotted in Fig. 10S of the ESI.[†] As the strain is varied from -5% to $+5\%$, a slight change in the m_s value occurs in both systems. The m_s on the Os ion varies from $1.12\mu_B$ to $1.18\mu_B$ as strain changes from -5% to $+5\%$. Within a similar strain limit, the m_s on Os/Ir changes from 0.9 to $0.95\mu_B$ / -1.35 to $-1.32\mu_B$ in the Ir-dop. case. Along with this, we computed T_C under $\pm 5\%$ biaxial ([110]) strain, as presented in Fig. 12 for both systems. It shows that under -5% strain, prs./Ir-dop. has the highest T_C of $245/233$ K, while $+5\%$ strain gives the lowest T_C of $138/129$ K. Under comp./tens. strain, structural distortions increase/decrease, which has a positive/negative effect on the T_C values, as discussed in previous theoretical works.^{5,39}

Panneerselvam *et al.*⁶⁷ found that altering the polar optical phonon scattering mechanism *via* strain in ScN significantly influences the variance in lattice thermal conductivity while minimally affecting the TE power factor values. Likewise, Zhang *et al.*⁶⁸ examined the impact of strain on two-dimensional materials, emphasizing that strain engineering can improve TE capabilities by altering the electronic structure. Additionally, Yu *et al.*⁶⁹ showed that strain engineering may markedly influence the κ_1 and heat flow in Bi_2Te_3 nano-films, therefore highlighting the significance of strain in TE materials. The κ_1 of the Bi_2Te_3 nano-film may be efficiently adjusted by the application of strain. A tensile strain of 6% can decrease heat conductivity by 50% ; however, a compressive strain of 4% can enhance κ_1 by 60% . Hence, we plotted the ZT values of N^\uparrow/N^\downarrow at room temp. (300 K) as a function of $\pm 5\%$ strain in Fig. 13 for both systems. The total ZT is similar in the N^\uparrow/N^\downarrow for the prs./Ir-dop. system because it is closely related to the real E_g value, in contrast to Fig. 10 (see Fig. 4). Therefore, we compare the ZT values with the calculated E_g (see Fig. 10) for the strained system, which reveals that ZT is substantially correlated with E_g , where ZT increases as E_g increases and *vice versa*. The computed ZT under $+2/+3\%$ tens. strain is $0.68/0.71$ for the prs. and $0.04/0.15$ for the Ir-dop. system at 300 K. Therefore, we computed the TE factors of the prs. system under $+2\%$ and $+3\%$ biaxial tens. strains, as shown in Fig. 11S of the ESI.[†] As E_g increases under $+2/+3\%$ tens. strain, the values of $\frac{\sigma}{\tau}$, $\frac{\kappa_e}{\tau}$, and χ are less than those of the unstrained system because there is a smaller number of charge

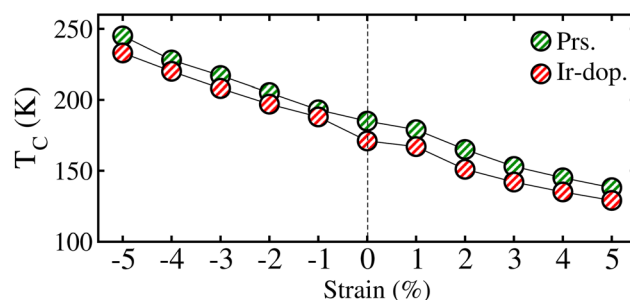


Fig. 12 Computed Curie temperature (T_C) under $\pm 5\%$ biaxial ([110]) strain for the prs. and Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures.



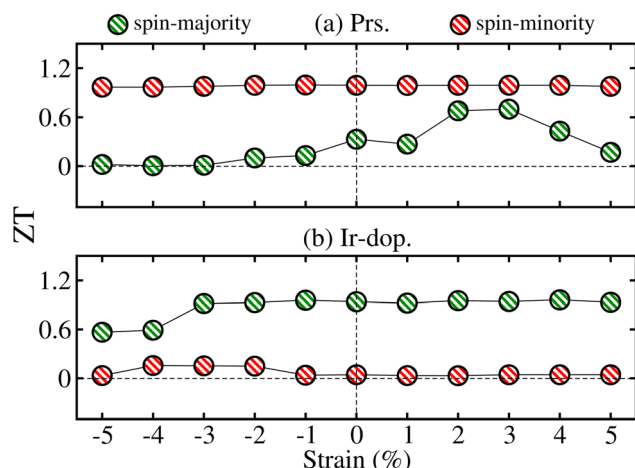


Fig. 13 Computed figure of merit (ZT) in the spin-majority and spin-minority channels for the (a) prs. and (b) Ir-dop. $\text{Sr}_2\text{CaOsO}_6$ structures as a function of $\pm 5\%$ biaxial ($[110]$) strain.

carriers across the E_F , but the trend remains the same with temp. as compared to the unstrained system. Also, the values of the PF at $+2/+3\%$ strain are less than that of the unstrained system, whereas the values of S also increase at $+2/+3\%$ tens. strain and it follows the same trend with temp. as followed in the unstrained system. Besides this, we calculated the $\frac{k_T}{\tau}$ of the prs. system under $+2/+3\%$ tens. strain, as presented in Fig. 12S of the ESI.† This shows that its value decreases with an increase in temperature, which is a required criterion for TE applications.

5 Conclusion

In summary, *ab initio* calculations were performed to examine the thermodynamic, mechanical, dynamical stability, thermoelectric, electronic structure, and magnetic properties of the unstrained and strained systems of the pristine (prs.)/Ir-doped (dop.) $\text{Sr}_2\text{CaOsO}_6$. It is predicted that both systems are thermodynamically and mechanically stable as they have negative formation energies and follow the Born criteria. Due to non-negative phonon frequency curves, they are dynamically stable as well. The unstrained prs./Ir-dop. system has a ferromagnetic (FM)/ferrimagnetic (FIM) ground state due to FM/antiferromagnetic (AFM) interactions between Os and Os/Os and Ir ions. The prs. motif is a semiconductor, while Ir-doping results in a half-metallic state (HM). The additional electrons produced by the dopant (Ir) cause a repulsion in the half-filled $\text{Os } t_{2g}$ in the spin-minority channel, which moves the Os bands towards the Fermi level, resulting in conductivity. Also, a large energy-gap of 1.15 eV in the non-metallic channel ensures the stability of the HM state of the system. Moreover, the computed ZT is 0.33/0.02 at 300 K for the prs./Ir-dop. motif. Further, our results indicate that the prs. system becomes half-metallic at a crucial value of -3% compressive strain. In contrast, the Ir-dop. motif retains its half-metallic nature against $\pm 5\%$ biaxial ($[110]$) strain. The prs./Ir-dop. structure has a high magneto-crystalline anisotropy energy (MAE) and

MAE constants (K) of 2.95/0.79 meV and $1.72/0.46 \times 10^7$ erg per cm^3 at the Curie temperature of 185/171 K, respectively. Interestingly, ZT values increase in the prs. system under $+2/+3\%$ tens. strain because the energy gap increases against these strain values. Hence, near-unity ZT values suggest that these materials are very suitable for magnetic memory devices and thermoelectric applications.

Data availability

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Author contributions

Samia Shahzadi: writing – original draft, investigations, formal analysis, data curation. Ihab Mohamed Moussa: validation, visualization, resources, formal analysis, funding. Sohail Mumtaz: visualization, validation, formal analysis. S. Nazir: writing – review and editing, validation, supervision, project administration, conceptualization.

Conflicts of interest

The authors declare no competing interests.

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