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## Out of stoichiometry CuCrO<sub>2</sub> films as a promising p-type TCO for transparent electronics†

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In this work, we report the enhancement of the functional properties of CuCrO<sub>2</sub>, a promising p-type transparent conductive oxide, achieved in out of stoichiometry CuCrO<sub>2</sub> thin films synthesized by aerosol-assisted chemical vapor deposition. Out of stoichiometry films consisting of Cr-deficient CuCrO<sub>2</sub>, i.e. Cu-rich CuCrO<sub>2</sub> phase, have a resistivity value of 0.05 Ω cm and an average transmittance of 58% in the visible range, resulting in a Gordon's figure of merit of 2200 μS. This is the highest ever published figure of merit among Cu-rich CuCrO<sub>2</sub> films synthesized by chemical methods. A remarkable result is that when further increasing the Cu/(Cu + Cr) ratio, the formation of CuO was not detected, allowing the synthesis of composite films formed by Cu<sub>2</sub>O and CuCrO<sub>2</sub> p-type oxides, which are more conductive than the Cu-rich CuCrO<sub>2</sub> phase. These nanocomposite films present an improved carrier mobility, with a resistivity value of around 0.02 Ω cm, and a reduced energy gap, with a transmittance of 52%, resulting in a figure of merit of 1400 μS. Both these thin films can find applications as a hole transport layer in various transparent optoelectronic devices where p-type TCOs are required, especially when synthesized by a solution-based process at a low temperature and ambient pressure over large surface areas.

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### Introduction

Transparent electronics and the conception of see-through devices require materials combining high electrical conductivity and optical transmittance in the visible range, properties fulfilled by transparent conductive oxides (TCOs). The figure of merit as reported by Gordon<sup>1</sup> (FoM<sup>G</sup>) can be used to quantify their performance. It is defined as the ratio of the electrical conductivity  $\sigma$  to the optical absorption  $\alpha$ , resulting in:

$$\text{FoM}^G = \frac{\sigma}{\alpha} = \frac{-1}{R_{\text{sh}} * \ln(T + R)} \quad (1)$$

with  $R_{\text{sh}}$  being the sheet resistance (Ωsq<sup>-1</sup>), and  $T$  and  $R$  being the average transmittance and reflectance, respectively, in the 400–800 nm range.

The massive breakthrough of transparent electronics is hindered by the lack of p-type TCOs with properties comparable to those of their n-type counterparts, the latter being characterized by an optical transmittance of around 80% in the visible range and a resistivity value as low as 0.001 Ω cm leading to a FoM<sup>G</sup>

value of more than 200 mS.<sup>2</sup> The development of p-type TCOs with high FoM<sup>G</sup> values allows the fabrication of fully transparent p–n junctions, as previously reported for Li-doped ZnO/ZnO,<sup>3</sup> and Sb-doped p-type SnO<sub>2</sub>/Sb-doped n-type SnO<sub>2</sub>,<sup>3,4</sup> although these devices were fabricated by physical deposition methods working at a high temperature.

In the hot topic of the synthesis of highly performing p-type TCOs, copper-based delafossite oxides with the general formula Cu<sup>1+</sup>M<sup>3+</sup>O<sub>2</sub> (M = Al,<sup>5</sup> Fe,<sup>6</sup> Ge,<sup>7</sup> Ga,<sup>8</sup> Cr,<sup>9</sup> Y<sup>10</sup> or Sc<sup>11</sup>) arise as promising materials since the first report on CuAlO<sub>2</sub><sup>5</sup> was characterized by p-type conductivity and good transparency. Among these, copper chromium oxide, CuCrO<sub>2</sub>, emerges as an ideal p-type TCO with a band gap of around 3.1–3.3 eV,<sup>12</sup> ensuring good transparency and a relatively good electrical resistivity ( $\rho = 1$  Ω cm) for stoichiometric films.<sup>13</sup> These optical and electrical properties make it suitable for a wide variety of optoelectronic devices, such as photovoltaic devices,<sup>14,15</sup> light<sup>16</sup> and gas sensors,<sup>17</sup> transparent diodes<sup>18</sup> and transistors.<sup>19</sup> Furthermore, this material is also attractive for other peculiar features such as its magnetic properties,<sup>20</sup> photocatalytic ability,<sup>21</sup> and thermoelectric properties.<sup>22</sup> CuCrO<sub>2</sub> films can be synthesized through different techniques such as sputtering,<sup>23</sup> chemical vapour deposition (CVD),<sup>9,24</sup> spray pyrolysis,<sup>25,26</sup> atomic layer deposition (ALD),<sup>13</sup> sol gel,<sup>27</sup> hydro-thermal synthesis,<sup>28</sup> molecular-beam epitaxy<sup>29</sup> and pulsed laser deposition (PLD).<sup>30</sup> Among them, chemical methods allow the

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deposition over large surface areas at a relatively low temperature (<400 °C), compatible with glass and even plastic substrates. These features are extremely appealing for industrial applications.

Concerning the transport mechanisms in CuCrO<sub>2</sub>, a small polaron hopping among Cu<sup>1+</sup>/Cu<sup>2+</sup> sites<sup>12</sup> was suggested as the conduction mechanism. Theoretical calculations<sup>12</sup> showed that the p-type conductivity is related to the formation of defects, such as Cu vacancies (V<sub>Cu</sub>), which have the lowest formation energy as confirmed by the experimental reports.<sup>24</sup> Furthermore, various studies<sup>30,31</sup> highlighted the possible formation of Cu anti-site defects (Cu<sub>Cr</sub>), *i.e.* Cu atoms occupying Cr sites. Intrinsic dopants are demonstrated to be responsible for the enhancement of the charge carrier density and, consequently, of the electrical properties. The kind of defects formed in the material is believed to be dependent on the growth conditions and finally on the deviation from the stoichiometry of CuCrO<sub>2</sub>.<sup>24–26,32</sup> For instance, the high conductivity of Cu-deficient CuCrO<sub>2</sub> films is attributed to the presence of Cu vacancies as detected by using scanning transmission electron microscopy (STEM).<sup>33</sup>

Several published studies concern the characterization of Cu-poor CuCrO<sub>2</sub> films. Crépellière *et al.*<sup>24</sup> reported the synthesis by the pulsed-injection metal-organic CVD (PI-MOCVD) of films with a copper content Cu/(Cu + Cr) of 33%, still preserving the crystalline delafossite phase. These films presented a resistivity value of around 0.06 Ω cm and an average transmittance of around 50% in the visible range, yielding the highest FoM<sup>G</sup> value of 2300 μS, to date, for this material. Lunca Popa *et al.*<sup>33</sup> reported the lowest resistivity value of 0.009 Ω cm achieved for the same Cu-poor composition, Cu<sub>0.66</sub>Cr<sub>1.33</sub>O<sub>2</sub>, synthesized by direct liquid injection MOCVD (DLI-MOCVD). Farrell *et al.*<sup>25</sup> reported a slightly lower FoM<sup>G</sup> value of 350 μS for Cu-deficient CuCrO<sub>2</sub> films with a Cu/(Cu + Cr) of ≈ 30–35% and oxygen in excess (Cu<sub>0.4</sub>Cr<sub>0.5</sub>) by using a non-vacuum technique, the spray pyrolysis. They reported a resistivity value of around 0.08 Ω cm and an average transmittance of 55%.

On the other hand, Cu-rich CuCrO<sub>2</sub> films, *i.e.* Cr-poor CuCrO<sub>2</sub> (CuCr<sub>1-x</sub>O<sub>2</sub>), have also been reported as a promising p-type TCO. Ling *et al.*<sup>34</sup> reported the synthesis of CuCr<sub>1-x</sub>O<sub>2</sub> by the solid-state reaction at a high temperature, showing a variation of the resistivity of two orders of magnitude for  $x = 0.1$  compared to the stoichiometric sample.<sup>34</sup> They attributed this enhancement to hybridization between Cu 3d and O 2p orbitals and the presence of a mixed-valence band Cu<sup>1+</sup>/Cu<sup>2+</sup>. Sidik *et al.*<sup>30</sup> reported the synthesis of CuCr<sub>1-x</sub>O<sub>2</sub> by PLD with a resistivity value of 0.04 Ω cm and an average optical transmittance of 60% for  $x = 0.03$ . The enhancement of the conductivity with Cr deficiency was linked to the presence of Cu atoms in Cr-vacancies to form Cu anti-site defects. Nevertheless, it was also reported that a high Cu content leads to the formation of a parasitic CuO<sup>24,27</sup> phase, detrimental for the optoelectronic properties of the film.

