

Effects of chemical pressure on the magnetic ground states of the osmate double perovskites SrCaCoOsO₆ and Ca₂CoOsO₆

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The magnetic ground state in the double perovskite system Sr_{2-x}Ca_xCoOsO₆ changes from an antiferromagnet ($x = 0$), to a spin glass ($x = 1$), to a ferrimagnet ($x = 2$) as the Ca content increases. This crossover is driven by chemical pressure effects that control the relative strength of magnetic exchange interactions. The synthesis, crystal structure, and magnetism of SrCaCoOsO₆ and Ca₂CoOsO₆ are investigated and compared with Sr₂CoOsO₆. Both compounds adopt a monoclinic crystal structure with rock-salt ordering of Co²⁺ and Os⁶⁺ and $a^-a^-b^+$ octahedral tilting, but the average Co–O–Os bond angle evolves from 158.0(3)° in SrCaCoOsO₆ to 150.54(9)° in Ca₂CoOsO₆ as the smaller Ca²⁺ ion replaces Sr²⁺. While this change may seem minor, it has a profound effect on the magnetism, changing the magnetic ground state from antiferromagnetic in Sr₂CoOsO₆ ($T_{N1} = 108$ K, $T_{N2} = 70$ K), to a spin glass in SrCaCoOsO₆ ($T_{f1} = 32$ K, $T_{f2} = 13$ K), to ferrimagnetic in Ca₂CoOsO₆ ($T_C = 145$ K). In the first two compounds the observation of two transitions is consistent with weak coupling between the Co and Os sublattices.

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I. INTRODUCTION

Magnetically frustrated materials have received considerable attention over the years owing to the manifestation of exotic magnetic states such as spin liquids, spin ices, and spin glasses [1–3]. In particular, the double perovskite structure, A₂BB'O₆, which consists of rock-salt ordered BO₆ and B'O₆ corner-connected octahedra [4], has been a rich vein of magnetically frustrated materials. When the B cation is diamagnetic, the resulting B' sublattice possesses a geometrically frustrated fcc topology. The competition between nearest-neighbor and next-nearest-neighbor exchanges can lead to various patterns of antiferromagnetic (AFM) order [5], as well as more exotic magnetic ground states, such as a spin-singlet state (i.e., Ba₂YMoO₆), incommensurate magnetic order (i.e., La₂NaOsO₆), and an unconventional spin glass (i.e., Sr₂MgReO₆) [6–9]. When both B and B' are magnetic transition-metal cations, the resulting B–O–B' superexchange pathway must also be considered. When the B–O–B' superexchange is strong, as is the case for Sr₂CrOsO₆ ($T_C = 725$ K), high-temperature ferrimagnetism is observed [10]. When it is very weak, as is the case for Sr₂CoOsO₆, AFM ordering of weakly coupled B and B' sublattices results [11,12]. When the B–O–B' interaction is intermediate between these two extremes, highly frustrated magnetic ground states and incommensurate magnetic order are expected to emerge.

Recently, it has been shown that the insulator Sr₂FeOsO₆ undergoes two antiferromagnetic transitions resulting from a complex interplay of competing exchange interactions [13,14]. Interestingly, Ca₂FeOsO₆, which contains the same Fe³⁺ and Os⁵⁺ ions, has a ferrimagnetic ground state [15,16]. The change in magnetic ground state is thought to occur because the longer-range Fe–O–Os–O–Fe and Os–O–Fe–O–Os

interactions become weaker as the Fe–O–Os bond angles become increasingly bent, which allows the antiparallel Fe–O–Os superexchange interaction to gain the upper hand [16]. In this study we explore the effect of chemical pressure on the magnetism of Sr₂CoOsO₆, where long-range Co–O–Os–O–Co and Os–O–Co–O–Os interactions have been shown to drive magnetic ordering, while nearest-neighbor Co–O–Os interactions are particularly weak [11,12]. Our results, summarized in Fig. 1, show that through the application of chemical pressure via substitution of Ca for Sr, the structure is distorted, impacting the relative strengths of the superexchange interactions, which in turn converts the ground state from antiferromagnetic to ferrimagnetic through a spin-glass region where exchange competition is strongest.

