

Ligand field-induced exotic dopant for infrared transparent electrode: W in rutile SnO₂

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Abstract

Transparent conductive oxides (TCOs) exhibiting high near-infrared (NIR) transmittance are one of the key materials for highly efficient thin-film solar cells with widened spectral sensitivity. To realize excellent NIR transparency in a TCO film, developing a dopant providing high mobility (μ) carriers at moderate concentration is quite important. Recent studies have revealed that *d*-block transition metals are promising dopants to enhance μ in conventional *s*-orbital-based host semiconductors such as In_2O_3 and SnO_2 . In a transition metal-doped TCO, ligand field splitting of the energy levels of the donor *d* states can render the transition metal an unprecedented dopant for realizing high μ in TCOs while controlling the carrier density. Herein, we demonstrate that W is such an “exotic” dopant for rutile SnO_2 , exhibiting high μ and thus allowing NIR transparency. A combination of electrical transport property measurements and hybrid density functional theory calculations revealed that W, a group-6 element, behaves as a singly charged donor (W^{5+}) showing minimized ionized impurity scattering in tetravalent cation-based SnO_2 due to the splitting of the W 5d t_{2g} -states. This splitting was realized not only by the octahedral crystal field but also hybridization with the O 2*p* orbitals in the rutile-type crystal structure.

Main Text

Transparent conductive oxides (TCOs) exhibiting high visible and near-infrared (NIR) transparency are increasingly in demand as transparent electrodes for next-generation thin-film solar cells with widened spectral sensitivity.[1–8] A key requirement for a TCO film with excellent NIR transparency is high Hall mobility (μ) for carrier electrons with control of the carrier density in a moderate range, which suppresses free-carrier reflection.[9] In a high- μ TCO, an appropriate amount of a singly charged donor must be doped into a host oxide semiconductor to minimize mobility-reducing ionized impurity scattering.[10] The conventional strategy adopts a *p*-block dopant in a group adjacent to the cation of the host oxide semiconductor on the periodic table (e.g., Al- or Ga-doped ZnO, Sn-doped In₂O₃, and Sb-doped SnO₂).[11] Recent studies, however, revealed that *d*-block (i.e., transition metal) dopants are preferable to *p*-block ones to enhance μ in typical *s*-orbital-based TCOs such as In₂O₃ [12–18] and SnO₂ [19–21]: In these TCOs, donor *d* states possess energy levels sufficiently high relative to the conduction band minimum (CBM) that they hybridize minimally with the host cation *s* states of the CBM, avoiding an increase in the electron effective mass induced by *p*-block dopants due to the hybridization between the *s* states of the dopant and the host cation.

In a transition metal-doped TCO, splitting of donor *d* states by the ligand field also plays a crucial role in determining the charge state of the donor, which extends the choice of dopant for achieving high μ . Group-6 Mo in In₂O₃ is such an example; Mo would be expected to behave as a multiply charged donor in trivalent cation-based In₂O₃. In Mo-doped In₂O₃, however, Jahn–Teller (J-T)-like distortion of MoO₆ octahedra splits Mo 4d *t*_{2g} states into an upper *d*_{xy} state and lower *d*_{xz} and *d*_{yz} states. While the energy level of the *d*_{xy} state is far above the CBM and releases a carrier electron to the CBM, the *d*_{xz}

and d_{yz} states are in the band gap and trap two electrons with high-spin configuration.[17,18] As a result, the Mo dopant serves as a singly charged donor (Mo^{4+}), fulfilling the above-mentioned criteria for a high- μ TCO.

In this study, we demonstrate that W is such an “exotic” dopant for SnO_2 , allowing high μ and thus NIR transparency. The electrical transport properties of epitaxial thin films of W-doped SnO_2 (WTO) and hybrid density functional theory (DFT) calculations revealed that group-6 W works as a singly charged donor (W^{5+}) in tetravalent cation-based SnO_2 . This unexpected W^{5+} state originates from the splitting of the W 5d t_{2g} -orbitals not only through crystal field based on the point charge model but also through hybridization between the W 5d-orbitals and the O 2p orbitals.

W-doped SnO_2 as a NIR transparent conductor

It was reported that polycrystalline WTO films grown on an anatase TiO_2 seed layer showed high μ of $>80 \text{ cm}^2\text{V}^{-1}\text{s}^{-1}$, leading to high transparency in the wide wavelength region of 400-1950 nm.[21] In polycrystalline WTO films, however, complex factors such as grain boundaries hinder the quantitative analysis of electrical properties. Thus, we fabricated epitaxial thin films of WTO to clarify the mechanism behind its high μ .