To date, there is no report on the enhancement of the functional properties of Cu-rich CuCrO<sub>2</sub> thin films synthesized by aerosol-assisted metal-organic CVD (AA-MOCVD), a chemical method at a low temperature and ambient pressure, with no post-treatment. Therefore, the present study aims to

evaluate the impact of the cationic ratio, Cu/(Cu + Cr), on the electrical and optical properties. In this work, Cu-rich CuCrO<sub>2</sub> thin films are highlighted as the best compromise between conductivity and transparency. To our knowledge, this is the first report that Cu-rich CuCrO<sub>2</sub> films, synthesized by a chemical method, display a figure of merit comparable to those of their Cu-poor counterparts. Furthermore, the film growth by AA-MOCVD creates an oxygen-poor environment that prevents the formation of the parasitic CuO phase. This allows the synthesis of composite materials formed by Cu<sub>2</sub>O and CuCrO<sub>2</sub>, displaying a conductivity higher than that of Cu-rich CuCrO<sub>2</sub> thin films due to the enhancement of the carrier mobility. In the hot topic of the synthesis of efficient p-type TCOs, the present work offers two appealing candidates for transparent electronic devices, where p-type TCOs deposited at low temperature and on a large surface area are required.

## Experimental methods

The precursor solution was composed of copper(II) acetylacetonate (Cu(acac)<sub>2</sub> - CuC<sub>10</sub>H<sub>14</sub>O<sub>4</sub>, Strem chemical) and chromium(III) acetylacetonate (Cr(acac)<sub>3</sub> - CrC<sub>15</sub>H<sub>21</sub>O<sub>6</sub>, Sigma Aldrich) dissolved in ethanol. Ethylenediamine (C<sub>2</sub>H<sub>8</sub>N<sub>2</sub>, Sigma Aldrich) was added to improve the solubility of precursors in ethanol with a molar concentration twice the total one. Alkaline earth borosilicate glass (Corning 1737) was used as the substrate. The substrates were cleaned by mechanical brushing using paper and various solvents such as acetone, isopropanol, and deionized water, then they were ultrasonically cleaned in isopropanol for 20 minutes, and they were rinsed with deionized water and dried with clean compressed air.

AA-MOCVD was performed in a home-made vertical flux cold wall reactor as depicted in the study by de Oliveira *et al.*<sup>35</sup> The deposition took place at atmospheric pressure ensured by an extracting system. A nitrogen trap was used for the condensation of the organic vapors after the reaction. Dry compressed air was used as the process gas with a total flow rate of 4800 sccm. The oxygen partial pressure was  $0.21 \times 10^5$  Pa. The substrate temperature was 350 °C for all depositions. Two series of samples were synthesized. Series 1: the deposition time was set to 60 minutes and the solution consumption rate was fixed at 2 ml min<sup>-1</sup>. Two total molar concentration solutions of (Cu + Cr) of 10 mM and 20 mM with Cu/(Cu + Cr) varying between 40% and 100% were used. Series 2: after calibration of the deposition rate in series 1, 4 samples with a thickness in the range of 75–85 nm were synthesized by tuning the deposition time. The total molar concentration was fixed to 10 mM with the Cu/(Cu + Cr) values of 40%, 50%, 70% and 100%.

The composition of the films was analysed by using energy-dispersive X-ray spectroscopy (EDS) measurements with an energy beam of 15 KeV and using an Oxford Inca Energy detector in a FEI Quanta 250 field-emission scanning electron microscope (FESEM). The reported values are the average of 5 points over a surface area of 6.25 cm<sup>2</sup>. Two scanning electron microscopes (SEM) were used for the top-view and cross-section



SEM observations. Series 1 was analysed using the FEI Quanta 250 FESEM microscope, and the samples of series 2 were observed using a FEG-ZEISS-Gemini 300 microscope.

X-ray diffraction (XRD) patterns were obtained using a Bruker D8 Advance diffractometer in the Bragg–Brentano configuration ( $\theta$ – $2\theta$ ) with Cu  $K_{\alpha 1}$  radiation ( $\lambda = 0.15406$  nm). Raman spectroscopy was performed at room temperature using a Jobin Yvon/Horiba LabRam spectrometer equipped with a liquid nitrogen-cooled charge-coupled detector. The excitation source was the 488 nm line of an Ar<sup>+</sup> laser; it was focused to a spot size close to 1  $\mu\text{m}^2$  by using a 100 $\times$  objective. The laser power at the sample surface was around 80  $\mu\text{W}$ .

High-resolution transmission electron microscopy (HRTEM) observations were performed using a JEOL 2010 LaB<sub>6</sub> instrument operating at 200 kV with a 0.19 nm point-to-point resolution. The sample lamellae were prepared in cross-section by tripod mechanical polishing and Argon ion milling until the perforation of the interface. The superficial roughness was measured by using atomic force microscopy (AFM) performed using a Veeco D3100 AFM on 1  $\mu\text{m}^2$  surface and the raw data were fitted by Gwydion software.

X-ray photoelectron spectroscopy (XPS) measurements were performed using a K-alpha spectrometer from Thermo Fisher Scientific with an Al  $K_{\alpha 1,2}$  (1486.6 eV) X-ray source. The core levels of Cu, Cr and O in the film were probed in the Cu 2p, Cr 2p, and O 1s energy ranges after Ar milling at 2 KeV for 3 minutes. The experimental data were fitted with the Advantage software from Thermo Fisher Scientific. The valence band spectrum was acquired by using XPS (VB-XPS) in the constant analyser energy mode using a step size of 0.1 eV in the range of –1 to 10 eV. Previously, the samples were subjected to Ar milling under the same conditions listed before. The binding energy (BE) scale of the spectrometer was calibrated by the positions of the peaks of Au 4f<sub>7/2</sub> ( $83.9 \pm 0.1$  eV) and Cu 2p<sub>3/2</sub> ( $932.8 \pm 0.1$  eV) core levels of the pure gold and copper metals.

The optical properties of the films were investigated by UV–vis–IR spectroscopy, using a Lambda 950 spectrophotometer from PerkinElmer equipped with an integration sphere, using a wavelength step of 5 nm. The total reflection was measured on the same instrument using PTFE/BaSO<sub>4</sub> and a black trap light for the 100% and 0% references, respectively. The electrical properties were measured by using a linear 4-probe system with a distance of 1 mm between the tips. The sheet resistance values correspond to the average of 5 points for each sample. The electronic transport properties were measured by the Hall effect at room temperature in a homemade setup with a magnetic field of 0.5 T.

