



Latest directions in p-type transparent conductor design

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Transparent conducting materials (TCMs) are crucial in the operation of modern opto-electronic devices, combining the lucrative properties of optical transparency and electronic conductivity. More than ever we rely on display and touch screens, energy efficient windows and solar cells in our day to day lives. The market for transparent electronics is projected to surpass \$3.8 billion by 2026 as the automotive industry seek to incorporate pop-up displays into driver windshields, and the prospect of touch-enabled transparent displays challenges the traditional mouse and keyboard mode of computer operation. However, these new technologies rely heavily on the development of high performance p-type TCMs, a task that has posed a significant challenge to researchers for decades. This review will cover the basic theory and design principles of transparent conductors, followed by an overview of early p-type TCMs and their shortcomings. We discuss the impact of high-throughput screening studies on materials discovery and critically assess the family of p-type halide perovskites that emerged from these, ruling them as unsuitable candidates for high-performance applications. We find that phosphides, selenides, tellurides and halides are the most promising emerging materials, capable of achieving greater valence band dispersion than traditional oxides, and we discuss the challenges facing these more exotic systems. The smorgasbord of materials presented in this review should guide experimental and computational scientists alike in the next phase of p-type transparent conductor research.

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1 Introduction

The dual properties of optical transparency and electronic conductivity position TCMs as one of the most important technologies of the 21st century, with an ever-increasing number of devices requiring invisible electrode layers. In touch and display screens, a transparent electrode above a liquid crystal display is critical to allow emitted light to reach the consumer; in solar cells, incoming light must be able to pass through charge transport layers to reach the active photovoltaic absorber layer below, where electron-hole pairs can be generated; and in energy efficient windows, light must be able to pass through entirely in order for the window to retain its primary function. All of these applications rely overwhelmingly on n-type transparent conductors such as Sn-doped In_2O_3 (ITO) and F-doped SnO_2 (FTO), degenerately doped post-transition metal oxides with wide optical band gaps and conductivity on the order of 10^4 S cm^{-1} .^{1,2} In contrast, p-type TCMs are still in their infancy, with many materials failing to combine acceptable levels of optical transparency with conductivities that are seriously competitive with their n-type counterparts. Closing this gap between p- and n-type technologies is crucial in order to meet the ambitious market projections for the transparent electronics industry.³ Transparent electronics rely on a transparent p-n junction, which in turn depends on the development of a high performance p-type TCM.^{4,5}

There have been multiple recent reviews on the topic of p-type TCMs. Zhang *et al.* give an excellent description of the

necessary underlying physics and a comprehensive overview of early directions in p-type material research, such as Cu-based delafossites, oxychalcogenides and oxides.⁶ These “prototypical” materials are also reviewed in detail, mainly from an experimental perspective, by Zhang *et al.*⁷ Cao *et al.* set out to describe precisely how to improve p-type conductivity in materials through defect engineering in “Design Principles of p-Type Transparent Conductive Materials”, an important resource for understanding the role that formation energies, ionisation potentials and charge compensation mechanisms play in the electronic behaviour of p-type materials.⁸ Hu *et al.* published a similarly titled “Design strategy for p-type transparent conducting oxides”, offering a greater range of recent examples from the literature.⁹ Fioretti and Morales-Masis offer a perspective on disperse valence band materials for p-type applications, with an insightful discussion on some of the experimental challenges that come with moving away from oxide-based chemistries.¹⁰ Finally, Shi *et al.* offer a comprehensive overview of both p- and n-type wide bandgap oxide semiconductors, with a particular focus on optoelectronic devices and applications, an invaluable resource on emerging thin-film transistor devices and OLED technologies,¹¹ while Wang *et al.* focus more intimately on developments in the p-type semiconductor device field.⁵

The field of organic optoelectronics is a key area of research associated with TCM technology, but is often omitted from discussions where inorganic materials are present (certainly from most of the above, with the notable exception of

