

A tunable amorphous p-type ternary oxide system: The highly mismatched alloy of copper tin oxide

Patrick J. M. Isherwood, Keith T. Butler, Aron Walsh, and John M. Walls

Citation: [Journal of Applied Physics](#) **118**, 105702 (2015); doi: 10.1063/1.4929752

View online: <http://dx.doi.org/10.1063/1.4929752>

View Table of Contents: <http://scitation.aip.org/content/aip/journal/jap/118/10?ver=pdfcov>

Published by the [AIP Publishing](#)

Articles you may be interested in

[Atomic layer deposition of tin oxide and zinc tin oxide using tetraethyltin and ozone](#)

J. Vac. Sci. Technol. A **33**, 021517 (2015); 10.1116/1.4907562

[Enhancement of p-type mobility in tin monoxide by native defects](#)

Appl. Phys. Lett. **102**, 212105 (2013); 10.1063/1.4808382

[Highly conductive p-type amorphous oxides from low-temperature solution processing](#)

Appl. Phys. Lett. **101**, 132104 (2012); 10.1063/1.4754608

[Electrical properties and stability of p-type ZnO film enhanced by alloying with S and heavy doping of Cu](#)

Appl. Phys. Lett. **97**, 142101 (2010); 10.1063/1.3496038

[Effect of an applied voltage during annealing on the resistivity and transparency of the amorphous tin oxide films](#)

J. Vac. Sci. Technol. A **21**, 1923 (2003); 10.1116/1.1619415

The logo for AIP APL Photonics is displayed. It features the letters 'AIP' in a large, white, sans-serif font on the left, followed by a vertical orange bar and the words 'APL Photonics' in a smaller, white, sans-serif font on the right. The background is a dark red with a bright yellow sunburst effect in the upper right corner.

APL Photonics is pleased to announce
Benjamin Eggleton as its Editor-in-Chief



A tunable amorphous p-type ternary oxide system: The highly mismatched alloy of copper tin oxide

Patrick J. M. Isherwood,^{1,a)} Keith T. Butler,² Aron Walsh,² and John M. Walls¹

¹*CREST, School of Electronic, Electrical and Systems Engineering, Loughborough University, Loughborough, Leicestershire LE11 3TU, United Kingdom*

²*Centre for Sustainable Technologies and Department of Chemistry, University of Bath, Claverton Down, Bath BA2 7AY, United Kingdom*

(Received 30 May 2015; accepted 17 August 2015; published online 9 September 2015)

The approach of combining two mismatched materials to form an amorphous alloy was used to synthesise ternary oxides of CuO and SnO₂. These materials were analysed across a range of compositions, and the electronic structure was modelled using density functional theory. In contrast to the gradual reduction in optical band gap, the films show a sharp reduction in both transparency and electrical resistivity with copper contents greater than 50%. Simulations indicate that this change is caused by a transition from a dominant Sn 5s to Cu 3d contribution to the upper valence band. A corresponding decrease in energetic disorder results in increased charge percolation pathways: a “compositional mobility edge.” Contributions from Cu(II) sub band-gap states are responsible for the reduction in optical transparency. © 2015 AIP Publishing LLC.

[<http://dx.doi.org/10.1063/1.4929752>]

I. INTRODUCTION

Transparent amorphous oxide semiconductors (TAOS) have opened up a huge array of possibilities for transparent, flexible opto-electronics devices.^{1–4} These amorphous materials possess the fundamental advantages of low-cost synthesis and smooth, uniform grain-boundary free films, enabling production in large-scale roll-to-roll processes. In order to fully exploit the technological avenues opened up by n-type TAOS, it is necessary to produce p-type counterparts. p-type TAOS would provide access to the multitude of semiconductor heterojunction technologies that have revolutionised science and technology in the latter part of the 20th century, but in low-cost, easily manufactured and highly versatile forms.

The task of developing p-type amorphous oxides faces two fundamental challenges: (i) materials capable of stabilising electron holes and (ii) ensuring sufficiently high hole mobility. The former relies on ensuring sufficiently low binding energy valence band maxima (VBM), whilst the latter in part depends on the orbital character of the valence band itself. The search for effective TAOS is a major challenge in contemporary materials design, as evidenced by a wide array of experimental^{5–8} and theoretical^{9–12} studies on the topic.