## Results and discussion

### Compositional analysis and growth rate

The Cu/(Cu + Cr) cationic content determined by EDS in the films from series 1 is reported in Fig. 1a for the two molar concentrations used as a function of the initial solution

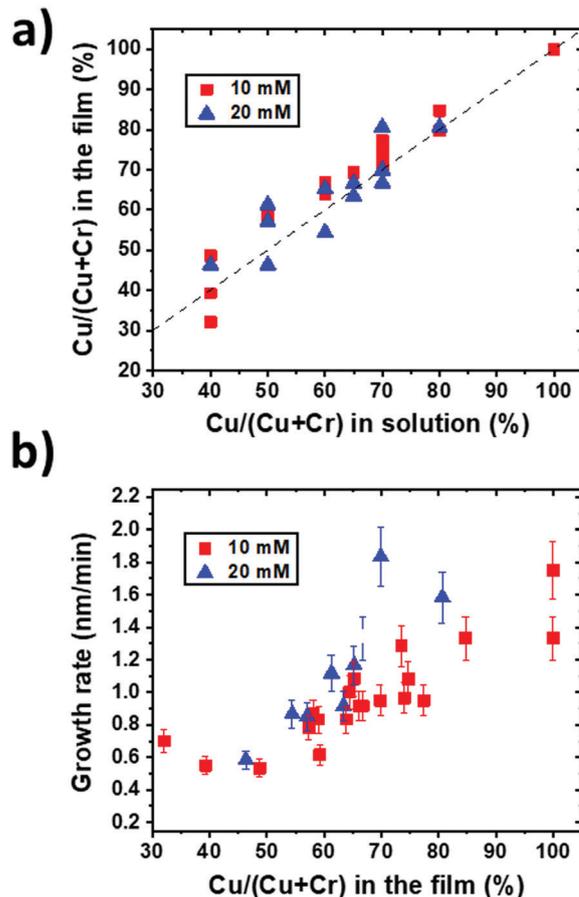


Fig. 1 (a) Cu/(Cu + Cr) cationic ratio in the film measured by EDS as a function of the cationic ratio in the solution for the depositions performed with 10 mM and 20 mM total cationic molar concentration solutions of series 1. (b) Growth rate as a function of the cationic ratio in the films for the depositions performed with 10 mM and 20 mM total cationic molar concentration solutions. The error bar represents the 10% relative error from SEM measurements.

composition. The composition reproducing the starting solution is indicated by the dashed line. This graph highlights that the incorporation of Cu in the films is generally slightly higher than the incorporation of Cr, suggesting a higher reactivity of the Cu precursor than the Cr one under these deposition conditions. This can be explained by a higher thermal stability of Cr(acac)<sub>3</sub> in comparison with that of Cu(acac)<sub>2</sub>,<sup>36</sup> implying a lower reactivity of the Cr precursor than the Cu one. Moreover, it is observable that the composition of the films is independent of the total molar concentration of the solution for the studied conditions. These results confirm the reproducibility of the process and the control over the composition of the films.

The film thickness measured from the SEM cross-section images allows calculating the deposition rate for different compositions. The growth rate as a function of the film composition is reported in Fig. 1b for the two total molar concentrations used for these experiments. In these deposition conditions, it is evidenced that the Cu-rich films grow faster, and generally, a higher total molar concentration leads to the



synthesis of thicker films but not at twice the thickness. When increasing the concentration of the Cu precursor, the higher growth rate suggests that the decomposition of  $\text{Cr}(\text{acac})_3$  and its reaction with Cu and O are catalysed by  $\text{Cu}(\text{acac})_2$  and its by-products, in agreement with the deposition of  $\text{CuCrO}_2$  by PI-MOCVD,<sup>24</sup> where a similar trend was found. Moreover, these results are coherent with the growth mechanism proposed for  $\text{CuCrO}_2$  deposited by spray pyrolysis<sup>25</sup> as the film formation occurs by the consecutive stacking of  $\text{CrO}_6$  octahedra and O–Cu–O dumbbells. Initially, Cr is adsorbed until the surface is saturated. Successively,  $\text{Cu}(\text{acac})_2$  decomposes and reacts with the Cr terminated surface until the surface is adequately Cu covered. Finally, the Cr atoms react with the Cu terminated surface allowing the film growth. A lack of Cu inhibits the formation of O–Cu–O dumbbells and the subsequent film growth, explaining the increasing growth rate with the Cu content.

These analyses allowed us to control the composition and thickness of the films and were used to synthesize the samples of series 2. The films were grown from the solution concentrations of  $\text{Cu}/(\text{Cu} + \text{Cr})$  of 40%, 50%, 70% and 100%. Several runs lead to film compositions corresponding to  $50 \pm 5\%$ ,  $63 \pm 5\%$ ,  $77 \pm 5\%$  and 100% of Cu. These samples will be labelled as  $\text{CuCrO}_2\text{:X}$  in the following, with X corresponding to the Cu content (in %) in the films as measured by EDS.

### Structural properties

The structural properties of the  $\text{CuCrO}_2\text{:X}$  films of series 2 were investigated by XRD. In Fig. 2a, XRD patterns were compared with the ICDD reference patterns of the rhombohedral  $\text{CuCrO}_2$  delafossite structure (space group  $R\bar{3}m$ , ICDD 04-010-3330) and cubic  $\text{Cu}_2\text{O}$  (space group  $Pn\bar{3}m$ , ICDD 00-005-0667), presented in the top windows. The stoichiometric  $\text{CuCrO}_2\text{:50\%}$  pattern, as shown in Fig. 2a(1), shows two wide peaks at  $2\theta = 36.4^\circ$  and  $2\theta = 62.4^\circ$ . They can be assigned to the (012) and (110) reflections of  $\text{CuCrO}_2$ . The broadness of these peaks indicates a poor crystallinity. The weak band in the 40–50° range is attributed to the glass substrate.<sup>37</sup> The Cu-rich  $\text{CuCrO}_2\text{:63\%}$  pattern (Fig. 2a(2)) shows only one narrow (012) peak, thus suggesting a higher crystallization and an increase of grain size. These results are in good agreement with the previous report on Cr-deficient  $\text{CuCrO}_2$ ,<sup>30</sup> where the authors reported a change in the preferred crystal orientation from the (006) plane to the (012) plane with an increase of Cr deficiency. Increasing the copper content up to the  $\text{CuCrO}_2\text{:77\%}$ , the composition (Fig. 2a(3)) leads to the additional formation of a  $\text{Cu}_2\text{O}$  cubic phase as stated by the presence of the  $\text{Cu}_2\text{O}$  (200) reflection at  $2\theta = 42^\circ$ , in agreement with the pattern of the  $\text{Cu}_2\text{O}$  film synthesized under the same deposition conditions (Fig. 2a(4)). This result highlights the deposition of the solid solution composed of  $\text{Cu}_2\text{O}$  and  $\text{CuCrO}_2$ . The overall poor crystallinity of our samples is attributed to the low deposition temperature and atmospheric pressure used during the synthesis. These results allow us to classify the thin films of series 2 in 4 categories: stoichiometric  $\text{CuCrO}_2$ , Cu-rich  $\text{CuCrO}_2$ ,  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  and  $\text{Cu}_2\text{O}$ .

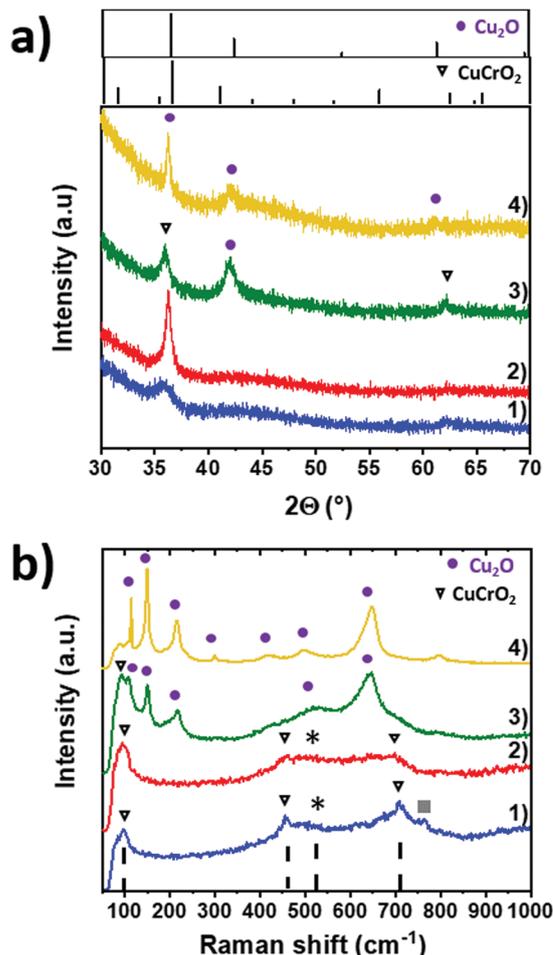


Fig. 2 (a) XRD pattern of the (1) stoichiometric  $\text{CuCrO}_2$ , (2) Cu-rich  $\text{CuCrO}_2$ , (3)  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  and (4)  $\text{Cu}_2\text{O}$  films. The top windows show the ICDD references (ICDD 04-010-3330) for  $\text{CuCrO}_2$  and (ICDD 00-005-0667) for  $\text{Cu}_2\text{O}$ . (b) Raman spectra of the same films.

