Strongly Coupled Magnetic and Electronic Transitions in Multivalent Strontium Cobaltites

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The topotactic phase transition in SrCoOₓ (x = 2.5–3.0) makes it possible to reversibly transit between the two distinct phases, i.e. the brownmillerite SrCoO₂.5 that is a room-temperature antiferromagnetic insulator (AFM-I) and the perovskite SrCoO₃ that is a ferromagnetic metal (FM-M), owing to their multiple valence states. For the intermediate x values, the two distinct phases are expected to strongly compete with each other. With oxidation of SrCoO₂.5, however, it has been conjectured that the magnetic transition is decoupled to the electronic phase transition, i.e., the AFM-to-FM transition occurs before the insulator-to-metal transition (IMT), which is still controversial. Here, we bridge the gap between the two-phase transitions by density-functional theory calculations combined with optical spectroscopy. We confirm that the IMT actually occurs concomitantly with the FM transition near the oxygen content x = 2.75. Strong charge-spin coupling drives the concurrent IMT and AFM-to-FM transition, which fosters the near room-T magnetic transition characteristic. Ultimately, our study demonstrates that SrCoOₓ is an intriguingly rare candidate for inducing coupled magnetic and electronic transition via fast and reversible redox reactions.

Coupling between magnetism and charge (or electricity) has triggered many fascinating phenomena, including colossal magnetoresistance1–3 and magnetoelectric effect4,5. It has been known that the strong competition between the antiferromagnetic super-exchange mechanism6 and the Zener double-exchange mechanism7 triggered by both doping and epitaxial strain brings out intriguing magnetoelectric phenomena and subsequent exotic phases.

Oxygen stoichiometry in the transition metal oxides (TMOs) plays an essential role in determining the physical properties, including optoelectronic and magnetic properties8–10. The multivalent nature of most transition metals often causes the formation of various solid TMO phases with different oxidation states, yielding intriguing oxygen concentration-dependent electronic and magnetic phase diagrams11–13. The oxygen content and consequent valence state of transition metals are also closely related to the ionic conduction and catalytic activities, which are critical in most cutting-edge energy storage and generation devices14–17. Therefore, exploring the role of oxygen defects in determining the electronic and magnetic properties would provide insight into identifying new possibilities for TMOs as energy materials.

Among TMOs, SrCoOₓ (SCO, 2.5 ≤ x ≤ 3.0) is an excellent candidate for studying the oxygen-content driven coupled phase transitions. Unlike LaCoOₓ, which shows good oxygen stability due to the robust Co³⁺ valence state18, SCO undergoes oxygen-content-dependent topotactic phase transitions. The latter accompany gigantic modifications in both its electronic and magnetic structures owing to the low oxygen sublattice stability12,19–21. It is worth mentioning that the brownmillerite SCO (x = 2.5, BM-SCO) exhibits a G-type AFM insulating phase with a high Neel temperature (TN = 540 K) and, the perovskite SCO (x = 3.0, P-SCO), which is a ferromagnetic metal, exhibits one of the highest Curie temperatures (TC = 305 K) among 3d transition metal oxides. However, due to the difficulty in synthesizing single phase perovskites, the coupling of the magnetic transition to the

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Received: 19 May 2017
Accepted: 13 October 2017
Published: xx xx xxxx
insulator-to-metal transition (IMT) and its abruptness have not been much explored, whereas a lot is known for other TMOs such as manganites. While the AFM in BM-SCO originates from superexchange in the insulating phase, the ferromagnetism in P-SCO originates from a distinct mechanism with itinerant holes mediating local Co spins. A study using neutron and x-ray scattering reported an abrupt transition in charge status from Co$^{3+}$ to Co$^{4+}$ in SCO at $x = x_0 = 2.82$ and in structure from orthorhombic to cubic at $x = x_1 = 2.75$. Therefore, understanding the microscopic picture of how the extreme phases (AFM-I vs FM-M) can be transformed via the reversible redox reactions is important both in fundamentals and new potentials of these materials.