II. EXPERIMENTAL

Powder samples of approximately 1.6 g of SrCaCoOsO₆ and Ca₂CoOsO₆ were synthesized by heating the reagents at 1000 °C for a period of 48 h in sealed silica tubes. PbO₂ was included separately as an *in situ* oxygen gas source as described in an earlier study [11]. Further synthetic details and precautions are elaborated in Ref. [17]. Phase purity of these products was established using laboratory x-ray diffraction. The high scattering contrast of Co and Os to x rays allowed us to confirm complete rock-salt ordering of Co and Os, while Sr and Ca are disordered on the A site in SrCaCoOsO₆. Neutron powder diffraction (NPD) measurements at 10 and 300 K for both materials were conducted on the POWGEN [18] beamline at the Spallation Neutron Source (SNS). Additional temperature-dependent neutron powder diffraction data were collected at 15-K intervals from 10 to 205 K on Ca₂CoOsO₆. Rietveld refinements were performed using the GSAS EXPGUI package [19,20].

dc magnetization measurements as a function of temperature were conducted utilizing a Quantum Design magnetic

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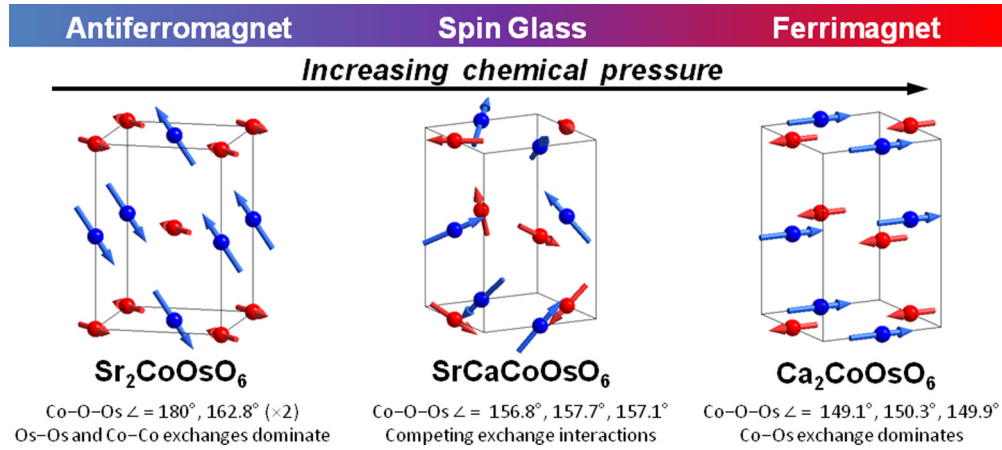


FIG. 1. (Color online) A schematic of the evolution of the magnetic ground state of $A_2\text{CoOsO}_6$ with chemical pressure due to the reduction of the Co–O–Os bond angle that accompanies increased octahedral tilting. Red and blue spheres correspond to Os and Co, respectively.

property measurement system (MPMS) superconducting quantum interference device (SQUID) magnetometer under zero-field-cooled and field-cooled conditions with a measuring field of 1 kOe for SrCaCoOsO₆ and 10 kOe for Ca₂CoOsO₆. dc magnetization measurements as a function of field were collected in a field range of ± 50 kOe at 5 and 300 K for Ca₂CoOsO₆. Similar data were collected for SrCaCoOsO₆ in a field range of ± 70 kOe at 2, 20, and 50 K. Heat capacity measurements were collected on a sintered pellet mounted with Apiezon grease using a Quantum Design physical property measurement system (PPMS) using the relaxation technique. ac magnetization studies at a variety of frequencies were conducted in the temperature range 3–60 K using a Quantum Design PPMS. dc magnetization relaxation measurements were conducted by cooling in an applied field of 10 kOe to specific temperatures before removing the field and collecting data as a function of time after field removal. X-ray magnetic circular dichroism (XMCD) measurements were conducted on APS beamlines 4-ID-C and 4-ID-D at the Co and Os $L_{3,2}$ edges at temperatures of 5 and 10 K, respectively, under fields of ± 35 kOe.

III. RESULTS AND DISCUSSION

Both materials crystallize with monoclinic $P2_1/n$ symmetry resulting from complete ordering of Co²⁺ and Os⁶⁺ ions and $a^-a^-c^+$ octahedral tilting [21], as defined using Glazer notation [22]. Refined room-temperature neutron powder diffraction patterns and depictions of the monoclinic crystal structure are given in Fig. 2. Select refined parameters and structural details at 10 and 300 K are listed in Table I. Further refined parameters, including variable temperature results, are available in Ref. [17]. No evidence for a structural phase transition was observed in either material over the temperature range examined. The average bond lengths for both CoO₆ and OsO₆ octahedra are comparable to those found in Sr₂CoOsO₆ [11], as well as other double perovskite oxides containing Os⁶⁺ or high-spin Co²⁺ [23,24], confirming that the electronic configuration of these cations does not change significantly upon substitution of Ca for Sr. The low-temperature Co–O–Os bond angles, which are dictated by the magnitude of octahedral tilting, take on average values of 157.2° in SrCaCoOsO₆ and

149.7° in Ca₂CoOsO₆. By comparison, the Co–O–Os bond angles in Sr₂CoOsO₆ are 180° in the c direction and 162.8° in the ab plane [11].