Raman spectroscopy was used on the same samples to confirm this hypothesis. Raman spectra are shown in Fig. 2b. Our stoichiometric and Cu-rich  $\text{CuCrO}_2$  samples are characterized by the three Raman modes at  $101\text{ cm}^{-1}$ ,  $460\text{ cm}^{-1}$ , and  $709\text{ cm}^{-1}$  (triangles in Fig. 2b(1) and (2)), assigned to the  $E_u$ ,  $E_g$ , and  $A_{1g}$  modes of  $\text{CuCrO}_2$ , respectively, in agreement with the previous report on this material.<sup>38</sup> Additional modes are attributed to the presence of defects able to relax the Raman selection rules. The first one labelled with \* and observed in the range of  $500\text{--}670\text{ cm}^{-1}$  is attributed to the presence of intrinsic defects.<sup>39</sup> Its intensity increases with Cu content, suggesting a greater amount of defects for Cu-rich  $\text{CuCrO}_2$  when compared to the stoichiometric  $\text{CuCrO}_2$ . The second one, at  $770\text{ cm}^{-1}$ , labelled as ■, has not been reported before and further investigations are required for its assignment.  $\text{Cu}_2\text{O}$  modes are expected at  $108\text{ cm}^{-1}$ ,  $149\text{ cm}^{-1}$ ,  $216\text{ cm}^{-1}$ ,  $495\text{ cm}^{-1}$  and  $649\text{ cm}^{-1}$ <sup>40</sup> and the  $\text{CuO}$  modes at  $297\text{ cm}^{-1}$ ,  $347\text{ cm}^{-1}$ , and  $632\text{ cm}^{-1}$ .<sup>41</sup> The absence of the  $\text{CuO}$  and  $\text{Cu}_2\text{O}$  Raman modes in these samples confirms that Cu-based oxides are either absent or amorphous and dispersed in  $\text{CuCrO}_2$  films, in agreement with the XRD results.



The successful deposition of the crystallized non-stoichiometric  $\text{CuCrO}_2$  without any detectable secondary phase in the films of the Cu composition of up to 63% is achieved. Further results presented in the next section (Fig. 4c) confirm the presence of  $\text{Cu}_2\text{O}$  in the films with a  $\text{Cu}/(\text{Cu} + \text{Cr})$  of  $> 65\%$ .

For the  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  composite film (Fig. 2b(3)), the Raman modes at  $99\text{ cm}^{-1}$  is attributed to the  $E_u$  mode of the delafossite phase. Additional modes are assigned to  $\text{Cu}_2\text{O}$ , as confirmed by the spectrum of the  $\text{Cu}_2\text{O}$  film deposited under the same conditions (Fig. 2b(4)). The formation of Cr oxides is excluded due to the Cu-rich/Cr-poor environment used for the growth.

Thus, the classification of the samples of series 2 is well confirmed, allowing us to label as the stoichiometric  $\text{CuCrO}_2$ , films with a Cu composition of  $\sim 50\%$ , Cu-rich  $\text{CuCrO}_2$  with a Cu composition of  $\sim 65\%$ ,  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  with a Cu composition of  $\sim 77\%$ , and  $\text{Cu}_2\text{O}$ .

The films corresponding to the out of stoichiometry composition of  $\text{CuCr}_{0.58}\text{O}_x$  ( $\text{Cu}/(\text{Cu} + \text{Cr}) = 63\%$ ) are characterized by a greater amount of defects in comparison with the stoichiometric compound. The preservation of the crystalline delafossite phase within a wide variation in the cationic ratio is consistent with previous reports on the  $\text{Cu}_{0.66}\text{Cr}_{1.33}\text{O}_2$  ( $\text{Cu}/(\text{Cu} + \text{Cr})$  of 33%) by DLI-MOCVD<sup>42</sup> and PI-MOCVD.<sup>24</sup> Moreover, it is important to note that  $\text{CuO}$  is not detected in any of our films, despite  $\text{Cu}^{2+}$  being present in the starting precursor. As described in the work by Lim *et al.*,<sup>43</sup> the gas phase decomposition of  $\text{Cu}(\text{acac})_2$  results in the formation of metallic copper condensing on the substrate. Then,  $\text{Cu}^0$  oxidizes to  $\text{Cu}_2\text{O}$ . When the oxygen partial pressure is too high during the growth,  $\text{Cu}_2\text{O}$  undergoes a secondary oxidation reaction to form  $\text{CuO}$ . The direct oxidation of  $\text{Cu}^0$  to  $\text{Cu}^{2+}$  is excluded because the Cu oxidation state transforms successively from  $\text{Cu}^0$  to  $\text{Cu}^{1+}$  and then to  $\text{Cu}^{2+}$ .<sup>44</sup> Thus, the absence of  $\text{Cu}^0$  and  $\text{CuO}$  in our films demonstrates that the used oxygen partial pressure is adequately high to completely oxidize  $\text{Cu}^0$  to  $\text{Cu}^{1+}$ , while still sufficiently low to prevent the formation of  $\text{CuO}$ . The reaction with Cr can also be a stabilising factor for  $\text{Cu}^{1+}$ . Furthermore, the deposition of a composite material formed solely by  $\text{Cu}_2\text{O}$  and  $\text{CuCrO}_2$  is in agreement with the phase diagram of the Cu–Cr–O system under these deposition conditions.<sup>45</sup> As shown by the thermodynamic results, the use of a low oxygen partial pressure allows the coexistence of  $\text{Cu}_2\text{O}$  and  $\text{CuCrO}_2$  phases, preventing the formation of  $\text{CuO}$ , which would

be detrimental for the synthesis of transparent conductive materials.

### Morphology and microstructure

Morphological and microstructural properties of the films are critical parameters for their integration into devices. The evolution of the morphology and grain size with the composition for the samples of series 2 with a similar thickness is presented in Fig. 3. Stoichiometric  $\text{CuCrO}_2$  (Fig. 3a) presents a dense and close-packed morphology with small well-defined grains, whereas an irregular grain shape with an overall increase in grain size is observed in Cu-rich  $\text{CuCrO}_2$  (Fig. 3b), in agreement with the XRD results. The  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  composite film (Fig. 3c) is composed of more rounded grains with a heterogeneous shape.

Cross-section images in insets clearly show the similar thickness of the three films and a variation of the superficial roughness with the  $\text{Cu}/(\text{Cu} + \text{Cr})$  cationic ratio, with a maximum roughness for Cu-rich  $\text{CuCrO}_2$ . The AFM measurements confirmed the trend of the roughness with values of root mean square roughness of  $0.99 \pm 0.15\text{ nm}$ ,  $2.08 \pm 0.35\text{ nm}$  and  $1.42 \pm 0.18\text{ nm}$  for  $\text{CuCrO}_2:59\%$ ,  $\text{CuCrO}_2:65\%$  and  $\text{Cu}_2\text{O} + \text{CuCrO}_2:77\%$  films, respectively.

The microstructure of two Cu-rich films,  $\text{CuCrO}_2:59\%$  and  $\text{CuCrO}_2:65\%$ , and of the  $\text{Cu}_2\text{O} + \text{CuCrO}_2:73\%$  composite film was analysed using TEM. HRTEM images and SAED patterns are presented in Fig. 4. The Cu-rich  $\text{CuCrO}_2$  thin films present a nanocolumnar microstructure with a broad orientation distribution, as reported previously by our group.<sup>16</sup> The increase of the Cu content leads to the growth of wider grains with higher vertical alignment, as visible by the comparison between  $\text{CuCrO}_2:59\%$  (Fig. 4a) and  $\text{CuCrO}_2:65\%$  (Fig. 4b). These results are in good agreement with SEM observations, as the growth of bigger nanocolumnar grains is likely related to the increase in the superficial roughness for the Cu-rich films. SAED patterns confirm that both films crystallize in the  $\text{CuCrO}_2$  structure without any detectable secondary phase, in agreement with the Raman and XRD results. This result corroborates that there is a wide compositional range, up to  $\text{Cu}/(\text{Cu} + \text{Cr}) = 65\%$ , in which the crystalline delafossite phase is preserved despite the variation of stoichiometry.