Figure 1(a) shows ω - 2θ X-ray diffraction (XRD) patterns for the epitaxial WTO thin films (W amount $x \leq 4.0 \times 10^{-2}$), where only the 002 diffraction peaks from SnO_2 and TiO_2 were observed. The reciprocal space map around the 112 diffraction peak for the WTO film indicated epitaxial growth of an almost fully relaxed film on the substrate (inset of Fig. 1(a)). Although the lattice constant change with W doping was very small due to the low dopant concentration (Supplementary Figure S1), the substitution of W for

Sn was validated by X-ray fluorescence holography (XFH).[22,23] The local environment around W reconstructed by XFH was almost equivalent to that of the host Sn sites (Supplementary Figure S2). Dark-field scanning transmission electron microscopy (STEM) and corresponding energy dispersive X-ray spectroscopy (EDS) measurements of the WTO film (Fig. 1(b), $x = 1.4 \times 10^{-2}$) confirmed that W atoms were homogeneously distributed inside the WTO thin films without any segregation or interdiffusion at the film/substrate interface.

The WTO epitaxial thin films showed excellent optical transparency and electrical conductivity. Figure 2(a) shows optical transmittance (T) and reflectance (R) spectra of WTO films with various values of x . Synchronized with the decrease in the sheet resistance induced by W doping (Fig. 2(b)), the IR transmittance monotonically decreased due to enhanced free carrier reflection. Nevertheless, high T was maintained in the wide wavelength region from ultraviolet to NIR (300–2000 nm) even for the WTO film with the lowest sheet resistance (13.0 Ω /sq. at $x = 1.4 \times 10^{-2}$), as reported for polycrystalline WTO films,[21] demonstrating the potential of WTO films as NIR transparent conductors.

Charge state of the W in WTO

According to the above-mentioned criteria for a high μ TCO, we speculate that the W ions in the WTO films exist as singly charged species (W^{5+}), for which ionized impurity scattering is weaker than that of doubly charged species (W^{6+}). Indeed, the carrier density (n_e) of the WTO films was close to the ideal values assuming 100% activation of W^{5+} (i.e., each W generated one electron) for $x \leq \sim 0.01$. In the WTO film with larger x , n_e saturated at $\sim 2.1 \times 10^{20} \text{ cm}^{-3}$ (Figure 3(a)) probably due to the formation of neutral W species (W^{4+}) as discussed later. Detailed analysis of μ as a function of n_e

also indicates the presence of the singly charged W^{5+} state, as described below. Figure 3(b) compares the experimental μ values of the WTO films measured at room temperature with theoretical ones calculated as $\mu_{\text{cal}} = (\mu_{\text{lat}}^{-1} + \mu_{\text{iis}}^{-1})^{-1}$ for both W^{5+} and W^{6+} states,[20] where μ_{lat} is the lattice mobility associated with phonon scattering [24] and μ_{iis} is the Hall mobility limited by ionized impurity scattering based on the Brooks–Herring–Dingle (BHD) formula [25]. As seen from the figure, μ increased with increasing n_e owing to the enhanced screening of dislocations and/or grain boundaries and reached $136 \text{ cm}^2\text{V}^{-1}\text{s}^{-1}$ for the WTO film with $n_e \sim 2.1 \times 10^{20} \text{ cm}^{-3}$ ($x = 1.4 \times 10^{-2}$), which was ~ 1.6 times higher than that of the polycrystalline film ($84 \text{ cm}^2\text{V}^{-1}\text{s}^{-1}$).[21] Remarkably, this μ value evidently exceeds μ_{cal} for W^{6+} and is comparable to that for W^{5+} , indicating that W is doped as W^{5+} , not W^{6+} . This feature was further corroborated by low-temperature electrical measurements, where phonon scattering can be ignored ($\mu_{\text{cal}} = \mu_{\text{iis}}$) (Supplementary Figure S3). At 10 K, the μ values agreed well with μ_{iis} for W^{5+} , which supports the conclusion that W exists as W^{5+} .