The issue of conductivity is related to the fact that the VBM in most oxide materials comprised O 2p orbitals. Oxygen is a relatively small atom with a high electronegativity, and the result is that the valence band is typically localised (in real space), flat (in reciprocal space), and narrow in band width.^{9,13} These properties result in hole localisation and hence high hole effective masses.^{9,14} One strategy that has proved successful in crystalline p-type oxides is to employ metal cations with electronic orbitals at similar energy to oxygen 2p states. The resulting hybridisation widens the valence band and causes hole delocalisation.^{13–15}

Owing to a reduced hole effective mass, mobility increases along with the conductivity of the material.¹⁴ Copper (both Cu⁺ and Cu²⁺) and tin (particularly Sn²⁺) are among the cheapest and most widely available metals for this purpose.^{9,14,16}

The binary copper oxides Cu₂O and CuO are two of the most studied p-type metal oxides.^{17–19} Cu₂O was one of the first materials found to exhibit the photovoltaic effect, and was being investigated for use in photovoltaics prior to Si.^{20,21} As p-type transparent conducting oxides (TCOs), neither are ideal. TCOs should have an optical band gap of 3 eV or more, thereby ensuring reasonable transmission of the visible spectrum.^{13,15} The band gaps of both copper oxides are too small.^{13,16} Other examples of p-type oxides include SnO, NiO, Cr₂O₃, and CuAlO₂.^{16,22–24} The development of p-type oxides with robust conductivity and appropriate band gaps for TCO applications remains a challenge.

The technique of alloying two highly mismatched materials has been shown to enable tailoring of the band gap of the resulting substance, and has been successfully employed in the III–V semiconductors, such as (In,Ga)As or Ga(As,N).^{25–27} Amorphous materials provide several benefits when compared to crystalline materials. Due to the lack of long-range order, they are typically more flexible and so can be used with a wider range of substrates.²⁸ They are also commonly deposited at lower temperatures since heat is not required to encourage the formation of a crystal lattice.^{15,28} Forming an amorphous material from highly mismatched materials ensures a large kinetic barrier for recrystallisation and hence a more robust disordered state.

In this study, we explore the formation of a ternary oxide mixture of Cu and Sn. Due to the mismatch between the crystal structures of CuO, Cu₂O, SnO, and SnO₂, an amorphous ternary oxide is formed. The distinct electron configurations of the two cations—Cu(II) is 3d⁹, whilst Sn(II) is

^{a)}Electronic mail: P.J.M.Isherwood@lboro.ac.uk

$5s^2$ —allows us to investigate the effects of altering the composition on the position and structure of the band edge, both of which are critical for designing p-type TAOS. We identify a “compositional mobility edge,” where a sharp change in the conductivity and transparency of the material is observed with a small change in Cu:Sn ratio, despite the gradual change in the band gap. This finding is explained by comparison with density functional theory (DFT) calculations of the evolution of the electronic density of states (DOS) with composition. The ability to affect gradual change in certain material properties whilst causing much more rapid alteration in others through the application of mismatched alloying is an important principle for designing p-type TAOS.

II. METHODOLOGY

A. Experimental

Thin-film deposition was carried out using an AJA International Orion 8 HV sputter coater equipped with an AJA 600 series radio frequency (RF) power supply and an Advanced Energy MDX 500 DC power supply. Films were deposited by co-sputtering from pre-formed ceramic metal oxide targets at a pressure of 1 mTorr (0.133 Pa) in a mixed oxygen-argon atmosphere. Film composition was varied by altering the power supplied to each deposition magnetron, thereby varying the deposition rate for the target material. SnO_2 was deposited using the DC power supply, and applied power was varied from 12.5 W to 50 W. CuO was deposited using the RF power supply, and applied power was varied from 72 W to 180 W. Argon flow rate was maintained at 7 standard cubic centimetres per minute (SCCM). Two sets of films were deposited, one with a pure oxygen flow rate of 1 SCCM and a second with an oxygen flow rate of 2 SCCM, resulting in oxygen partial pressures of 0.017 Pa and 0.03 Pa, respectively.

Film characterisation involved measurement of transmission, film thickness, sheet resistance and Hall mobility, compositional and crystal structure analysis, and detailed observation of both film surfaces and in cross-section using SEM and TEM. Transmission data were used to calculate the film band gap using the Tauc method.²⁹ Transmission was measured using a Cary Varian 5000 spectrophotometer. Thickness was measured using an Ambios XP2 stylus profilometer, and Hall mobility measurements were carried out using an Ecopia HMS 3000 Hall effect device. Carrier type was confirmed by exploiting the Seebeck effect and measuring the output Voltage. A four-point probe was used to measure sheet resistance, and these data were used to verify the resistivities measured using the Hall effect. Crystal structure was examined using X-ray diffraction (XRD) using a Bruker D2 Phaser benchtop diffractometer equipped with a Cu- K_α X-ray gun and a LynxeyeTM detector. The beam slit was 1 mm wide, and the antiscatter plate was positioned 3 mm above the sample. SEM was carried out using a Carl Zeiss Leo 1530 VP field emission gun scanning electron microscope. Aperture size was 30 μm and the operating voltage was 5 kV. TEM was conducted using a FEI Tecnai F20 field emission gun transmission electron microscope equipped with a bright field detector at an operating voltage