For the  $\text{Cu}_2\text{O} + \text{CuCrO}_2:73\%$  nanocomposite film (Fig. 4c), the formation of the secondary  $\text{Cu}_2\text{O}$  phase results in more

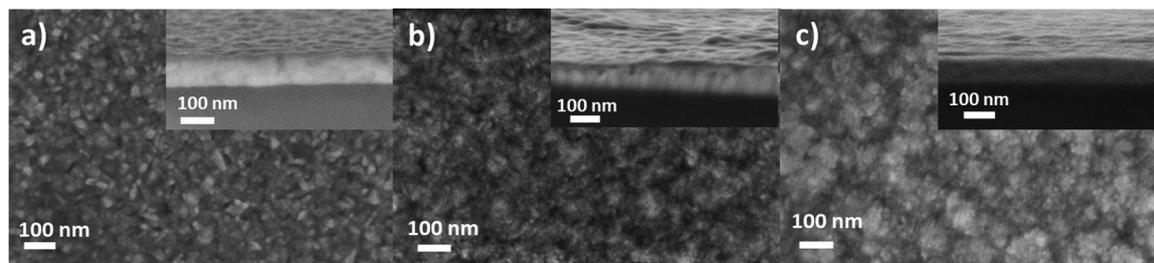


Fig. 3 Top view and cross-section (inset) SEM micrographs of the films of series 2. (a) The stoichiometric  $\text{CuCrO}_2$ , (b) Cu-rich  $\text{CuCrO}_2$  and (c)  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  samples. The scale bar is given on each micrograph.



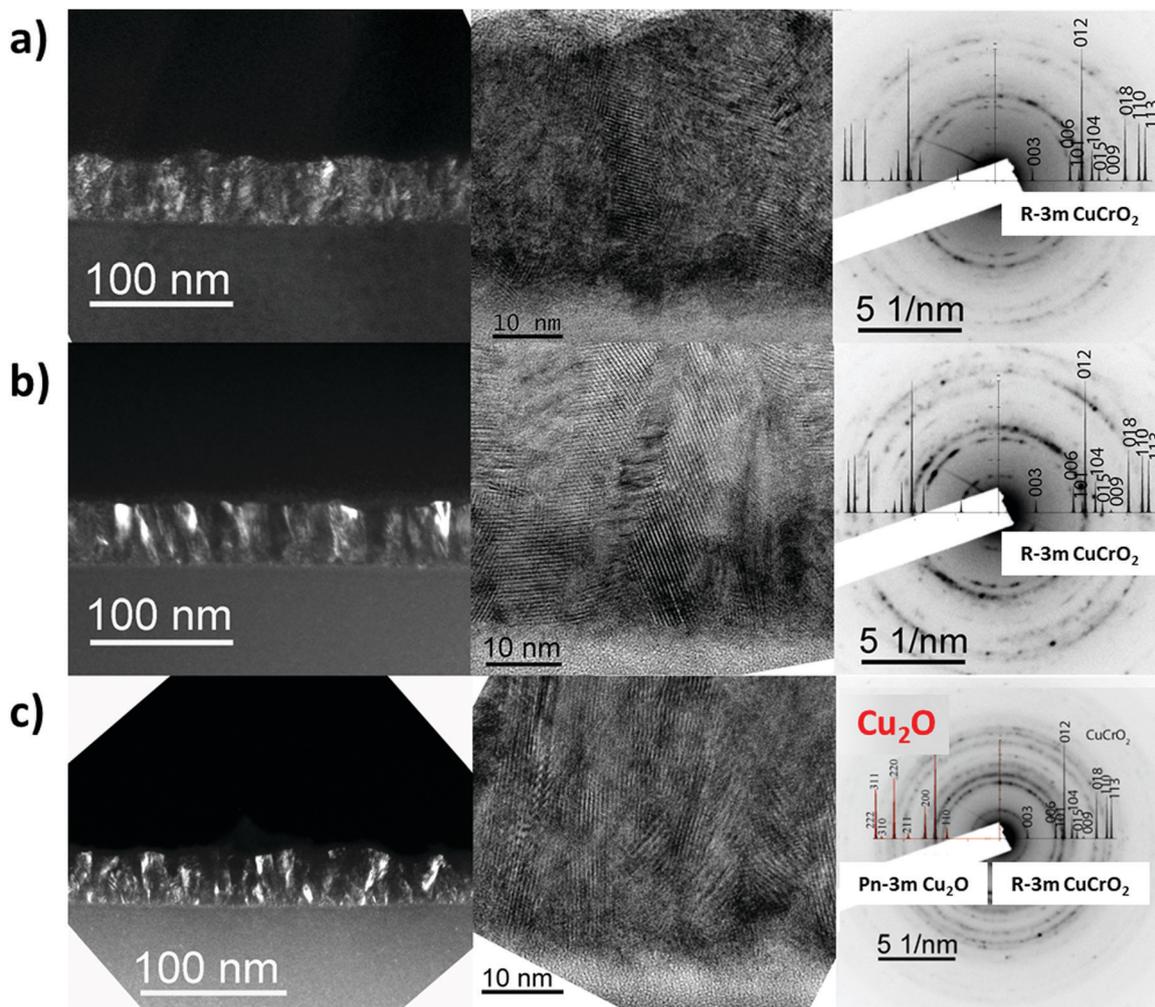


Fig. 4 Microstructural characterisation by TEM of the Cu-rich  $\text{CuCrO}_2$  films. From the left, cross-section HRTEM in the dark mode, cross-section HRTEM in the bright mode and SAED patterns for the (a)  $\text{CuCrO}_2$ :59%, (b)  $\text{CuCrO}_2$ :65% and (c)  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  ( $\text{Cu}/(\text{Cu} + \text{Cr}) = 73\%$ ) samples. The corresponding SAED patterns identifying the  $\text{CuCrO}_2$  rhombohedral delafossite phase and the  $\text{Cu}_2\text{O}$  cubic phase are presented along with the theoretical diffraction patterns of the present phases as computed with the software JEMS.

rounded grains. The film is composed of both  $\text{Cu}_2\text{O}$  and  $\text{CuCrO}_2$  as confirmed by SAED, which is consistent with the Raman and XRD results.

### Surface chemistry analysis

XPS was used to determine the chemical state of the different elements in the bulk of the films by the analysis of Cu 2p (Fig. 5a) and Cr 2p (Fig. 5b) contributions. The evolution of the intensity with  $\text{Cu}/(\text{Cu} + \text{Cr})$ , increasing for Cu 2p<sub>3/2</sub> and decreasing for Cr 2p<sub>3/2</sub>, suggests an increase in the Cu content and a reduction in the Cr content, in agreement with the EDS quantification. The spectra from the Cu 2p<sub>3/2</sub> core level (Fig. 5a) are similar for all the samples. They are characterized by a moderate intensity of Cu(II) satellite peaks between 940 eV and 945 eV. This feature is associated with the  $\text{Cu}^{2+}$  state. The main peak is centred at 932.6 eV. The fitting of this peak for the three samples is shown in Fig. S1 in the ESI.† This contribution can be ambiguously attributed to  $\text{Cu}^0$  and  $\text{Cu}^{1+}$  due to their close binding energies centred at 932.5 eV and 932.6 eV, respectively. The width

of the peak ( $\text{FWHM} \approx 1.2$  eV) suggests the presence of Cu in the monovalent state. Besides, a small contribution of  $\text{Cu}^{2+}$  ( $\text{FWHM} \approx 2.0$  eV), centred at 933.7 eV, is present in the bulk of the samples as confirmed by the asymmetry of the Cu 2p<sub>3/2</sub> peak towards higher binding energies.