Formation of singly charged W^{5+} in the WTO films was rationalized by DFT calculations. Figure 3(c) displays the thermodynamic transition level diagram under *O-poor* conditions, which describes the formation energies as a function of Fermi energy and thus provides intuitive information on defect concentrations for the tungsten species W_{Sn} and W_{i} as well as the dominant intrinsic defects, V_{Sn} and V_{O} . [26,27] This diagram clearly shows that (1) the dominant *n*-type defect in WTO is W_{Sn} across the range of chemical potentials, and (2) W_{Sn} acts as a one-electron donor (i.e., singly charged W^{5+}) if the Fermi energy is lower than the (+1/0) transition level occurring around 0.29 eV above the CBM. Notably, the energy of the (+1/0) transition level agrees well with the Burstein-Moss shift ($\sim 0.30 \text{ eV}$) of the WTO films with saturated n_e of $\sim 2.1 \times 10^{20} \text{ cm}^{-3}$

(Supplementary Figure S4), suggesting that an increase in the density of neutral impurity states limits the n_e in heavily W-doped thin films.

Impact of ligand field on the electronic structure of WTO

To understand more closely the role of W as a dopant in SnO₂, unfolded supercell band structures and density of states for the WTO were calculated (Fig. 4). Based on experimental evidence, substitutional W (W_{Sn}) in the +1 (ionized) charge state was considered. It was assumed that W possessed a high-spin state (with two electrons in the spin-up component), not a low-spin (with one electron in the spin-up and the other in the spin-down component) state because the high-spin state has ~0.2 eV lower formation energy than the low-spin state. In the spin-up bands (Fig. 4(a)), a filled band of W 5*d* character forms a mid-gap state around 1.63 eV above the valence band maximum (VBM), consistent with the experimental result that W was incorporated as W⁵⁺. Other clear W-related features can be seen in the localized bands of W 5*d* character starting around 1.39 eV above the CBM in both the spin-up and spin-down components, which release one electron into the CBM. Furthermore, these W-related bands showed negligible contribution at the CBM (Fig. 4(b)). This implies that the CBM effective mass is not perturbed through doping, resulting in very high mobility, as seen in other transition metal-doped high- μ TCOs such as Ta-doped SnO₂ [19] and Zr- or Mo-doped In₂O₃ [17,18]. A slightly smaller electron effective mass for the WTO films than that of *p*-block element (Sb)-doped SnO₂ was indeed confirmed by Drude fitting of the optical reflectance spectra (Fig. 2(c)).

The band structures of WTO mentioned above clearly indicated that the high μ stems from the splitting of W 5*d* states. In the case of Mo-doped In₂O₃, the splitting of

Mo $4d t_{2g}$ states is caused by the crystal field of J-T-like distorted MoO_6 octahedra as mentioned above.[17,18] However, the splitting of the W $5d$ states in WTO *cannot* be explained by the same scenario, because similar splitting of the W $5d$ states occurs even in an undistorted octahedral coordination (Supplementary Figure S5). The mechanism behind the splitting of W $5d$ states can be understood by considering hybridization between the W $5d$ -orbitals and the O $2p$ orbitals in addition to the octahedral crystal field. As illustrated in a molecular orbital diagram (Fig. 4(c)), the d states of a transition metal in the rutile structure near the Fermi level generally consist of two states originating from the t_{2g} orbitals in the octahedral crystal field [28]: One $d_{||}$ orbital directed along the c axis is almost non-bonding, while two d_{\perp} orbitals are hybridized with O $2p_{\pi}$ orbitals and form anti-bonding states. According to this diagram, the splitting of W $5d$ states in WTO can be rationalized as follows. The lowest energy W-related band (band 1 in Fig 4(a)) corresponds to $d_{||}$, and the higher energy bands (bands 2 and 3) to d_{\perp} . The partial charge densities of these bands (Fig. 4(d)) verified this assignment, where the non-bonding feature of the $d_{||}$ state (band 1) and the anti-bonding hybridization between the d_{\perp} and the O $2p_{\pi}$ orbitals (bands 2 and 3) are visualized.

These results indicated that not only the crystal field based on the point charge model but also hybridization with neighboring oxygen orbitals, which depends on the host crystal structure, should be considered to develop an exotic transition metal dopant for a high- μ TCO: For example, it is predicted that W_{Sn} would *not* be a high- μ dopant for Sn-based TCOs with the perovskite structure,[29] such as BaSnO_3 [30,31] and SrSnO_3 ,[32,33] because W $5d t_{2g}$ states show no further splitting due to their non-bonding nature in the perovskite lattice. As a result, doubly charged W^{6+} becomes the most stable state, which exhibits stronger ionized impurity scattering.