of 200 kV. TEM samples were prepared by focused ion beam milling using a dual beam FEI Nova 600 Nanolab scanning electron microscope. A standard *in-situ* lift-out method was used to prepare cross-sectional samples. A thin platinum cover layer was deposited to define the sample surface and to homogenize the final sample thinning. Samples were thinned to 75 nm. Compositional analysis was carried out using a Thermo Scientific K-AlphaTM X-ray photoelectron spectrometer equipped with a monochromated Al- K_α X-ray gun ($h\nu = 1486.6\text{ eV}$), an EX06 ion source, and a 180° double focusing hemispherical 128 channel analyser. Measurements were run at a base pressure of 3×10^{-7} mBar. The survey scan range was 0 to 1350 eV with a pass energy of 200 eV. Step size was 1 eV, and data were collected over 10 scans, with 10 ms dwell time per step. High-resolution scans were run with a pass energy of 50 eV. Step size was 0.1 eV, and data were collected over five scans with 50 ms dwell time per step. Samples were subjected to a 75 s low power Ar+ etch (200 eV beam energy) prior to measurement so as to remove any adventitious surface contamination. Peaks were identified and fitted using the Thermo Advantage analysis suite, using Smart (a Shirley variant) background subtraction.

B. Computational

All DFT calculations were performed within periodic boundary conditions using the Vienna *Ab Initio* Simulation Package (VASP)³⁰ and the projector augmented wave formalism.^{31,32} To represent the electron density, a plane wave basis with a kinetic cut-off energy of 500 eV was used. We employed the PBESol and HSE06 exchange and correlation functionals.^{33,34}

Amorphous structures were obtained by following the “melt and quench” scheme using *ab initio* molecular dynamics. The starting point was a random configuration with a stoichiometry chosen to match the experimentally determined atomic composition³⁵ in a simulation cell set to a volume obtained by interpolation between the volumes of the respective binary phases. The atoms in the random configuration were packed according to cut-off ratios to avoid spurious interactions and their positions were determined by the RandomGenerator package.³⁶ The simulations were then run for 20 ps at 3000 K, with a timestep of 2 ps and the semi-local PBESol functional. The resultant configuration was then cooled by a simulated annealing procedure over 60 ps to 0 K. At 0 K, the structure was locally optimised with respect to both ionic positions and cell volume. Finally, the electronic structure of the systems obtained was calculated using the screened non-local HSE06 hybrid functional.³⁴ The importance of an accurate treatment of electron self-interaction (which we achieve by inclusion of Hartree-Fock exchange) has been shown for amorphous oxides.³⁷

The resulting amorphous structures are also analysed using the R.I.N.G.S. package³⁸ to identify bonding motifs and under-coordinated centers. These are particularly important in amorphous semiconductors as trap states,^{39–42} unlike crystalline materials where point⁴³ and extended defects⁴⁴ tend to dominate.

III. RESULTS AND DISCUSSION

A. Structural characteristics

Analysis of the XRD patterns confirms that all of the mixed oxide films are amorphous, with no peaks present. Several samples were tested over longer timescales with the same result (Figure 1). TEM cross-sections also indicate that these materials are amorphous, with no crystal structure being visible (Figure 1).

SEM analysis showed that the films exhibit a significant number of surface structures and surface topology (Figure 2). Whilst the cause of this is not known, it is possible that it is related to partial phase separation. Films deposited at high temperatures (above 200 °C) were found to exhibit large-scale phase separation which was visible without magnification (Figure 2). For the samples studied herein, the separation is minimal.

B. Optical and electronic properties

It was found that the addition of CuO ($E_g = 1.2$ eV) to SnO₂ ($E_g = 3.6$ eV) caused a significant reduction in the band gap of the resulting material. The shift in band gap with increasing copper content was found to be non-linear (Figure 3). The addition of more oxygen to the deposition environment was found to have no significant impact.