The Cr 2p<sub>3/2</sub> core level (Fig. 5b) is centred at a binding energy of 576 eV corresponding to  $\text{Cr}^{3+}$  in agreement with the previous report on  $\text{CuCrO}_2$ .<sup>24</sup> Nevertheless, the distinction among the Cr species is complicated due to the presence of the Cu Auger peak, Cu LMM, centred at 569.9 eV. Again, the position of this Auger peak suggests the presence of  $\text{Cu}^{1+}$  rather than the metallic Cu,<sup>46</sup> which should be centred at 568.3 eV.<sup>47</sup> These results indicate that Cu and Cr are mainly in the 1+ and 3+ oxidation states, respectively, despite the stoichiometry variation. The O 1s peak (not shown here) was centred at around 530 eV.

The existence of  $\text{Cu}^{2+}$  does not necessarily imply the formation of  $\text{CuO}$ , not detected in our films, because it has been reported that  $\text{Cu}^{2+}$  can be present in the planar triangular



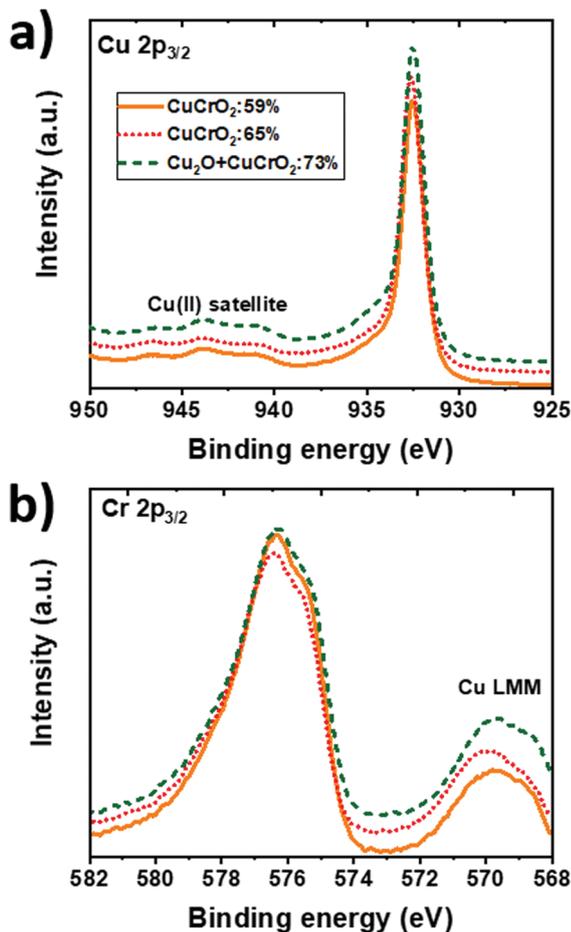


Fig. 5 XPS spectra in the (a) Cu  $2p_{3/2}$  and (b) Cr  $2p_{3/2}$  orbitals energy ranges for CuCrO<sub>2</sub>:59%, CuCrO<sub>2</sub>:65% and Cu<sub>2</sub>O + CuCrO<sub>2</sub> (Cu/(Cu + Cr) = 73%). The spectra were shifted for clarity.

network of the lattice, balanced by oxygen interstitial.<sup>18</sup> This is in agreement with the results obtained by the combustion synthesis of CuCrO<sub>2+x</sub>,<sup>18</sup> where the pure delafossite phase was detected despite the higher amount of Cu<sup>2+</sup> species (37% Cu<sup>1+</sup> vs. 63% Cu<sup>2+</sup>). The presence of Cu<sup>2+</sup> in our samples can be induced by the Cr deficiency, leading to an excess of holes at the Cu sites. This is supported by Ling *et al.*<sup>34</sup> in the case of films obtained by the solid state reaction due to the fact that the Cu<sup>2+</sup> content increases when the Cr content decreases. In our case, because of the O-poor environment used during the growth, the formation of Cu<sup>2+</sup> is not favoured.

Table 1 presents the contribution in % of Cu<sup>1+</sup> and Cu<sup>2+</sup> to the Cu  $2p_{3/2}$  spectra. The FWHM of the fitted peaks is also included in brackets. This procedure leads to a maximum value of the Cu<sup>1+</sup>/Cu<sup>2+</sup> ratio of CuCrO<sub>2</sub>:65%. As proposed by Ling *et al.*,<sup>34</sup> this fact suggests a favourable electronic structure with an improvement of hybridization between Cu 3d and O 2p orbitals. A more recent work by Singh *et al.*<sup>48</sup> confirmed that the conduction mechanism takes place among Cu<sup>1+</sup>-O-Cu<sup>2+</sup> sites, rather than directly between Cu<sup>1+</sup> and Cu<sup>2+</sup> sites. We can speculate that in both cases, with the conduction taking place *via* direct Cu<sup>1+</sup>-Cu<sup>2+</sup> or *via* Cu<sup>1+</sup>-O-Cu<sup>2+</sup> sites, the maximization

Table 1 Fitting results of the Cu  $2p_{3/2}$  XPS peak. Reported information are the cationic ratio in the film as probed by EDS and distribution in the Cu  $2p_{3/2}$  core level of monovalent (Cu<sup>1+</sup>) and bivalent (Cu<sup>2+</sup>) copper ions (in %). For each component, the corresponding FWHM of the fitted peak is reported in the brackets

| Cu/(Cu + Cr) in the film by EDS (%) | Cu <sup>1+</sup> (%) (FWHM) | Cu <sup>2+</sup> (%) (FWHM) |
|-------------------------------------|-----------------------------|-----------------------------|
| 59                                  | 80 (1.26 eV)                | 20 (2.05 eV)                |
| 65                                  | 87 (1.25 eV)                | 13 (2.05 eV)                |
| 73                                  | 79 (1.22 eV)                | 21 (2.04 eV)                |

of the Cu<sup>1+</sup>/Cu<sup>2+</sup> ratio is required to increase the number of hopping sites and, thus, the film conductivity.

VB-XPS was performed on the Cu-rich film with a Cu/(Cu + Cr) of 65% (Fig. S2 in the ESI<sup>†</sup>) and was compared with the curve resulting from the band theory, adapted by the work of Yokobori *et al.*<sup>49</sup> The theoretical spectra shows that the maximum of the valence band is dominated by the Cr 3d states (labelled with  $\alpha$ ) with a contribution attributed to the Cu 3d states (indicated with  $\beta$ ), responsible for the peak with the highest intensity. Additional contributions at higher binding energies (labelled with  $\gamma$ ) are attributed to the O 2p orbital. We can distinguish the 3 contributions in the experimental spectra, although the resolution of the spectrometer was too low to obtain additional information. The reduced intensity of the Cr states compared to that of the Cu ones can be attributed to the lower cross-section of the trivalent cation compared to Cu.<sup>50</sup> However, we may speculate that the variation in intensities can also be linked to the Cu abundance/Cr deficiency in our films, leading to the limited presence of the shoulder of the Cr 3d states at about 1.5 eV. Consequently, we can assert that the Fermi level remains close to the top of the valence band, in agreement with the high charge density of delafossite oxides and with the previous report of VB-XPS for this compound.<sup>29</sup> The use of a higher resolution XPS spectrometer will be essential to obtain a proper quantification of the energy difference between the Fermi level and the valence band edge.

Electrical measurements during annealing up to 200 °C under N<sub>2</sub>/O<sub>2</sub> with 20% O<sub>2</sub> (not shown here) were performed on a 200 nm thick sample with a Cu/(Cu + Cr) of 61%. The resistivity variation with the temperature was fitted using the Arrhenius equation, leading to an activation energy of 76 meV. This value ranges around 3 kBT, highlighting a highly degenerate semiconductor. This result is in good agreement with the proximity of the Fermi level with the top of the valence band as confirmed by our VB-XPS spectra.