Conclusion

We have demonstrated that WTO is a high- μ TCO showing excellent transparency in a wide wavelength range from ultraviolet (300 nm) to NIR (2000 nm). Comprehensive analysis of electrical transport properties and DFT calculations revealed that a singly charged W^{5+} state with minimized ionized impurity scattering is stabilized by the splitting of $W 5d t_{2g}$ states in WTO. This splitting stems not only from the octahedral crystal field but also from hybridization with the O $2p$ orbitals surrounding the dopants, whose contribution has not been considered in a transition metal-doped TCO so far. Hybridization between the metal d orbital and O $2p$ orbitals is a well-established concept to understand the electronic structure of a transition metal oxide, and thus would provide a new guide for designing a novel high- μ dopant to obtain a TCO with NIR transparency.

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Author Contribution

Y. H. and S. N. conceived the project. M. F. fabricated the WTO films and characterized their optical and electrical transport properties. B. W. and D. Scanlon performed the computational calculations. M. F., K. K. and K. H. measured the XFH. F. M., Y. H., Y. S., and D. Sekiba performed the RBS measurements. F. M., Y. H., B. W. K.K. and T. H. wrote the manuscript with the inputs from all authors.

Methods

Sample preparation and characterization

Thin film deposition: (001)-oriented WTO films with thicknesses of 130–150 nm were grown on TiO₂ (001) and Al₂O₃ (10–10) (m-plane) by pulsed laser deposition (PLD) using a KrF excimer laser. Sintered pellets of WTO with nominal W/(Sn+W) ratio of $6 \times 10^{-4} - 5 \times 10^{-2}$ were used as PLD targets. The repetition rate and laser fluence were fixed at 2 Hz and $1-2 \text{ J}\cdot\text{cm}^{-2}$, respectively. The typical growth rate was 0.18–0.21 Å per shot. The base pressure of the growth chamber was maintained at 1×10^{-8} Torr. The oxygen partial pressure and substrate temperature during film growth were set at 1×10^{-2} Torr and 500 °C, respectively.

Structure and chemical composition: The crystal structures of the films were evaluated by XRD measurements using a four-circle diffractometer (Bruker AXS, D8 DISCOVER). The W composition of the WTO films, x in $\text{W}_x\text{Sn}_{1-x}\text{O}_2$, was evaluated by using energy dispersive X-ray spectroscopy coupled with scanning electron microscopy (SEM-EDX) (JEOL, JSM-7100F with JED-2300). The electron accelerating voltage was set at 5.0 kV to reduce the background signal from the substrate. The results of SEM-EDX measurements were calibrated by a calibration curve prepared based on the x values of a series of WTO films determined by Rutherford backscattering spectrometry (RBS). The RBS measurements were performed with a $^{35}\text{Cl}^{7+}$ beam accelerated to 38.4 MeV generated by a 5-MV tandem accelerator (Micro Analysis Laboratory, The University of Tokyo [MALT]).[34] The incident angle of the $^{35}\text{Cl}^{7+}$ beam was set as 75° from the surface normal and backscattered particles were collected at 150° from the beam incident direction. A W $L\alpha$ X-ray fluorescence (8.4 keV) hologram of the WTO (001) epitaxial film ($x = 1.4 \times 10^{-2}$) was measured at the BL13XU beamline of SPring-8 of the Japan

Synchrotron Radiation Research Institute. Details of the experimental setup have been reported elsewhere.[22] The ranges of the exit and azimuthal angles, θ and φ , were $\theta = 0-75^\circ$ and $\varphi = 0-360^\circ$, and the angles were varied in steps of 1° and 0.25° , respectively. The energy of the incident X-rays was set to 12.2 keV–15.7 keV with energy steps of 0.5 keV. The atomic image around W atoms was reconstructed from these holograms by using the Barton algorithm.[35] The cross-sectional microstructure and elemental maps of the films were observed by using a scanning transmission electron microscope (JEOL JEM-ARM200F, 200 kV) with an EDX detector (Thermo Fisher Scientific, NSS). The specimen for TEM observation was prepared by using a focused ion beam system (Hitachi High-Tech, XVision 200TB).