Figure 3(a) shows the temperature dependence of the dc magnetization of Ca₂CoOsO₆ as measured in a SQUID magnetometer in an applied field of 10 kOe. Ca₂CoOsO₆ exhibits a sharp rise in magnetization with a critical temperature of approximately 145 K. A Curie-Weiss fit was performed in the temperature range 350–400 K, resulting in an effective moment of 4.16 μ_B and Weiss constant of +5 K. However, it should be noted that ideal Curie-Weiss behavior may not be fully achieved within the temperature range measured (see Ref. [17]). The field dependence up to 50 kOe at 5 K produces a hysteresis loop with an M_{sat} of 1.77 μ_B per formula unit [see Fig. 3(b)]. For the electronic configurations of high-spin d^7 for Co²⁺ ($S = 3/2$) and d^2 for Os⁶⁺ ($S = 1$), a ferromagnetic M_{sat}

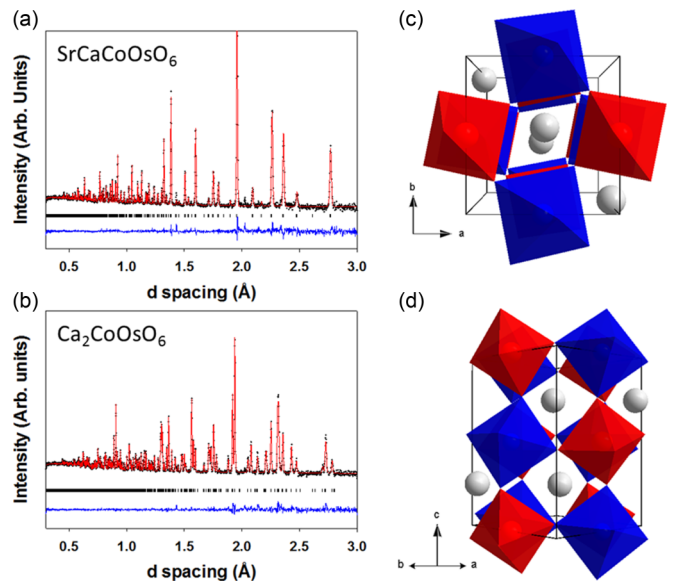


FIG. 2. (Color online) Refined room-temperature neutron powder diffraction patterns of (a) SrCaCoOsO₆ and (b) Ca₂CoOsO₆. The crystal structure of Ca₂CoOsO₆ is shown looking down the (c) 001 and (d) 110 directions to illustrate the $a^-a^-b^+$ pattern of octahedral tilting that leads to $P2_1/n$ symmetry.

TABLE I. Select structural features as determined from Rietveld refinements of neutron powder diffraction data. The \angle symbol represents a bond angle.

Temperature (K)	$\text{Ca}_2\text{CoOsO}_6$		SrCaCoOsO_6	
	10	300	10	300
Space group	$P2_1/n$	$P2_1/n$	$P2_1/n$	$P2_1/n$
$a(\text{\AA})$	5.3753(1)	5.39236(9)	5.5211(2)	5.5219(3)
$b(\text{\AA})$	5.5683(1)	5.55632(9)	5.5393(2)	5.5410(2)
$c(\text{\AA})$	7.6362(2)	7.6632(13)	7.7942(3)	7.8125(3)
β	89.755(1)	89.831(1)	89.926(6)	89.917(5)
$R_{wp}(\%)$	2.90	2.08	2.97	3.06
Co–O1 ($\times 2$)	2.080(2)	2.076(1)	2.069(3)	2.061(4)
Co–O2 ($\times 2$)	2.072(2)	2.063(2)	2.053(3)	2.051(4)
Co–O3 ($\times 2$)	2.032(2)	2.031(2)	2.049(4)	2.063(5)
Os–O1 ($\times 2$)	1.935(1)	1.934(2)	1.919(3)	1.926(3)
Os–O2 ($\times 2$)	1.932(2)	1.936(2)	1.929(3)	1.927(5)
Os–O3 ($\times 2$)	1.922(2)	1.929(2)	1.928(4)	1.921(4)
$\angle\text{Co–O1–Os}$	149.07(9)	149.86(9)	156.8(3)	157.7(3)
$\angle\text{Co–O2–Os}$	150.29(9)	150.97(9)	157.7(3)	158.9(3)
$\angle\text{Co–O3–Os}$	149.86(8)	150.78(8)	157.1(2)	157.4(3)