Here, we present a combined study of density-functional theory (DFT) calculations and optical spectroscopy to reveal the coupling between itinerancy of charge carriers and magnetic transition mediated by the oxygen concentration in SCO. The DFT results predict that charge-spin coupling is sufficiently strong that the IMT is triggered by the magnetic transition. It is also found that the Drude peak in optical spectroscopy (indication of metallicity) occurs concomitantly with the formation of strong sigma bonding, facilitating the itinerant hole conduction and the stabilization of consequent ferromagnetism. Thus, we conclude that the strongly coupled nature may be responsible for the abrupt transition from AFM to FM.

**Result**

**IMT with oxidation revealed by optical conductivity.** Figure 1 shows oxygen content ($x$) dependent optical conductivity spectra ($\sigma_1(\omega)$) computed using DFT calculations (Fig. 1(a)) and recorded by spectroscopic ellipsometry (Fig. 1(b)). The overall qualitative absorption features ($\alpha$, $\beta$, $\gamma$, and $\delta$ peaks) are consistently revealed in both theoretical and experimental data. BM-SCO (see the spectrum for $x = 2.5$) shows an insulating behavior.
with two optical absorption peaks, \( \alpha \) and \( \beta \), representing Mott and charge-transfer gaps, respectively. Among
them, \( \beta \) is the dominant absorption with a greater intensity. As shown in Fig. 2(a), the \( \beta \) peak shows possible transitions from oxygen 2\( p \) to Co 3\( d \) (Mott) and \( p-d \) (charge-transfer) excitations, respectively. A large charge-transfer excitation, \( \beta \), occurs due to the orbital mixing in BM, while the excitation split into \( \gamma \) and \( \delta \) due to the separation between \( \sigma \) and \( \pi \) of oxygen 2\( p \) orbitals. \( p^\sigma \) and \( p^\pi \) denote \( \sigma \) and \( \pi \) bonding of oxygen \( p \) orbitals. In BM-SCO, the overall \( e_g \) and \( t_{2g} \) distributions are rather similar because of mutual mixing induced by vacancy-induced lattice distortions. The electronic distribution becomes clearly separated in P-SCO due to the higher structural symmetry.

**Figure 2.** Electronic structure for BM- and P-SCO. Computed PDOS (eV\(^{-1}\)) and electronic pictograms of (a) BM- (SrCoO\(_{2.5}\)) and (b) P-SCO (SrCoO\(_3\)). Blue and purple arrows denote \( d-d \) (Mott) and \( p-d \) (charge-transfer) excitations, respectively. A large charge-transfer excitation, \( \beta \), occurs due to the orbital mixing in BM, while the excitation split into \( \gamma \) and \( \delta \) due to the separation between \( \sigma \) and \( \pi \) of oxygen 2\( p \) orbitals. \( p^\sigma \) and \( p^\pi \) denote \( \sigma \) and \( \pi \) bonding of oxygen \( p \) orbitals. In BM-SCO, the overall \( e_g \) and \( t_{2g} \) distributions are rather similar because of mutual mixing induced by vacancy-induced lattice distortions. The electronic distribution becomes clearly separated in P-SCO due to the higher structural symmetry.
insulating $\beta$ peak into two major peaks ($\gamma$ and $\delta$) is strongly related to IMT with oxidation. The FM for highly oxidized SCO is induced by the itinerant hole conduction$^{22}$, as the formation of $\sigma$-bond and subsequent straight bonding character of Co-O-Co are advantageous to the itinerancy and FM.