The change in band gap of an alloy can usually be described by a quadratic bowing parameter, following:

$$E_g(A_xC_{1-x}) = xE_g(A) + (1-x)E_g(C) - bx(1-x), \quad (1)$$

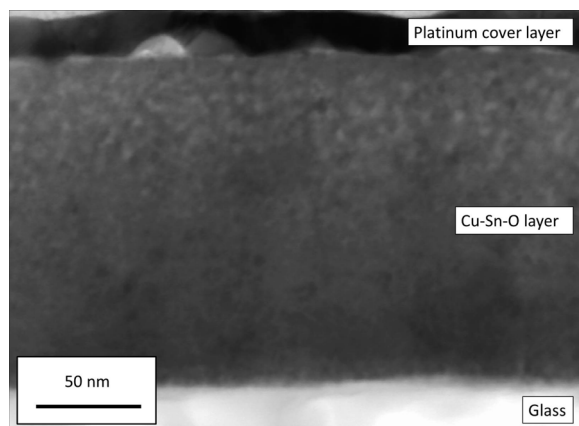
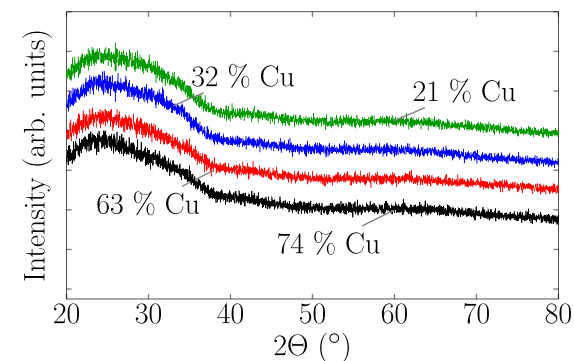


FIG. 1. (Upper) XRD patterns for thin-films with varying copper content. (Lower) TEM image of a Cu-Sn-O alloy film.

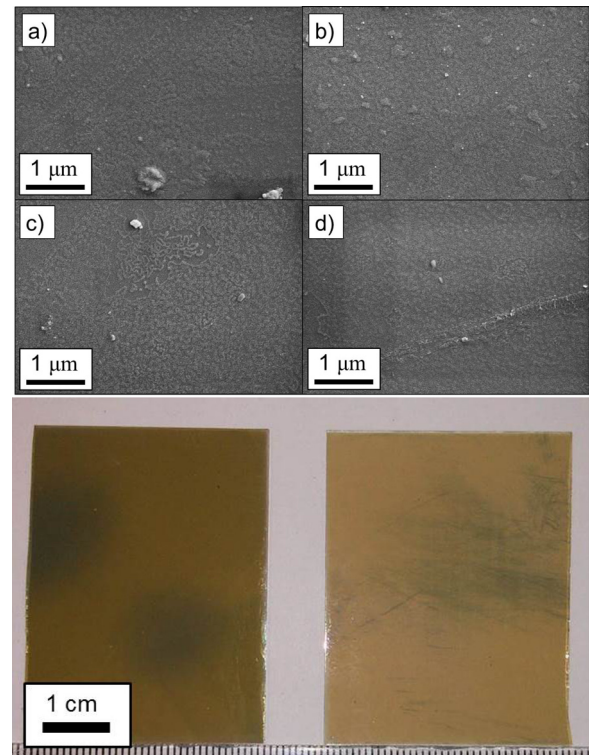


FIG. 2. (Upper) SEM photomicrographs of mixed copper tin oxide films. Films have copper contents of 74% (a), 63% (b), 32% (c), and 21% (d). (Lower) Films deposited at 300 °C showing discolouration indicative of large-scale phase separation.

where $E_g(A)$ and $E_g(C)$ are the band gaps of the two end-member compounds, x is the proportion of semiconductor A that is present in the alloy, and b is the bowing parameter. It was not possible to find a successful fit to measured data using the binary oxides as end member materials. By instead extrapolating the curve to the axes at either end of the sequence, it was possible to extract a compositionally independent bowing parameter of -1.5 eV. This large bowing is consistent with the mismatch between the charge state and

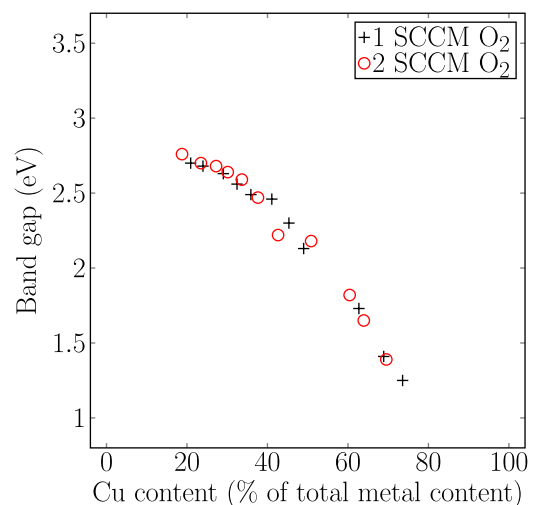


FIG. 3. Optical band gap against film copper content (as a percentage of total metal content) for both oxygen deposition environments.

structural preferences of Cu and Sn, and is typical of the values observed in other mismatched alloys.^{45,46}

Contrary to the bandgap trends, the averaged optical transmissions for each film did not show the same smooth change across the composition range. Instead, there is a gradual decline in transmission with increasing copper content up to a critical value of around 50% of the total metal content, beyond which the transmission drops rapidly (Figure 4). Low energy optical transitions can be associated with Cu(II) 3d⁹ centres.