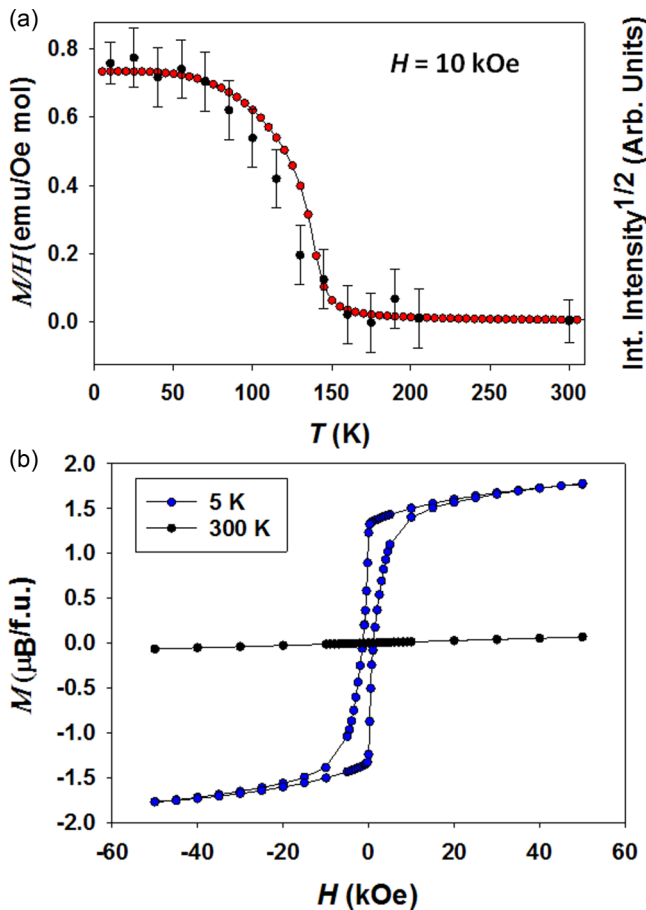


FIG. 3. (Color online) (a) The field-cooled temperature dependence of the dc magnetization of $\text{Ca}_2\text{CoOsO}_6$ measured at 10 kOe (red circles, left axis) and the square root of the total integrated intensity of the $(\bar{1}01)$, (110) , and (011) reflections, which increase due to magnetic scattering below the ordering temperature (black circles, right axis). (b) The field-dependent magnetization of $\text{Ca}_2\text{CoOsO}_6$ measured at 5 and 300 K.

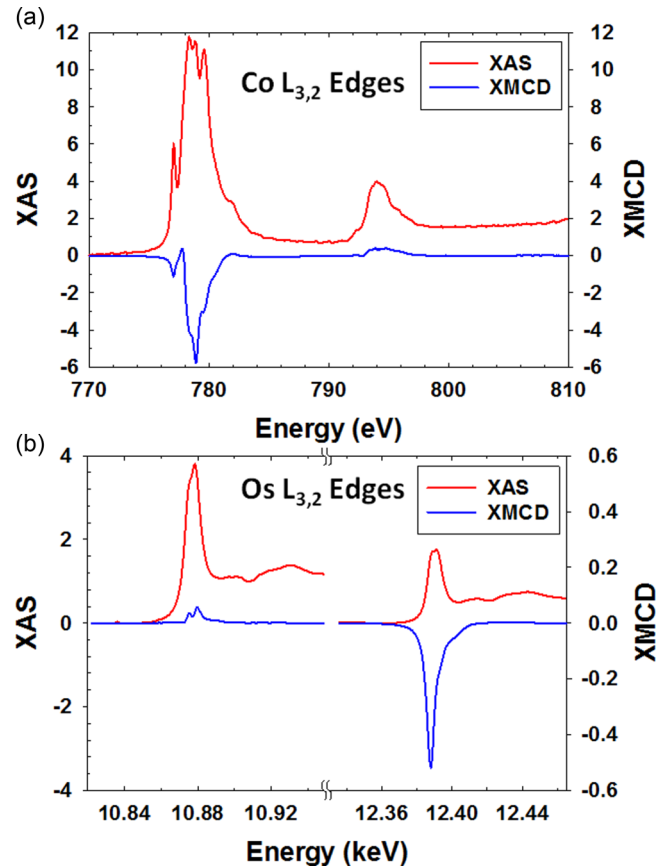


FIG. 4. (Color online) XAS and XMCD data for $\text{Ca}_2\text{CoOsO}_6$ measured at the (a) Co and (b) Os $L_{3,2}$ edges.

should be in the vicinity of $5 \mu_B$ per formula unit, assuming spin-only moments. On the other hand, an M_{sat} of only $1 \mu_B$ per formula unit is expected for a ferrimagnetic ground state. The observation of an experimental M_{sat} larger than the expected spin-only value can be attributed to the presence of spin-orbit coupling that increases the Co^{2+} moment and decreases the Os^{6+} moment. As discussed later, noncollinear magnetic configurations which might also lead to an increase in M_{sat} are ruled out by the absence of new Bragg peaks in neutron powder diffraction patterns collected below the Curie temperature.