**Possibility of magnetically-driven IMT.** In order to understand the main cause for IMT at $x = 2.75$, we comparatively calculated $\sigma_e(\omega)$ of relaxed SrCoO$_{2.75}$ for both FM and AFM configurations. The AFM configuration clearly shows an insulating optical spectrum [(see marked $x = 2.75$ (AFM) in Fig. 1(a))] as for BM with the dominant $\beta$ peak, whereas the $\beta$ peak disappears with the emergence of Drude peak for FM configuration [(see marked $x = 2.75$ (FM) in Fig. 1(a)] resembling that of P-SCO. We confirm that the Drude peak begins to appear from $x = 2.75$ in PV'-SCO (2.75 $< x < 2.8$) as in Fig. 1(b). Note that the oxidation state was confirmed by comparing x-ray absorption spectroscopy data with known spectra from bulk. This comparison clearly shows that electronic property, including insulating/metal character, is strongly subject to magnetic ordering; and metallic characters should be induced by the emergence of FM ordering at $x \sim 2.75$.

In particular, we find the $\beta$ peak of our DFT calculations is about 1 eV higher than that of experiment. When we use smaller $U$, the difference gets smaller, which means the estimation of $p$-$d$ transition is affected by the choice of $U$.

Figure 2 shows PDOS data calculated to explore the origin of optical absorption peaks. BM-SCO with 1D oxygen vacancy channels forms a fairly distorted structure and, thus, gives rise to an orbital mixing of $\pi$ and $\sigma$ bonding between oxygen and Co (Fig. 2(a)), which yields the strong $\beta$ peak indicative of an insulating state. However, by filling the 1D oxygen vacancies with oxidation, the overall structure evolves into the cubic symmetry, and $\sigma$- and $\pi$-bonding are clearly separated. Consequently, $\sigma$ bonding hybridization character is enhanced, so that the bandwidth of Co $e_g$ spreads up to 4 eV as shown in Fig. 2(b). Therefore, a straighter and easier path for the hole mediation of Co spin is formed and induces Zener-type double exchange ferromagnetism. Distinct from other cubic perovskites (SrMoO$_3$, $M = V$, Cr, Mn, and Fe), SrCoO$_3$ is a unique FM, exhibiting a high Curie temperature (near room temperature), driven by the strong double exchange. The evolution from Co$^{3+}$ to Co$^{4+}$ lowers the electronic energy states, increasing the overlap with oxygen. This change may help sustain FM close to room temperature. Moreover, these coupled characters, i.e. the AFM super-exchange in insulator and the FM double exchange in metal, reinforce the possibility of the coupled magnetic IMT at around $x = 2.75$.

**Drastic transition.** To check the character of the possibly coupled magnetic IMT, we carried out further DFT calculations and compared with experimental results as shown in Fig. 3. We used different compositions of SrCoO$_x$ ($x = 2.5, 2.75, 2.875,$ and 3.0) and in particular searched for different vacancy configurations for SrCoO$_{2.75}$ and SrCoO$_{2.875}$ to find the lowest energy structure. For every oxidation phase with the lowest energy, we calculated the energy difference between the G-AFM and FM relaxed phases as shown in Fig. 3(a). The energy difference shows an S-shape with a rather rapid increase around the transition ($x = 2.75$). It is worth mentioning that this rapidly increasing tendency from $x = 2.75$ does not change with a reasonable choice of $U$ as we have tested, for instance, with $U_{\text{eff}} = 2.5$ eV. Moreover, as summarized in Fig. 3(b), we see the AFM-I phase is transformed to the FM-M phase at around 2.75. Therefore, a small modification in oxidation near $x = 2.75$ can induce a large change in electronic and magnetic properties. This transition can be further supported by a sudden increase in the $\delta$ peak upon oxidation (see Fig. 1), which is responsible for the increase in sigma bonding. Potze et al.$^{22}$ proposed that an increase in sigma bonding can induce the double exchange interaction, resulting in enhanced FM$^{24}$. Consistently, we previously observed coexistence of the two phases, i.e. BM-SCO and PV-SCO, with an increased FM response upon oxidation without any intermediate phase$^{25}$. As shown in Table 1, increase in bond-angle of Co-O-Co induces formation of the straight $\sigma$-bond that drives the concurrent metallicity.