Optical and electrical transport properties: The optical transmittance T and reflectance R of the films were measured by using a UV/visible/near-infrared spectrophotometer (JASCO, V-670) and an FT-IR spectrometer (JASCO, FT/IR-4100). Electrical transport properties of the films were measured using the six-probe method with the standard Hall bar geometry (1 mm \times 2.4 mm). A laboratory-constructed system equipped with a 2 T electromagnet (at room temperature) and a commercially available system (Quantum Design, physical property measurement system (PPMS Model 6000), at low temperature) was used for the measurements. Ag or In electrodes were used for ohmic contacts. Current-voltage characteristics and Hall voltage-magnetic field characteristics were measured repeatedly (at least twice) to confirm the reliability and reproducibility of the measurements.

Computational methodology

Theoretical and modeling methods: Computational calculations were carried out using density functional theory (DFT) within the Vienna *ab-initio* Simulation Package (VASP).[36–39] The hybrid functional PBE0 formalized by Adamo and Barone [40] was used in order to provide an accurate description of the band gap and electronic properties relative to experiment as shown previously for SnO₂ and other tin-based oxides.[19,26,27,41–44] To describe the interaction between the core and valence electrons, the projector augmented wave (PAW) method [45] was employed. Tungsten defects were simulated in a $2 \times 2 \times 3$ (72 atoms) supercell expansion of a geometrically optimized unit cell of SnO₂. [27] Each defect supercell was subjected to structural relaxation to a force convergence criterion of 0.01 eV Å⁻¹ using a plane-wave energy cut-off of 400 eV and a Γ -centered $2 \times 2 \times 2$ k -point grid. The tungsten related limiting phases, W and WO₃ were also relaxed to the same accuracy and with Γ -centered k -point meshes of $18 \times 18 \times 18$ and $4 \times 4 \times 3$, respectively.

Defect Formalism: For a defect in charge state ‘ q ’ the formation energy $\Delta H_f(D, q)$ can be described as follows:

$$\Delta H_f(D, q) = (E^{D, q} - E^H) \pm \sum_i n_i (E_i + \mu_i) + q (E_{Fermi} + \varepsilon_{VBM}^H + v_{pot}) + E_{corr}[q] \quad (1)$$

where $E^{D, q}$ and E^H refer to the total energy of the defect (in charge state ‘ q ’) and undoped (host) supercells, respectively. The chemical potentials and elemental energies are considered as well, where n is the number of species ‘ i ’ added to or taken away from an external reservoir, E_i is energy of species i , and μ_i is the associated chemical potential. E_{Fermi} is related to the VBM of the host (ε_{VBM}^H) and v^{pot} is a term aligning the electrostatic potential of the defective supercell to that of the host. The last term in equation (1), $E_{corr}[q]$, is a post-processing correction pertaining to the finite size of the supercell and consists of two parts: an image charge correction and a band filling correction. The image

charge correction is applied due to the long-range nature of the Coulombic interaction of a defect with its periodic images. The scheme implemented herein is that formalized by Lany and Zunger [46] and adapted for non-cubic systems by Hine and Murphy.[47] Due to unrealistic band-filling present in finite-sized supercells, a correction by Lany and Zunger [46] is applied to recover the “dilute-limit”.

The formation of defects in SnO₂ can be defined within two limits arising from the formation enthalpy of SnO₂ ($\Delta H_f[\text{SnO}_2] = -5.27\text{eV}$, exp. = -5.98 eV [48]) which are the *Sn-rich/O-poor* and *Sn-poor/O-rich* bounds. The *Sn-rich/O-poor* boundary is restricted by the formation of Sn_(s) such that the chemical potentials of Sn and O are $\Delta\mu_{\text{Sn}} \leq 0\text{ eV}$ and $\Delta\mu_{\text{O}} \geq 2.64\text{ eV}$ respectively. The *Sn-poor/O-rich* limit is limited by the formation of O_{2(g)}, and thus the chemical potentials are now $\Delta\mu_{\text{Sn}} \leq -5.27\text{ eV}$ and $\Delta\mu_{\text{O}} \geq 0\text{ eV}$ for Sn and O, respectively. Considering the growth condition of the WTO film, the *Sn-rich/O-poor* condition was focused on in this study. Further limits were placed on the dopant (W) chemical potentials due to the formation of WO₃ ($\Delta H_f[\text{WO}_3] = -8.39\text{ eV}$ exp = -8.73 eV [48]). Within the *Sn-rich/O-poor* and *Sn-poor/O-rich* limits, the chemical potential limits of tungsten in SnO₂ are $\Delta\mu_{\text{W}} \leq -0.48\text{ eV}$ and $\Delta\mu_{\text{W}} \geq -8.39\text{ eV}$, respectively. The thermodynamic transition levels were calculated using the following equation:

$$\varepsilon_D \frac{q}{q'} = \frac{\Delta H_f(D, q) - \Delta H_f(D, q')}{q' - q} \quad (2)$$