The relative sign of the XMCD signal, shown as in Fig. 4, is reversed between Co and Os, indicating that the moments localized on these elements are oriented antiparallel to one another. Application of the sum rules yields a negative moment on Os and a positive moment on Co, meaning that upon application of a field, the Co moment aligns parallel to the field direction while Os moments align antiparallel to the external magnetic field. This is consistent with a ferrimagnetic configuration, as suggested by the previously described magnetization data. Neglecting T_z , the magnetic dipole operator, the calculated ratio of the orbital moment to the spin moment, m_l/m_s , is +44% and -28.5% for Co and Os, respectively, consistent with expectations for positive and negative orbital contributions for the high-spin d^7 and d^2 electronic configurations. The spin and orbital moment contributions on Os calculated from the sum rules [25]

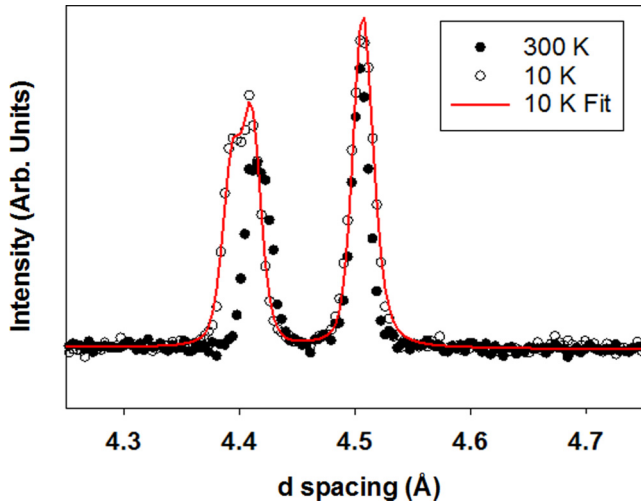


FIG. 5. (Color online) High d -spacing reflections of $\text{Ca}_2\text{CoOsO}_6$ at 300 K (filled circles) and 10 K (open circles) showing the increased intensity due to magnetic scattering, with fit to 10-K data shown as the red curve. The larger increase in intensity of the 101 as compared to the 011 peak is due to the orientation of magnetic moments along the b direction.

are $1.97(17)$ and $-0.56(4) \mu_B$, respectively, resulting in a substantial Os moment, in sharp contrast to the results from NPD refinements. The spin and orbital moments found in this way on Co are $1.8 \mu_B$ and $0.8 \mu_B$, respectively. Additionally, the x-ray absorption spectroscopy (XAS) spectrum of the Co $L_{3,2}$ edges is consistent with the assignment of high-spin Co^{2+} [26,27].

Neutron diffraction measurements of $\text{Ca}_2\text{CoOsO}_6$ taken at intervals of 15 K up to 205 K confirm the magnetic ordering. Plotted against the right axis of Fig. 3(a) is the integrated intensity of the $(\bar{1}01)$, (101) , and (011) reflections, which contain the strongest magnetic contribution. The intensity of these reflections is clearly rising as the sample is cooled below the Curie temperature. Figure 5 shows these peaks at 10 and 300 K for comparison to highlight the increased intensity from the magnetic contribution. No additional purely magnetic reflections were detected at low temperatures, excluding more complicated canted magnetic structure models. Magnetic refinements were conducted using the magnetic form factor coefficients for Os^{6+} as given by Kobayashi *et al.* [28]. Due to the high degree of correlation between the Co and Os moments, it was not possible to meaningfully refine the Co and Os moments independently. If one refines the data with only ferromagnetically aligned spins on cobalt (see Fig. 5), a moment of $2.33(8) \mu_B$ oriented parallel to the b axis is obtained. While this value is not unreasonable for a high-spin Co^{2+} ion, the lack of a moment on Os is inconsistent with the XMCD data and the total moment is not in agreement with the measured value of M_{sat} ($1.77 \mu_B$ per formula unit).

To probe the sensitivity of the NPD data to different combinations of Co and Os moments, a series of refinements were performed using fixed Os moments. The results show that fits of comparable quality can be obtained with relatively small Os moments aligned either parallel or antiparallel to a more substantial Co moment (see [17]). Discarding solutions