### Electrical properties

The impact of the stoichiometry on the electrical properties of the CuCrO<sub>2-x</sub> thin films is reported in Fig. 6 for all our samples (series 1 and series 2). The variation of the resistivity values with Cu/(Cu + Cr) in the film follows a U-shape curve, which was fitted using an exponential quadratic law as a guide for the eyes. The values related to the 4 sample categories of series 2 were marked with arrows. They prove that, for a given composition, the thickness variations are not strongly affecting the electrical properties. It is important to note that the minimum of the curve



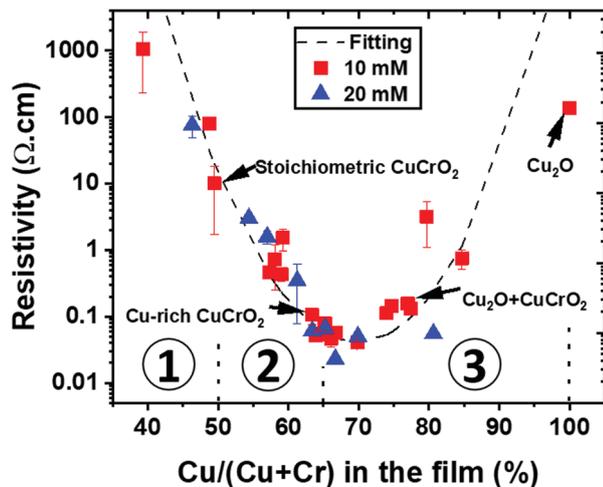


Fig. 6 Variation of the resistivity with the cationic ratio in the out of stoichiometry  $\text{CuCrO}_2$  thin films. The values corresponding to the samples of series 2 are indicated by the arrows.

does not correspond to the stoichiometric samples but to the Cu-rich  $\text{CuCrO}_2$  ones. The lowest resistivity values, below  $0.1 \Omega \text{ cm}$ , were obtained for the Cu-rich  $\text{CuCrO}_2$  films with a  $\text{Cu}/(\text{Cu} + \text{Cr})$  value between 60 and 70%, corresponding to the Cu-rich  $\text{CuCrO}_2$  and  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  composite films, respectively. For more clarity, the graph has been divided in three regions: region 1, from Cu-poor  $\text{CuCrO}_2$  to the stoichiometric one, *i.e.* a  $\text{Cu}/(\text{Cu} + \text{Cr})$  value from 40% to 50%; region 2, Cu-rich  $\text{CuCrO}_2$  with a  $\text{Cu}/(\text{Cu} + \text{Cr})$  from 51% to 65%; and region 3 from the composite  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  to the pure  $\text{Cu}_2\text{O}$  corresponding to a  $\text{Cu}/(\text{Cu} + \text{Cr})$  from 66% to 100%. In region 1, Cu-poor  $\text{CuCrO}_2$  shows a very high resistivity value of around  $1000 \Omega \text{ cm}$ , reduced down to  $10 \Omega \text{ cm}$  for the stoichiometric samples, comparable with the values reported in the literature.<sup>13</sup> In region 2, the resistivity is reduced by two orders of magnitude, reaching values below  $0.1 \Omega \text{ cm}$  for Cu-rich  $\text{CuCrO}_2$  films with a  $\text{Cu}/(\text{Cu} + \text{Cr})$  value between 60 and 65%. The lowest resistivity value of  $0.02 \Omega \text{ cm}$  is achieved in region 3 for the  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  composite film with 67% of Cu. The electrical properties are then deteriorated up to the value around  $100 \Omega \text{ cm}$  for the pure  $\text{Cu}_2\text{O}$ .

The low conductivity of samples in region 1 can be attributed to their low crystallinity of the films for this composition. Although this is only a qualitative explanation, the enhancement of the conductivity of two orders of magnitude for samples with a composition in the 50–65% range can be confirmed through simultaneous factors as established by Chen *et al.*<sup>51</sup> On the one hand, the increase in the cationic ratio can result in the formation of defects such as Cu vacancy and Cu antisite that have a lower formation energy than others.<sup>52</sup> In our Cu-rich/Cr-deficient conditions, we infer that the additional Cu atoms might be placed in the Cr vacancies to form Cu antisite defects ( $\text{Cu}_{\text{Cr}}$ ). This will be balanced by the generation of holes, leading to an increase of the charge carrier density in the films,<sup>51</sup> thus resulting in a lower resistivity of the film. Other defect species can play a major role in the electrical properties.

In region 2, the films within the 60–65% compositional range show an improved crystallinity as indicated by the higher intensity of the (012) reflection in XRD and the growth of bigger grains, as observed by SEM and TEM. This is congruent with the work by Sidik *et al.*<sup>30</sup> where Cr deficiency was demonstrated to enhance the crystallinity of the films. Furthermore, as stated before, the enhancement of the conductivity can be attributed to the favourable electronic structure of these films through the maximization of the  $\text{Cu}^{1+}/\text{Cu}^{2+}$  ratio, as proved by XPS. Nevertheless, the lowest resistivity value of  $0.02 \Omega \text{ cm}$  was achieved in region 3 for the  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  composite film with 67% of Cu. Increasing the Cu content to above 65% favours, in our deposition conditions, the formation of the  $\text{Cu}_2\text{O}$  phase in the delafossite phase that is more conductive, thus increasing the electrical resistivity up to the values obtained for the pure  $\text{Cu}_2\text{O}$ . This increase can be attributed to the change in grain shape and degradation of the film crystallinity. In comparison, the films deposited by PI-MOCVD at low pressure with a similar composition resulted in the synthesis of  $\text{CuO} + \text{Cu}_2\text{O} + \text{CuCrO}_2$ ,<sup>24</sup> with a resistivity higher than that of the delafossite phase out of stoichiometry. This finally demonstrates that the fact to prevent the formation of CuO, as reported in our study, is promising for the synthesis of highly conductive films.

Out of stoichiometry  $\text{CuCrO}_2$  films with no detectable secondary phase are characterized by a low carrier mobility, being under the detection limit of our experimental setup. This allows setting an upper limit of  $0.1 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$  for these films. The Hall effect measurements were successfully performed for  $\text{Cu}_2\text{O} + \text{CuCrO}_2$ :73% and pure  $\text{Cu}_2\text{O}$ , confirming the p-type behaviour of these films. The mobility values of  $0.65 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$  and  $5 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$  were obtained for these two samples, respectively. The corresponding charge carrier density values were  $9.3 \times 10^{18} \text{ cm}^{-3}$  and  $1.5 \times 10^{16} \text{ cm}^{-3}$ , respectively. These results reveal an improvement in mobility by the formation of  $\text{Cu}_2\text{O}$ . This finding allowed us to conclude that the lowest resistivity achieved for  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  with 67% of Cu is due to the combination of the good mobility of  $\text{Cu}_2\text{O}$  and the high charge density achieved for  $\text{CuCrO}_2$ .

### Optical characterization and figure of merit

The optical properties of the samples of series 2 were probed by UV-vis spectroscopy and the results are reported in Fig. 7. As observed in Fig. 7a, the total transmittance is reduced when increasing the copper content. The average value in the 400–800 nm visible range is around 63% for the stoichiometric  $\text{CuCrO}_2$  film and decreases down to 45% for  $\text{Cu}_2\text{O}$ . The reduced slope of the transmittance *versus* wavelength curve observed for the  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  sample is a further indication of the secondary phase in the film.<sup>53</sup> The presence of another phase will create different optical transition levels, finally resulting in a smoother slope of the transmittance curve. The reflectance (Fig. 7b) does not follow any clear trend, with an average value in the 400–800 nm range varying from 26% for the  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  composite film to 32% for the Cu-rich  $\text{CuCrO}_2$  film. The shift of the reflectance minimum could be attributed to a