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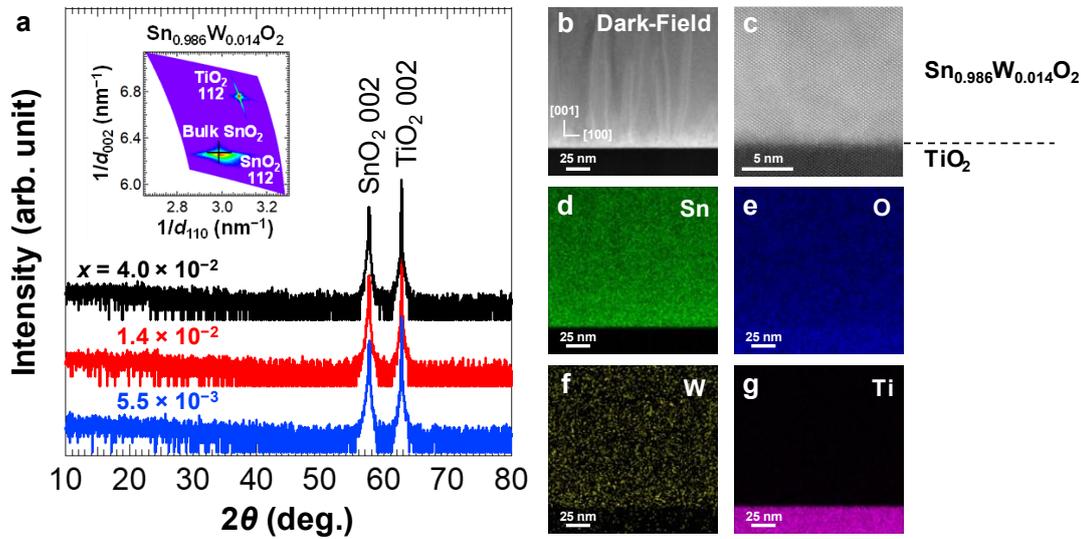


Figure 1. (a) ω - 2θ XRD patterns of $W_xSn_{1-x}O_2$ (001) films epitaxially grown on TiO_2 (001) substrates. Inset shows a reciprocal space map around the 112 diffraction peak. (b,c) Dark field STEM images of the $W_{0.014}Sn_{0.986}O_2$ film and (d-g) EDS elemental maps corresponding to the image (b).