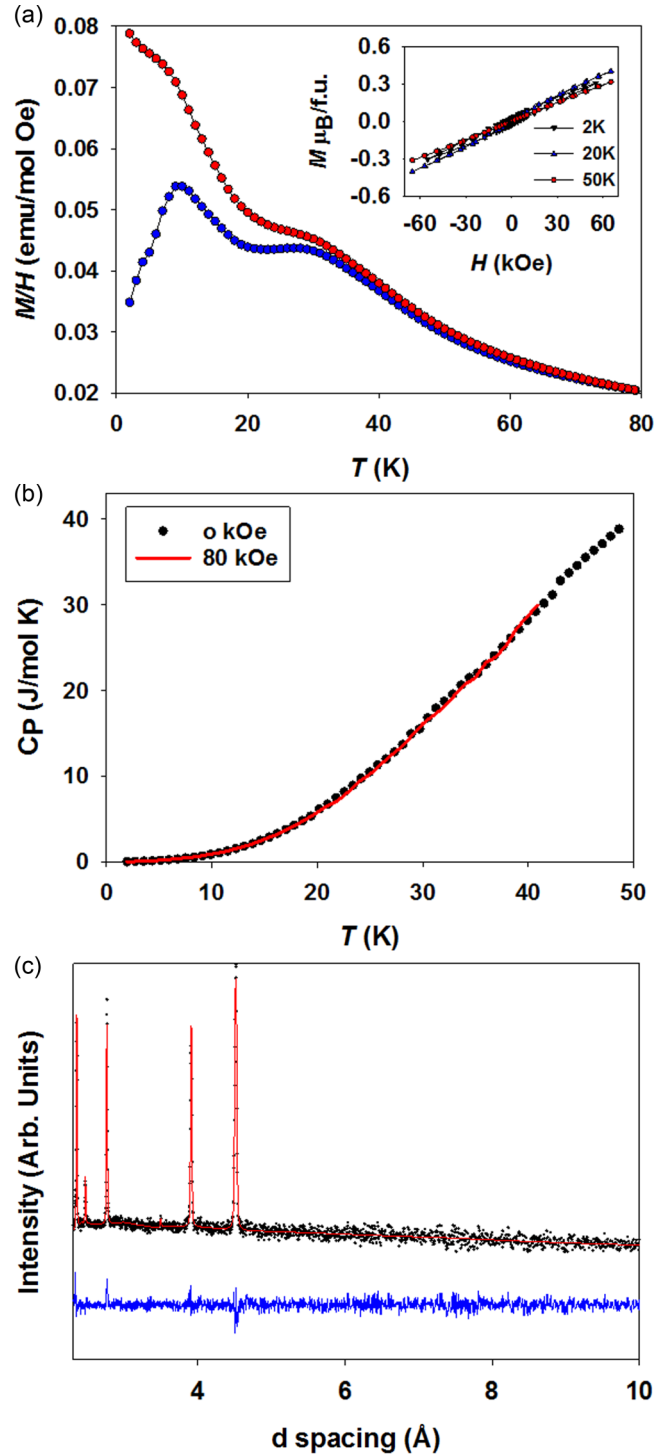


FIG. 6. (Color online) (a) The field-cooled (red) and zero-field-cooled (blue) temperature dependence of the dc magnetization of SrCaCoOsO_6 measured in 1 kOe, with field dependence at 2, 20, and 50 K shown in the inset; (b) heat capacity in 0- (black symbols) and 80-kOe (red line) fields; and (c) the high d -spacing region of the neutron powder diffraction pattern at 10 K (black dots). The agreement of the fit (red curve) to the data obtained using a nuclear-only model signals a lack of long-range magnetic order.

where the Co and Os moments are aligned parallel to each other as well as those where the Co moment falls below $2 \mu_B$ suggests that the Os moment lies somewhere between 0 and

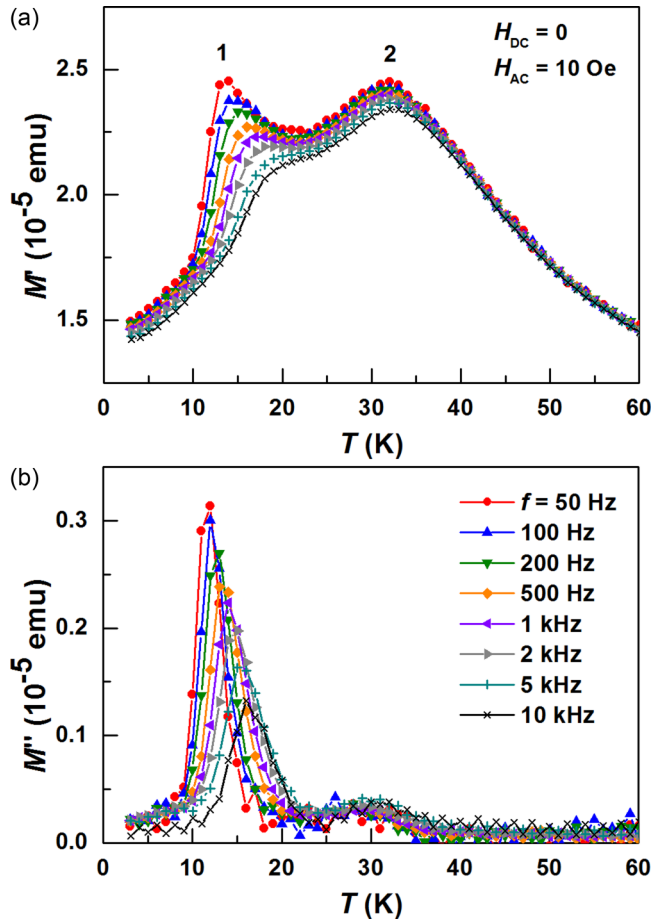


FIG. 7. (Color online) Results from frequency-dependent magnetization measurements on SrCaCoOsO₆. The real (M') and imaginary (M'') parts of the ac magnetization are shown in (a) and (b), respectively. Measurements were performed in zero dc field, with an ac field of 10 Oe.