Electrical resistivity was found to mirror the change shown by average transmission. Resistivity is roughly constant with increasing copper content up to the same critical point of around 50% of the total metal content, with further increases causing an exponential reduction in the resistivity (Figure 4). Unfortunately, it was not possible to obtain reliable carrier concentration and mobility data due to the limitations of the Hall system when measuring low mobility materials. Seebeck tests confirmed that all films showed p-type electrical behaviour.

C. Materials modelling

Pair correlation functions (PCFs) of the various samples show that the average Cu–O bond length is 2.5 Å, whereas the average for Sn–O bonds is 2.8 Å. By integrating over the first peak of the PCFs, we are able to obtain the average coordination number of various species present. This analysis shows that increasing Cu content generally leads to increased oxygen coordination. Under-coordinated species in amorphous semiconductors have been shown to be the cause of sub-gap states^{39,47} and are also detrimental to charge carrier lifetimes,⁴¹ therefore the enhanced coordination in Cu

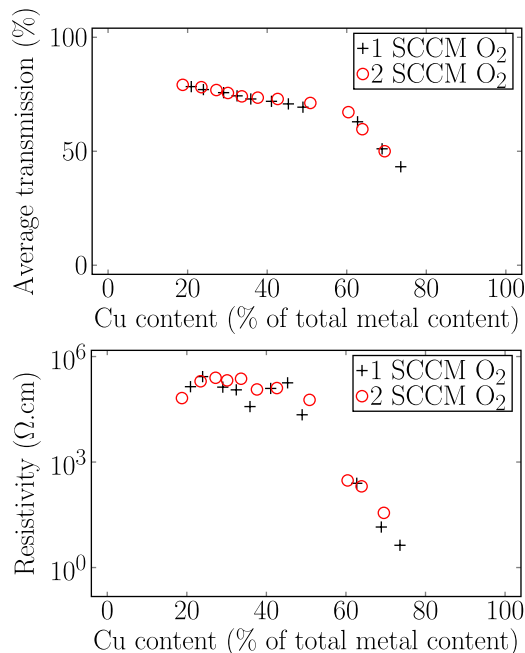


FIG. 4. (Upper) Average optical transmission against copper content (as a percentage of total metal content). (Lower) Film resistivity against film copper content.

rich samples should lead to decreased oxygen related sub-gap states.

The species resolved density of states (PDOS) is shown in Figure 5, which demonstrates the distinct electronic structure of the samples in Cu-rich and Sn-rich regions. The upper valence states become dominated by Cu 3d character in the Cu rich region as also confirmed in the electron density plots. In the Cu-rich samples, the edge of the density of states is steeper; however, there are noticeable gap states indicative of the empty d band of Cu(II). The presence of these gap states is consistent with the sharp change in rate of decrease in the transmission when Cu content is over 50%, with no equivalent change in bandgap. Sample absorption curves³⁵ have been shown to indicate the existence of sub-gap states.⁴⁸ For conclusive experimental evidence, the use of hard X-ray XPS would be necessary; however, this is beyond the scope of the current study.

To further understand the abrupt change in conductivity, when the Cu content is greater than 50%, we have further analysed the simulated amorphous phases in terms of disorder and fluctuations in the local Madelung potential (which largely determines the carrier energies in oxide systems⁴⁹) disorder. The site Madelung potential (V_M) is calculated by integrating the Hartree potential (V_H) over the ionic site