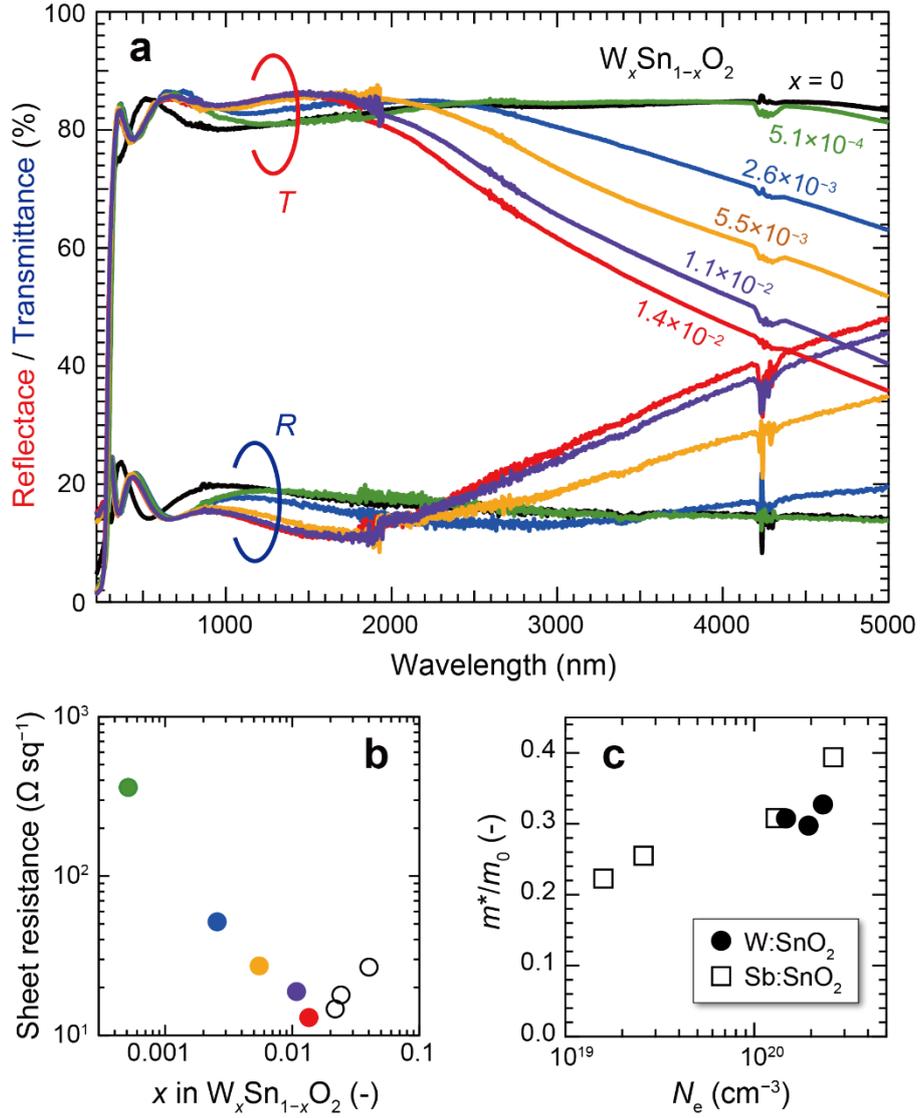


Figure 2. (a) Optical reflectance R and transmittance T spectra and (b) sheet resistance of WTO (001) epitaxial films grown on $m\text{-Al}_2\text{O}_3$ substrates. (c) Effective masses of electrons in the WTO (001) films determined by Drude fitting of the reflection spectra in the NIR region. Effective masses for Sb:SnO₂[49] are also plotted for comparison.