$0.4 \mu_B$. The reasons why the Os moment estimated from the neutron refinements is significantly smaller than the moment estimated from XMCD is unclear.

The dc magnetization of SrCaCoOsO₆ reveals two cusps, each with a field-cooled-zero-field-cooled divergence at 10 and 27 K [see Fig. 6(a)]. The Curie-Weiss fit between 200 and 350 K (see [17]) resulted in an effective moment of $5.51 \mu_B$ and Weiss constant of -146 K. The effective moment is somewhat higher than the calculated spin-only value of $4.80 \mu_B$; however, in light of the XMCD measurements on Ca₂CoOsO₆, this increase in effective moment is likely due to a positive orbital contribution that increases the cobalt moment more than the negative orbital contribution decreases the osmium moment [23]. No discernible anomalies are apparent in the heat capacity, with or without an applied field of 80 kOe, as measured by a PPMS and shown in Fig. 6(b), nor is any magnetic scattering due to long-range ordering apparent in the 10-K neutron powder diffraction pattern shown in Fig. 6(c).

As the dc magnetization and heat capacity results are indicative of a glassy transition, ac magnetization measurements were performed as a function of temperature and frequency with an applied dc field of 0 Oe and ac field

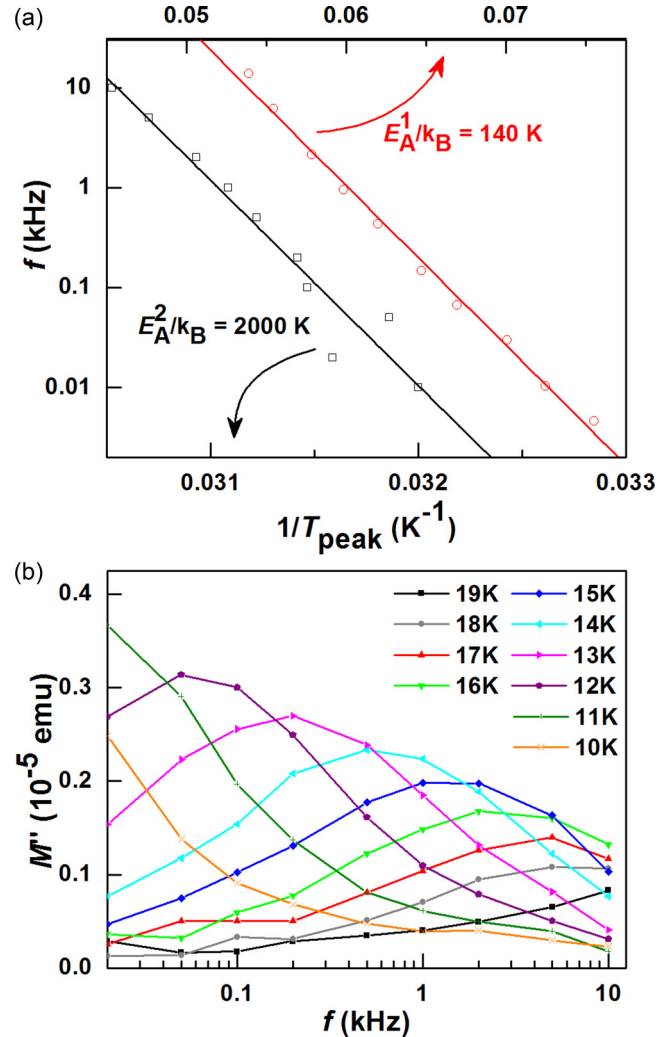


FIG. 8. (Color online) (a) Fits as described in the text to the temperatures T_{peak} at which the peaks in M' are observed in the ac magnetization data of SrCaCoOsO₆ as a function of temperature. (b) Frequency dependence of M'' for the lower temperature transition.

of 10 Oe. The resulting real and imaginary components, displayed in Figs. 7(a) and 7(b), respectively, show a strong frequency dependence at the lower temperature transition, $T_{f1} = 13$ K, as well as a weaker frequency dependence at the higher temperature transition, $T_{f2} = 32$ K. The frequency- and temperature-dependent behaviors observed here support the identification of both peaks observed in the dc magnetization measurements as classical spin-glass freezing events [3,29]. Activation energies (E_A) associated with these transitions were determined by modeling the relationship between the temperature at which the real part of the magnetization M' peaks (T_{peak}) and the frequency (f) using $f(T_{\text{peak}}) = f_0 \exp(-E_A/k_B T)$, where k_B is the Boltzmann constant. The results are shown in Fig. 8(a). An order of magnitude larger activation energy is observed for the higher temperature transition, indicating that the glassy state existing between T_{f1} and T_{f2} is much more “rigid” than the one formed upon cooling below T_{f1} , which is also consistent with the significantly smaller dissipation observed at T_{f2} as measured by the imaginary part of the magnetization M'' [Fig. 7(b)]. The frequency dependence and