Table 1 Optoelectronic properties of p-type TCMs from the literature. Values with ^a are calculated from DFT using the HSE06 functional, ^busing GGA+U. *d* = film thickness, nm; *T* = transmission; E_g^{ind} = indirect band gap (calculated), eV; E_g^{opt} = optical band gap, eV; σ = conductivity, S cm^{-1} ; *n* = carrier concentration, cm^{-3} ; μ = mobility, $\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$. CuI* data represents best quality sample from ref. 80, in text data is the average from the study

Material	Deposition	<i>d</i>	<i>T</i> %	E_g^{ind}	E_g^{opt}	σ	<i>n</i>	μ	Ref.
CuAlO ₂	PLD	500	28	3.52 ^a	3.5, 4.08 ^a	1	1.3×10^{17}	10.4	31 and 41
CuCrO ₂	Sputtering	150	66	3.06 ^a	3.28, 3.75 ^a	0.01	3.4×10^{17}	2.2	43 and 45
Mg:N-doped CuCrO ₂	Sputtering	150	69	—	3.52	278	1.2×10^{21}	0.006	45
CuScO ₂	Sputtering	110	40	—	3.3	30	—	—	47
CuYO ₂	PVD	330	50	—	3.5	0.025	—	—	47
CuGaO ₂	PLD	500	80	—	3.6	0.063	1.7×10^{18}	0.23	48
CuInO ₂ p-type	PLD	170	50	—	3.9	0.0028	—	—	50
SrCu ₂ O ₂	PLD	150, 700–880	—, 80	1.6 ^b	3.3, 2.1	0.0039, 0.01	1×10^{13} , 1×10^{15}	—, 6.7	53, 81 and 82
Sr-doped LaCuOS	Sputtering	150	60	—	3.1	20	—	—	58
Mg-doped LaCuOSE	Sputtering	40	—	—	2.8, 2.72 ^a	910	1.7×10^{21}	3.5	62 and 63
La _{0.75} Y _{0.25} CuOS	Solvothermal	250	76	—	3.06	89.3	6.6×10^{20}	0.85	64
Cu ₂ O	Sputtering	1000	—	—	2.0, 2.12 ^a	38	1×10^{14}	256	65 and 66
Ni-doped Cr ₂ O ₃	PLD	—	40	—	3.3	28	—	—	69
SnO p-type	PLD, Sputtering	100, 15	—, 92	0.61 ^a	2.6, 2.65, 2.6 ^a	—	$1, 2.2 \times 10^{17}$	7, 18.71	18 and 71
NiO	Sputtering	100	70	—	—, 4.1 ^a	9	5×10^{17}	28	75 and 78
ZnCo ₂ O ₄	PLD	100–300	26	3.96 ^a	2.26	0.39	—	—	83 and 84
ZnRh ₂ O ₄	PLD	100–300	55	2.88 ^a	2.74	2.75	—	—	83 and 84
ZnIr ₂ O ₄	PLD	100–300	61	2.37 ^a	2.97	3.39	—	—	83 and 84
Sr-doped LaCrO ₃	MBE	50	43	—	4.6	56	7.5×10^{21}	0.04	85
K-doped Ba ₂ BiTaO ₆	PLD	120	90	—	4.5, 3.6 ^a	0.005	1×10^{14}	30	86 and 87
K-doped BaSnO ₃	PLD	100	—	—	—, 3.40 ^a	4.8×10^{-7}	1.0×10^{13}	0.30	88 and 89
Cs ₄ CdSb ₂ Cl ₁₂	Solvothermal	—	40	—	3.11 ^a , 3.30	2×10^{-6}	—	—	90 and 91
Bilayer TeO ₂	Eutectic melt	1.5	—	—	3.7, 3.62 ^a	316	1.4×10^{19}	141	92
CaCuP	Sintering	—	—	1.23 ^a	2.78, 2.71 ^a	2×10^{-3}	—	—	93
BP	CVD	—	—	2 ^a	4 ^a	2800	5×10^{19}	350, 900 ^a	94–96
CuI*	MBE	283	—	—	3.1 ^a	0.89	1.06×10^{18}	110	80 and 97
(Cu ₂ S ₂)(Sr ₃ Sc ₂ O ₅)	Sintering	—	—	—	3.1, 3.06 ^a	2.8, 2.0 ^a	1×10^{17} , 1×10^{18a}	150	40 and 98
(Cu ₂ S ₂)(Ca ₃ Al ₂ O ₅)	—	—	—	—	3.17 ^a	1767 ^a	1×10^{21a}	—	40
(Cu ₂ S ₂)(Ba ₃ Sc ₂ O ₅)	Sintering	—	—	—	3.04 ^a	3.24, 3.24 ^a	1×10^{21a}	—	40