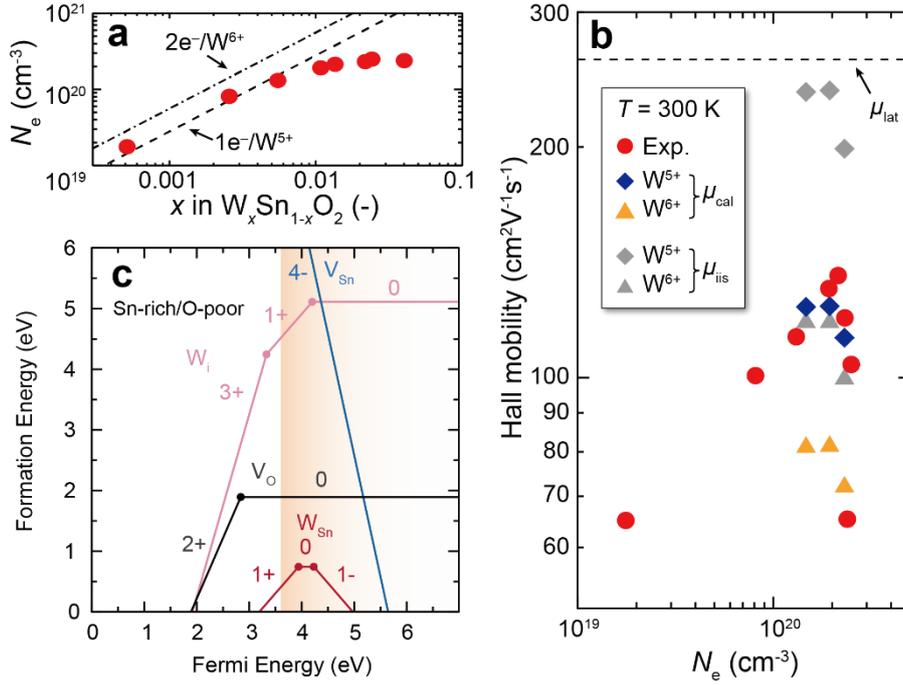


Figure 3. (a) Carrier density of WTO (001) epitaxial films on TiO₂ (001) substrates. The broken line and the dash-dotted line represent the expected n_e when all the doped W ions generate one electron per W (100% doping efficiency for W^{5+}) and two electrons per W (100% doping efficiency for W^{6+}), respectively. (b) Comparison of μ for the WTO films at 300 K (circles) with those theoretically predicted from the contribution of intrinsic scattering, μ_{cal} (diamonds) for both W^{5+} and W^{6+} . Hall mobilities limited by ionized impurity scattering, μ_{iis} , are calculated with the Brooks–Herring–Dingle (BHD) formula and plotted for comparison. m^* evaluated by Drude fitting of the optical reflectance spectra was used for calculating μ_{iis} . (c) The calculated thermodynamic transition levels under the Sn-rich/O-poor regime. The Fermi level ranges from 0 eV (VBM) to 7 eV (3.4 eV above the CBM), with the conduction band region depicted by the orange gradient.

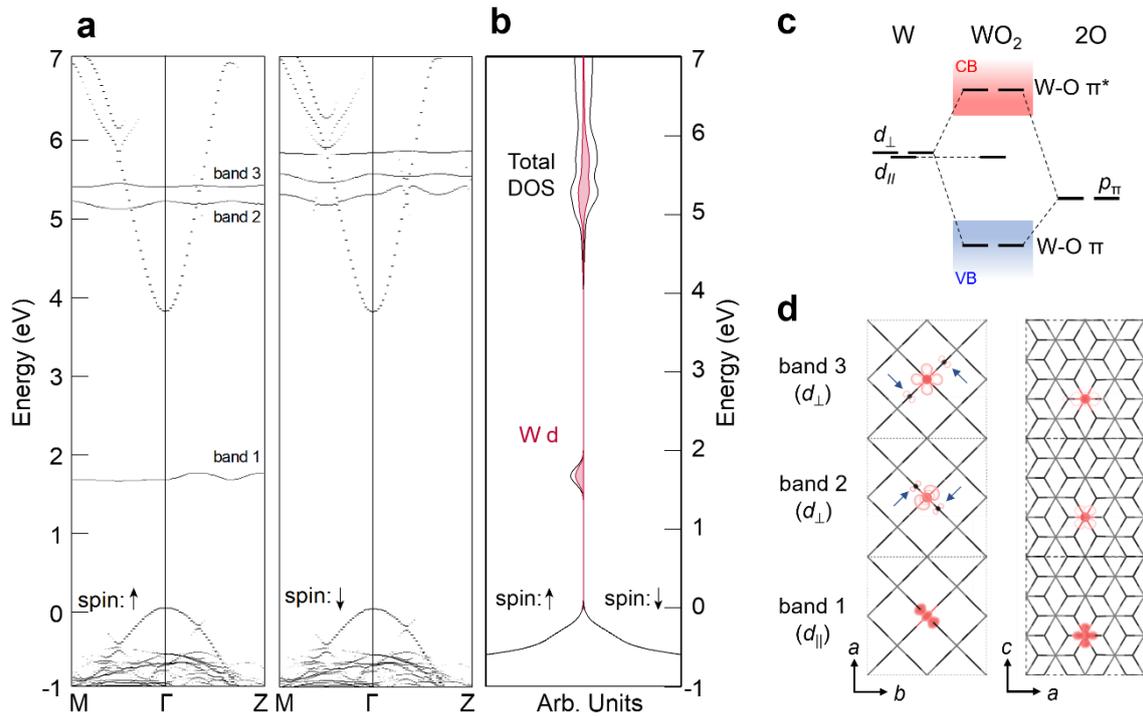


Figure 4. (a) Spin-up (left) and spin-down (right) components of the unfolded band structures and (b) density of states (DOS) for WTO in its ionized form ($1+$ charge state), in which the VBM is set to 0 eV. The DOS relating to W states is colored in (b), and the corresponding bands in (a) are numbered as 1-3. (c) Schematic diagram of W $5d t_{2g}$ states in WTO considering hybridization with the O $2p$ orbitals. Valence and conduction bands are shaded in blue and red, respectively. (d) Partial charge densities of the W $5d$ states (bands 1-3 in (a)) projected along (left) c -axis and (right) b -axis. Black arrows in the left panels indicate contribution of O $2p_{\pi}$ orbitals hybridized with d_{\perp} states of W.