$$V_M = \int_V V_H dr, \quad (2)$$

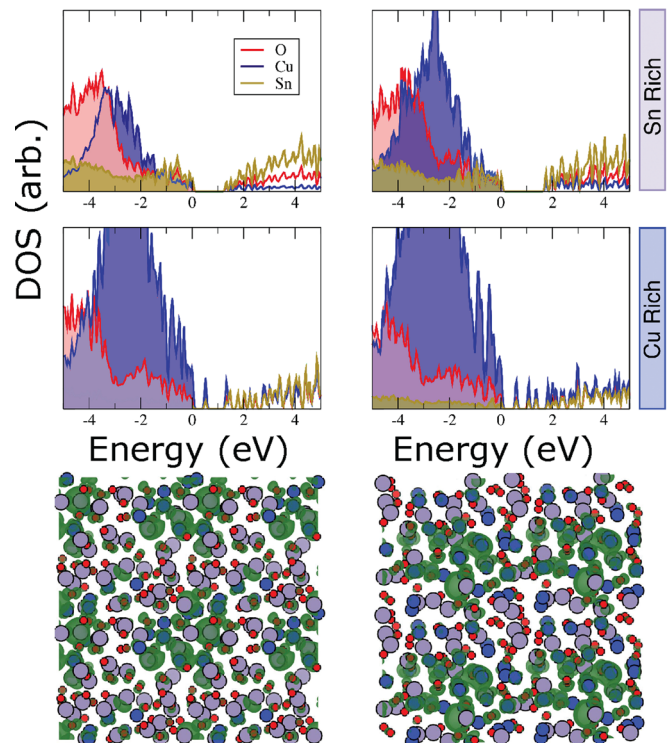


FIG. 5. (Upper) Electronic density of states and (lower) real space electronic distribution of the upper valence band of a Cu-rich and Sn-rich alloys. The stoichiometries of the samples for the DOS are for the Sn rich samples 20:74:105 (left) and 16:33:52 (right), and for the Cu rich sample 33:20:49 (left) and 74:20:105 (right). The density isosurface is plotted at $1 \times 10^{-3} \text{ e}\text{\AA}^{-3}$. Cu atoms are blue, Sn atoms are grey, and O atoms are red.

TABLE I. Results from DFT simulations showing the x:y:z is stoichiometry in $\text{Cu}_x\text{Sn}_y\text{O}_z$, the average metal to oxygen separation in Å (where M is the dominant metal cation), the average oxygen coordination number ($\langle O_{cn} \rangle$), the dominant metal contribution to the valence band maximum (VBM), and the standard deviation in the site Madelung potentials (ΔV_{Cu} and ΔV_{Sn} in V).

x:y:z	$d(\text{M}-\text{O})$	$\langle O_{cn} \rangle$	VBM	ΔV_{Cu}	ΔV_{Sn}
20:74:105	2.8	3.42	Sn	0.4	1.0
16:33:52	2.8	3.36	Sn	0.4	1.0
33:20:49	2.5	3.54	Cu	0.3	0.9
74:20:105	2.5	3.67	Cu	0.6	0.9

which we perform using the MacroDensity package.⁵⁰ This allows us to obtain the variation in the Madelung potential resolved by atomic site, as presented in Table I.

The p-type conductivity will relate to the cationic sites that define the top of the valence band, i.e., the Cu sites in Cu rich samples. Following the Anderson localisation model, if there is energetic disorder across these sites, then only a fraction is deemed “favourable” for conduction. Conduction is said not to occur if the difference in potential between sites exceeds γV . If the spread of the disorder is defined as W , then the fraction of sites participating in conduction pathways is⁵¹

$$p \approx \frac{2\gamma V}{W}. \quad (3)$$

There will be a percolation problem and the hole will become localised if p is less than a certain threshold.

For the copper tin oxides studied here, the variance in the Sn site Madelung potential is most pronounced and a spread in potential of ≈ 1 V will lead to poor conductivity (see Table I). The variance in the Cu site potential is almost half that of Sn (≈ 0.5 V). Therefore, the mobility edge of the alloy can be controlled with composition.

The concept of a compositional mobility edge explains the unusual simultaneous decrease in resistivity and optical transmission. At the same point where the band edge becomes dominated by Cu states, which offer a percolation path through the material for holes, the sub gap states associated with unfilled Cu orbitals also appear. If the sub-gap Cu d-bands are key to obtaining a low film resistivity,⁵² then it is extremely unlikely that a p-type Cu(II) oxide based material could be synthesised which is simultaneously optically transparent and semiconducting. However, the principles outlined here suggest further routes for the development of ternary (and higher order) conductive p-type amorphous materials. In particular, the existence of compositional edges for some properties (conductivity) and not for others (bandgap) demonstrates the ability to independently control such key physical properties.

IV. CONCLUSIONS

From a combined experimental and computational study of ternary Cu–Sn oxides, all alloy compositions were found to be amorphous. The optical band gap was found to decrease gradually with increasing copper content, and a bowing parameter of -1.5 eV was calculated. In contrast,

both optical transmission and electrical resistivity were found to show little change at lower Cu content. There is a sharp transition as the Cu content reaches 50%, with further increases causing dramatic reductions in both resistivity and transmission. Simulations of the amorphous phases from first-principles indicate that this transition occurs at the point where the metal ion defining the valence band edge switches from Sn to Cu. This results in reduced energetic disorder and an increase in the number of charge percolation pathways, a “compositional mobility edge.” Due to the reduction in optical transmission associated with unfilled Cu d bands forming sub-gap states, Cu(II)-based amorphous oxide alloys of this type are unlikely to be good candidates as amorphous transparent conducting oxides.