We further compared the calculated magnetic moments with the experimental values. The change in experimental magnetic moments shown in Fig. 3(b) is in good agreement with the theoretical results, showing a rather rapid increase at around $x = 2.75$. A magnetic transition is known to occur at around $x = 2.75$ and MIT at $x = 2.9^{24}$. Figure 3(c) and (d) show stable vacancy positions at $x = 2.5$ and $x = 2.75$. Based on the calculated structures, we conjecture that oxidation from $x = 2.5$ to $x = 2.75$ fills V1 in $z = 0.25$ plane and V2 in $z = 0.75$ plane, the most distant vacancies in the BM unit-cell. The simultaneous filling of the distant oxygen vacancy sites circumvents vacancy clustering so is advantageous to fast and effective oxidation. The measured magnetic moments of Co1 and Co3 sites of AFM BM phase are about 3.1 $\mu_B$/Co and 2.9 $\mu_B$/Co (Fig. 9 of ref.$^{25}$), respectively. Our calculated moments are 2.9 $\mu_B$/Co and 2.8 $\mu_B$/Co, respectively.

**Magnetically-driven metallic state.** Figure 4 shows total DOS for BM and two intermediates ($x = 2.75$ and 2.875) close to the transition. We computed for both AFM and FM phases to check the coupling of magnetism and metallicity. In SrCoO$_x$ ($x = 2.5 - 2.875$), the metallic character always appears with FM ordering at the intermediate phases, whereas they are insulating with AFM at 2.5 and 2.75. The proposed double exchange interaction$^{26}$ can be facilitated with the structure starting at $x = 2.75$ with oxidation$^{21}$. In Fig. 4(d), we show a schematic of the phase diagram from our optical measurements and DFT results compared with the previous $dc$-measurements$^{24}$. Although our combined optical study with DFT calculations shows a coupled magnetic IMT at $x = 2.75$, the $dc$ transport measurement shows the IMT at around $x = 2.9$. Thus, this offset of the IMT could be associated with extrinsic effects, such as phase separation by forming puddles of conducting charges, domain boundaries, etc., which can reduce electrical conductivity yielding poor electronic conduction in $dc$-measurements. Similar discrepancy between the optical study and $dc$-measurement was reported in colloossal magnetoresistive (CMR) (La,Sr)MnO$_3$.$^{25}$
Conclusions

In summary, we provide experimental and computational evidence that FM transition induces concurrent IMT in SrCoO\(_x\). First, the room-\(T\) AFM to high-\(T\) FM transition is a rare magnetic transition, so that such a distinct transition requires additional driving force such as IMT as happened in CMR. Also, a rather rapid change is found in magnetic energy and moment at around \(x = 2.75\), which may be affected by the IMT. Secondly, our DFT study reveals that formation of the straight \(\sigma\)-bond drives the concurrent metallicity. Thus, the orthorhombic to cubic transition at \(x = 2.75\) (ref.\(^{23}\)) forming straight bonds can drive the metallic state at the same composition. Lastly, our DFT results confirm that the metallic state is always accompanied by FM ordering regardless of the composition at \(x = 2.75\).
degree of oxidation in SrCoO$_{x}$, so the FM transition at $x = 2.75$ should trigger IMT. We further note that IMT in TMOs has been recognized as an important ingredient for many technological applications. We show that oxygen intercalation, which is known to occur as low as 200 °C in SrCoO, oxygen sponges can be a powerful means to induce IMT without complicated chemical doping of cations. This vacancy-induced, magnetically coupled IMT can be compared with pressure-induced, magnetically coupled IMT in rare-earth double perovskites. It is also worth noting that TMOs have many order parameters, including spin, charge, lattice, and orbital. Since the strong interactions among the order parameters can be delicately controlled by oxygen content, the fundamental understanding of the coupling between the magnetic and electronic properties reported here will bring a tremendous impact on tailoring diverse functionalities.