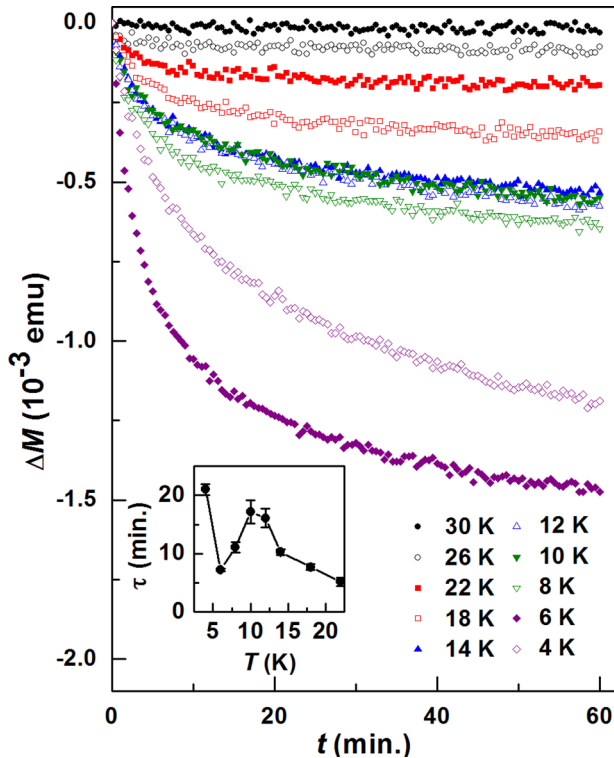


FIG. 9. (Color online) Time dependence of the dc magnetic moment of SrCaCoOsO₆ measured after cooling from 50 K in a magnetic field of 10 kOe and then reducing the field to zero. The inset contains the time constant determined from fits to the data.

the magnitude of M'' at T_{f1} is sufficiently strong to allow an examination of how the characteristic relaxation time varies with frequency [30], as shown in Fig. 8(b). This plot shows a single, characteristic peak in frequency which varies smoothly with temperature around T_{f1} .

Since spin-glass behavior is confirmed by the ac magnetic measurement, the time dependence of the dc magnetization was examined. Magnetic moment measurements were made after cooling from 50 K to the measurement temperature in a magnetic field of 10 kOe and then reducing the field to zero. The results, shown in Fig. 9, were fit with a stretched exponential of the form $M \sim \exp[-(t/\tau)^b]$, where τ is the time constant associated with the relaxation of the magnetic moment [3,29]. In general, it is expected that the time constant will increase upon cooling deeper into the spin-glass state. The temperature dependence of the fitted time constants is shown in the inset of Fig. 9. A sharp decrease in τ is observed upon cooling below 10 K, indicating the onset of

a second glassy state with faster relaxation rates, consistent with the ac results discussed above. Thus, the magnetic data indicate the formation of two independent spin-glass states in SrCaCoOsO₆. In the context of what is known about the properties of Sr₂CoOsO₆ [11], one can speculate that they correspond to independent freezing of the Co and Os sublattices.

IV. CONCLUSIONS

As was the case for the Sr₂FeOsO₆ system [15,16], the competing magnetic interactions are extremely sensitive to changes in the Co–O–Os bond angles in response to substitution of Ca for Sr. As the chemical pressure is increased and the bond angles are bent, the Co–O–Os–O–Co and Os–O–Co–O–Os exchange pathway strengths are weakened in favor of Co–O–Os superexchange. In the fully Ca-substituted end member, Ca₂CoOsO₆, the competition between the aforementioned long-range bond exchange pathways and the shorter AFM Co–O–Os exchange is dominated by the latter, resulting in ferrimagnetic order at the moderate T_C of 145 K. In SrCaCoOsO₆, where the octahedral tilting is intermediate, the relative strengths of these exchanges are comparable and spin-glass states emerge. Interestingly, it appears that the Co–O–Os superexchange pathway remains weak enough to allow the Co and Os sublattices to freeze independently of one another. It is remarkable that these subtle changes in bond angle can produce such dramatic changes in the magnetic ground state of these and related mixed $3d/5d$ transition-metal oxides. Further theoretical and experimental study of these systems is likely to yield additional unusual magnetic phenomena and is needed to increase our understanding of the competing superexchange interactions that are responsible for this rich magnetic phase space.

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