ACKNOWLEDGMENTS

The authors are grateful to RCUK for financial support through the SuperSolar Hub (EPSRC Grant No. EP/J017361/1). P.J.M.I. would like to acknowledge the invaluable help and support of Ali Abbas with structural characterisation. The research at Bath has been supported by the EPSRC (Grant Nos. EP/K016288/1 and EP/M009580/1). This work benefited from access to both the University of Bath’s High Performance Computing Facility and ARCHER, the UK’s national high-performance computing service, which is funded by the Office of Science and Technology through EPSRC’s High End Computing Programme (Grant No. EP/L000202).

- ¹E. Fortunato, P. Barquinha, and R. Martins, *Adv. Mater.* **24**, 2945 (2012).
- ²V. Pecunia, K. Banger, and H. Sirringhaus, *Adv. Electron. Mater.* **1**, 1400024 (2015).
- ³Y.-H. Kim, J.-S. Heo, T.-H. Kim, S. Park, M.-H. Yoon, J. Kim, M. S. Oh, G.-R. Yi, Y.-Y. Noh, and S. K. Park, *Nature* **489**, 128 (2012).
- ⁴J. F. Wager, *Science* **300**, 1245 (2003).
- ⁵N. Masó and A. R. West, *Chem. Mater.* **24**, 2127 (2012).
- ⁶U. A. Joshi and P. A. Muggard, *J. Phys. Chem. Lett.* **3**, 1577 (2012).
- ⁷A. J. Leenheer, J. D. Perkins, M. F. A. M. van Hest, J. J. Berry, R. P. O’Hayre, and D. S. Ginley, *Phys. Rev. B* **77**, 115215 (2008).
- ⁸J. Robertson and Y. Guo, *Appl. Phys. Lett.* **104**, 162102 (2014).
- ⁹G. Hautier, A. Miglio, G. Ceder, G.-M. Rignanese, and X. Gonze, *Nat. Commun.* **4**, 2292 (2013).
- ¹⁰J. B. Varley, V. Lordi, A. Miglio, and G. Hautier, *Phys. Rev. B* **90**, 045205 (2014).
- ¹¹H. Peng, A. Zakutayev, S. Lany, T. R. Paudel, M. D’Avezac, P. F. Ndione, J. D. Perkins, D. S. Ginley, A. R. Nagaraja, N. H. Perry, T. O. Mason, and A. Zunger, *Adv. Funct. Mater.* **23**, 5267 (2013).
- ¹²D. O. Scanlon, K. G. Godinho, B. J. Morgan, and G. W. Watson, *J. Chem. Phys.* **132**, 024707 (2010).
- ¹³H. Yanagi, S.-I. Inoue, K. Ueda, H. Kawazoe, H. Hosono, and N. Hamada, *J. Appl. Phys.* **88**, 4159 (2000).
- ¹⁴A. Kudo, H. Yanagi, H. Hosono, and H. Kawazoe, *Appl. Phys. Lett.* **73**, 220 (1998).
- ¹⁵*Handbook of Transparent Conductors*, edited by D. S. Ginley, H. Hosono, and D. C. Paine (Springer, London, 2010).
- ¹⁶J. A. Caraveo-Frescas, P. K. Nayak, H. A. Al-Jawhari, D. B. Granato, U. Schwingenschlögl, and H. N. Alshareef, *ACS Nano* **7**, 5160 (2013).
- ¹⁷J. W. Hodby, T. E. Jenkins, C. Schwab, H. Tamura, and D. Trivich, *J. Phys. C: Solid State Phys.* **9**, 1429 (1976).
- ¹⁸D. Scanlon, B. Morgan, G. Watson, and A. Walsh, *Phys. Rev. Lett.* **103**, 096405 (2009).
- ¹⁹T. Sander, C. T. Reindl, M. Giar, B. Eifert, M. Heinemann, C. Heiliger, and P. J. Klar, *Phys. Rev. B* **90**, 045203 (2014).
- ²⁰N. F. Mott, *Proc. R. Soc. London, Ser. A* **171**, 281 (1939).
- ²¹A. W. Copeland, O. D. Black, and A. B. Garrett, *Chem. Rev.* **31**, 177 (1942).