Methods

First-principles calculations. DFT calculations were performed within the generalized gradient approximation GGA $+ U$ method with the Perdew–Becke–Erzenhof parameterization as implemented in the Vienna ab initio Simulation Package (VASP-5.2). The Projector augmented wave (PAW) potentials include ten valence electrons for Sr ($4s^24p^6$), nine for Co ($3d^84s^1$), and six for oxygen ($2s^22p^4$). The wave functions are expanded in a plane wave basis with 500 eV energy cutoff. For BM-SCO, a $3 \times 1 \times 3$ Monkhorst-Pack $k$-point grid was used for relaxation and $6 \times 2 \times 6$ grid was used for optical property and density of states. For P-SCO, a $14 \times 14 \times 14$ grid was used. We use the Dudarev implementation with on-site Coulomb interaction $U = 4.5$ eV and on-site exchange interaction $J_H = 1.0$ eV to treat the localized $d$ electron states in Co with $U_{dd} = 3.5$ eV. A similar choice of $U$ was successfully applied to BM-SCO for analysis of its electronic and magnetic properties. To find the minimum energy configuration in the given compositions ($x = 2.5, 2.75$, and $2.875$), we compared total energies of different vacancy configuration and chose the lowest energy configuration where the two oxygen vacancies are most distant in unit-cell. The calculated lattice parameters are shown in Table 2.

Experimental intensity is lower than theoretical one because extrinsic effects in experimental conditions may reduce the optical conductivity. Also, positions of the frequency ($\omega$) values in the theoretical peak depend on the choice of $U$. Therefore, a smaller $U$ may shift lower the peaks and gives rise to better agreement with experimental results.
Table 2. Lattice information calculated from DFT. c-lattice parameters of calculated SrCoO$_x$ (x = 2.5, 2.75, 2.875, 3.0) epitaxially-strained on LSAT substrate (a = b = 3.87 Å).

| x     | AFM | FM |
|-------|-----|----|
| 2.5   | 5.746 | 5.804 |
| 2.75  | 5.627 | 5.616 |
| 2.875 | 5.591 | 5.585 |
| 3.0   | 5.527 | 5.584 |

Sample preparation and spectroscopic ellipsometry. We used pulsed laser epitaxy to grow epitaxially strained SCO thin films on (001) (LaAlO$_3$)$_{0.3}$(SrAl$_{0.5}$Ta$_{0.5}$O$_3$)$_{0.7}$ (LSAT) substrates. We kept the sample thickness as 55 Å. Different annealing conditions were applied to systematically change the oxidation state of the BM-SCO thin films. P-SCO was also directly grown in O$_2$ + O$_3$ (5%) environment. Detailed sample preparation and other properties can be found elsewhere$_{10}$. Spectroscopic ellipsometry was performed using an ellipsometer (M-2000, J. A. Woollam Co., Inc.) at 0.4 and 5.4 eV at an incident angle of ~60°, 70°, and 80°. Simple two-layer (film/substrate) model fit was used to successfully deduce the complex dielectric functions of thin films.

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(theoretical calculations). W.S.C., H.J., and H.N.L. were supported by the U.S. Department of Energy, Office of Science, Basic Energy Sciences, Materials Sciences and Engineering Division (synthesis and experimental characterization). W.S.C. was in part supported by Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Science, ICT and Future Planning (NRF-2017R1A2B4011083) (optical data analysis). Computation resources were supported by the Supercomputing Center/Korea Institute of Science and Technology Information with supercomputing resources including technical support (KSC-2017-C3-0018).

Author Contributions
J.H.L. proposed the possibility of the transition. H.J. fabricated the samples and provided basic concept of the phase transition and related phenomena. W.S.C. performed optical spectroscopic measurements. J.H.L., M.S.Y., H.-J.L. carried out first-principles DFT calculations. M.S.Y. supplied computational resources. J.H.S. and J.N. helped analyse the data. H.N.L. supervised all the experimental work. J.H.L., W.S.C., H.J., M.S.Y., and H.N.L. conceived the overall idea and participated in writing and revising the manuscript.

Additional Information
Competing Interests: The authors declare that they have no competing interests.

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