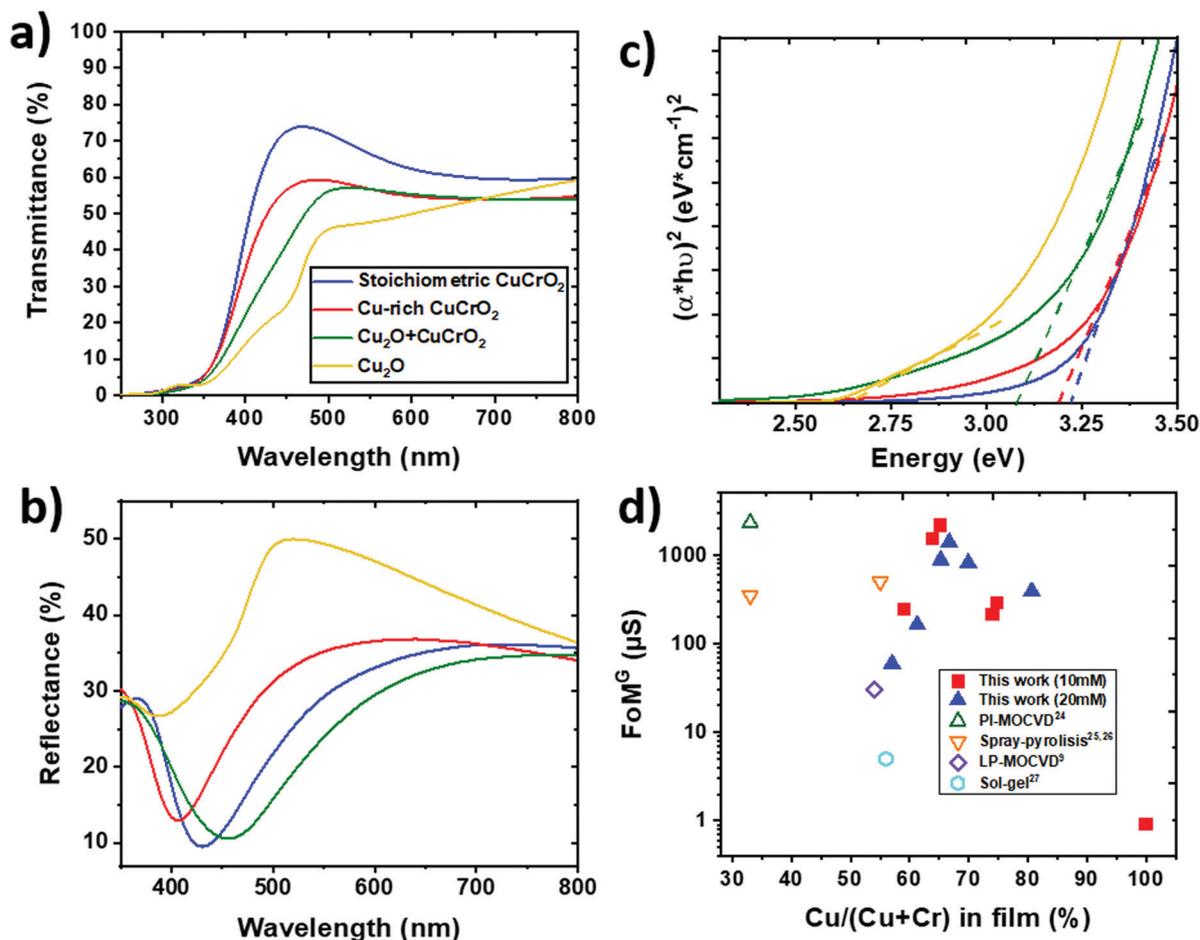


Fig. 7 Optical properties and figure of merit for the out of stoichiometry  $\text{CuCrO}_2$  films of series 2. (a) Transmittance, (b) reflectance, (c) Tauc plot, and (d)  $\text{FoM}^{\text{G}}$  by Gordon,  $\text{FoM}^{\text{G}}$ , for the various compositions and the two total molar concentrations that were used (series 1) and the values estimated from the literature for various chemical deposition techniques.

variation of the refractive index with the composition. The absorbance was modelled taking into account multiple reflections within the thin films and a non-absorbing substrate.<sup>24</sup> It should be noted that the direct band gap in  $\text{CuCrO}_2$  is responsible for the optical properties, whilst the indirect energy gap influences the electrical properties.<sup>54</sup> A Tauc plot was used to estimate the direct energy gap (Fig. 7c), and the estimated values are around 3.2 and 3.15 eV for the stoichiometric  $\text{CuCrO}_2$  and Cu-rich  $\text{CuCrO}_2$ , respectively, in agreement with the literature.<sup>12,25</sup> The lowest optical properties obtained when increasing  $\text{Cu}/(\text{Cu} + \text{Cr})$  suggests the stronger interaction between  $\text{Cu } 3d^{10}$  electrons responsible for the reduction of the energy gap and the overall transparency.<sup>55</sup> The formation of  $\text{Cu}_2\text{O}$  leads to an additional optical transition at around 2.5 eV, further reducing the energy gap, as observable for  $\text{Cu}_2\text{O} + \text{CuCrO}_2$  with a value of around 3.1 eV. An energy gap of around 2.6 eV was obtained for the pure  $\text{Cu}_2\text{O}$ , in agreement with previous studies on  $\text{Cu}_2\text{O}$  films deposited by CVD.<sup>56</sup> The presence of  $\text{CuO}$  would lead to the appearance of another optical transition at around 1.6 eV, not observable in our samples.

Despite thickness variations, the  $\text{FoM}^{\text{G}}$  value was calculated from eqn (1) to compare the properties of our films with the

literature; the results are reported in Fig. 7d. The  $\text{FoM}^{\text{G}}$  values from the literature were estimated with a 0% reflectivity when this value is not reported in the study. The AA-MOCVD Cu-rich  $\text{CuCrO}_2$ :65% films showed a promising  $\text{FoM}^{\text{G}}$  value of 2200  $\mu\text{S}$ , comparable with the value of 2300  $\mu\text{S}$ , the highest for this material, obtained by the Cu-deficient  $\text{CuCrO}_2$  films deposited by PI-MOCVD.<sup>24</sup> The AA-MOCVD growth method presents many advantages as it is performed at atmospheric pressure and at lower temperature than the PI-MOCVD method. Besides, the  $\text{Cu}_2\text{O} + \text{CuCrO}_2$ :67% composite films with higher conductivity achieved a good  $\text{FoM}^{\text{G}}$  value of 1400  $\mu\text{S}$ , despite a reduction in their optical performances.

## Conclusions

AA-MOCVD is a versatile and reproducible procedure working at a low temperature (350 °C) and atmospheric pressure, which enables the deposition of highly transparent and conductive  $\text{CuCrO}_2$ -based thin films.

The deposition of the out of stoichiometry  $\text{CuCrO}_2$  films without any detectable secondary phase is achieved up to a



composition of Cu/(Cu + Cr) of 65%. Single-phase Cu-rich CuCrO<sub>2</sub> films present a nanocolumnar microstructure with a resistivity value lower than 0.1 Ω cm, and a wide band gap of around 3.1–3.2 eV. The optimal composition was found for Cu-rich CuCrO<sub>2</sub>:65% with a resistivity value of 0.05 Ω cm and an average transmittance of 55% in the 400–800 nm range, resulting in a high p-type FoM<sup>G</sup> value of 2200 μS. The enhancement of p-type conductivity is attributed to the formation of the intrinsic defects with Cu, occupying the Cr vacancies and creating Cu antisite defects, and to the further improvement of the film crystallinity.

In summary, we highlight the successful deposition of the Cr-deficient CuCrO<sub>2</sub> by AA-MOCVD with a figure of merit comparable with the one of the Cu-deficient CuCrO<sub>2</sub> films. Besides, the growth in an oxygen-poor environment has been demonstrated to impede the formation of CuO, allowing the synthesis of nanocomposite films composed exclusively of Cu<sub>2</sub>O and CuCrO<sub>2</sub>. These films are characterized by a carrier mobility of 0.65 cm<sup>2</sup> V<sup>-1</sup> s<sup>-1</sup>, higher than the Cu-rich CuCrO<sub>2</sub> films. In this study, the lowest resistivity value of 0.02 Ω cm was achieved for the Cu<sub>2</sub>O + CuCrO<sub>2</sub> films with 67% of Cu. Despite a reduced average transmittance of 52%, the performance of the nanocomposite films led to a FoM<sup>G</sup> value of 1400 μS.

As revealed in the present work, the modulation of the optical and electrical properties, such as mobility, resistivity, total transmittance, and band gap, is achieved by tuning the stoichiometry of the films. This allowed us to synthesize two appealing candidates for hole transport layers in the thin film, organic, and perovskite solar cells, as well as for organic light-emitted diodes (OLEDs) and thin film transistors (TFTs) as these Cu-rich CuCrO<sub>2</sub> thin films can be incorporated into these devices.

## Conflicts of interest

There are no conflicts to declare.

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