- ²²H. Kawazoe, M. Yasukawa, H. Hyodo, M. Kurita, H. Yanagi, and H. Hosono, *Nature* **389**, 939 (1997).
- ²³H. Sato, T. Minami, S. Takata, and Y. Yamada, *Thin Solid Films* **236**, 27 (1993).
- ²⁴E. Arca, K. Fleischer, and I. V. Shvets, *Appl. Phys. Lett.* **99**, 111910 (2011).
- ²⁵T. S. Kim, T. V. Cuong, C. S. Park, J. Y. Park, H. J. Lee, E. Suh, and C. Hong, *J. Korean Phys. Soc.* **43**, 273 (2003).
- ²⁶S. V. Novikov, C. R. Staddon, A. V. Akimov, R. P. Campion, N. Zainal, A. J. Kent, C. T. Foxon, C. H. Chen, K. M. Yu, and W. Walukiewicz, *J. Cryst. Growth* **311**, 3417 (2009).
- ²⁷C. Caetano, M. Marques, L. G. Ferreira, and L. K. Teles, *Appl. Phys. Lett.* **94**, 241914 (2009).
- ²⁸S. Narushima, H. Mizoguchi, K. I. Shimizu, K. Ueda, H. Ohta, M. Hirano, T. Kamiya, and H. Hosono, *Adv. Mater.* **15**, 1409 (2003).
- ²⁹J. Tauc, *Mater. Res. Bull.* **3**, 37 (1968).
- ³⁰G. Kresse, *Phys. Rev. B* **54**, 11169 (1996).
- ³¹P. E. Blöchl, *Phys. Rev. B* **50**, 17953 (1994).
- ³²G. Kresse, *Phys. Rev. B* **59**, 1758 (1999).
- ³³J. P. Perdew, A. Ruzsinszky, G. I. Csonka, O. A. Vydrov, G. E. Scuseria, L. A. Constantin, X. Zhou, and K. Burke, *Phys. Rev. Lett.* **100**, 136406 (2008).
- ³⁴A. V. Krukau, O. A. Vydrov, A. F. Izmaylov, and G. E. Scuseria, *J. Chem. Phys.* **125**, 224106 (2006).
- ³⁵See supplemental material at <http://dx.doi.org/10.1063/1.4929752> for further information on stoichiometries and oxidation state data used for model construction, and for representative absorption curves for deposited films.
- ³⁶See <https://github.com/keeto/AmorphousStructureGenerator> for keeto/AmorphousStructureGenerator (last accessed August 05, 2013).
- ³⁷W. Körner, D. F. Urban, D. M. Ramo, P. D. Bristowe, and C. Elsässer, *Phys. Rev. B* **90**, 195142 (2014).
- ³⁸S. L. Roux and P. Jund, *Comput. Mater. Sci.* **49**, 70 (2010).
- ³⁹S. Sallis, K. T. Butler, N. F. Quackenbush, D. S. Williams, M. Junda, D. A. Fischer, J. C. Woicik, N. J. Podraza, B. E. White, A. Walsh, and L. F. J. Piper, *Appl. Phys. Lett.* **104**, 232108 (2014).
- ⁴⁰W. Körner and C. Elsässer, *Thin Solid Films* **555**, 81 (2014).
- ⁴¹M. W. Lamers, K. T. Butler, J. H. Harding, and A. Weeber, *Sol. Energy Mater. Sol. Cells* **106**, 17 (2012).
- ⁴²K. T. Butler, M. P. W. E. Lamers, A. W. Weeber, and J. H. Harding, *J. Appl. Phys.* **110**, 124905 (2011).
- ⁴³K. G. Godinho, J. J. Carey, B. J. Morgan, D. O. Scanlon, and G. W. Watson, *J. Mater. Chem.* **20**, 1086 (2010).
- ⁴⁴S.-H. Yoo, K. T. Butler, A. Soon, A. Abbas, J. M. Walls, and A. Walsh, *Appl. Phys. Lett.* **105**, 062104 (2014).
- ⁴⁵Y. Z. Zhu, G. D. Chen, H. Ye, A. Walsh, C. Y. Moon, and S. H. Wei, *Phys. Rev. B* **77**, 245209 (2008).
- ⁴⁶N. Tit, I. M. Obaidat, and H. Alawadhi, *J. Phys.: Condens. Matter* **21**, 075802 (2009).
- ⁴⁷S. Sallis, N. F. Quackenbush, D. S. Williams, M. Senger, J. C. Woicik, B. E. White, and L. F. J. Piper, *Phys. Status Solidi A* **212**, 1471 (2015).
- ⁴⁸K. Nomura, T. Kamiya, H. Yanagi, E. Ikenaga, K. Yang, K. Kobayashi, M. Hirano, and H. Hosono, *Appl. Phys. Lett.* **92**, 202117 (2008).
- ⁴⁹A. Walsh and K. T. Butler, *Acc. Chem. Res.* **47**, 364 (2014).
- ⁵⁰See <https://github.com/WMD-Bath/MacroDensity> for WMD-Bath/MacroDensity (last accessed August 05, 2013).
- ⁵¹*Models of Disorder: The Theoretical Physics Of Homogeneously Disordered Systems*, edited by J. Ziman (Cambridge University Press, Cambridge, 1979).
- ⁵²Y. Peng, Z. Zhang, T. Viet Pham, Y. Zhao, P. Wu, and J. Wang, *J. Appl. Phys.* **111**, 103708 (2012).