Shi *et al.*)¹¹. Indeed, this vast topic deserves its own spotlight, and so the reader is directed to an incredibly comprehensive piece by Ostroverkhova for an overview of the underlying physics, materials and applications associated with organic optoelectronics,¹² and also to Lee *et al.* for a particular focus on flexible organic optoelectronic device applications.¹³

In this review, we aim to combine and condense many of the themes touched upon by existing reviews, while offering a fresh perspective on emerging materials, particularly those from high-throughput screening studies. We outline the basic requirements for a TCM in Section 2, providing a clear point of reference for the materials discussed in the later sections. We then offer an overview of the prototypical p-type TCMs in Section 3, spotlighting Cu-based delafossites, oxychalcogenides and oxides. Understanding the strengths and limitations of these materials spurred on a wider search for p-type TCMs, aided by the advent of large scale materials screening, with spinels, perovskites, phosphides and complex oxychalcogenides coming to the attention of researchers. These more exotic materials are analysed in Section 4 and represent the latest efforts in p-type TCM research. A summary of the properties of all the materials discussed can be found in Table 1. We then provide a short update on current p-type TCM applications in Section 5 before presenting some concluding thoughts in Section 6.

2 Design principles

2.1 Electronic structure requirements

2.1.1 A strict definition of the “band gap”. The fundamental gap is the energy required for the transition of an electron from the valence band maximum (VBM) to conduction band minimum (CBM). If the VBM and CBM are described by the same wave-vector k in reciprocal space, then this transition is direct, and an incoming photon with sufficient energy can excite an electron from the VBM to the CBM. If the VBM and CBM are described by different wave-vectors, then the transition is indirect. For an incoming photon to excite an electron across an indirect band

gap, a change in crystal momentum (provided by an internal phonon process – a photon is not capable of inducing this change itself) is required. As there is no guarantee that a phonon will be instantaneously available when a photon is absorbed, the intensity of indirect absorption is lower than that of direct absorption. These concepts are illustrated in Fig. 1a.

The optical gap is the energy of a transition measured from optical experimental techniques,^{14,15} and can be larger or smaller than the fundamental gap. From the perspective of a theorist, the optical gap is the first direct allowed transition, corresponding to the band gap observed experimentally. If there is an excitonic state within the band gap of a material, formed by the Coulombic attraction of an electron–hole pair, then an electron can be excited from the VBM to this lower energy state rather than the CBM, giving an optical gap that is smaller than the fundamental gap. Opposite behaviour can be observed when the transition from the VBM to CBM is forbidden due to Laporte selection rules. Forbidden transitions will experience very weak intensity, such as d–d transitions, whereas the first allowed transition will experience a strong absorption onset, causing the optical gap to be larger than the fundamental gap.¹⁶ Selection rules governing symmetry, parity and spin can all have an effect on determining the first allowed transition from occupied to unoccupied states, and the optical transition matrix can lend great insight into which transitions are allowed and forbidden.

Another phenomenon that can affect the magnitude of the optical band gap in transparent conductors is the Moss–Burstein shift, illustrated for an n-type transparent conductor in Fig. 1b. An electron donor will raise the Fermi level above the conduction band minimum, increasing the band gap by E^{MB} – this is well established in CdO.¹⁷

2.1.2 Band gap requirements for transparency. Armed with our definitions, we can now consider the requirements for the band gap of a transparent conductor. To ensure optical transparency, the optical band gap must be larger than 3.1 eV, corresponding to the maximum wavelength of visible photons. This has the implication that a direct fundamental gap with a symmetry forbidden transition can be less than 3.1 eV, as is the case of highly successful

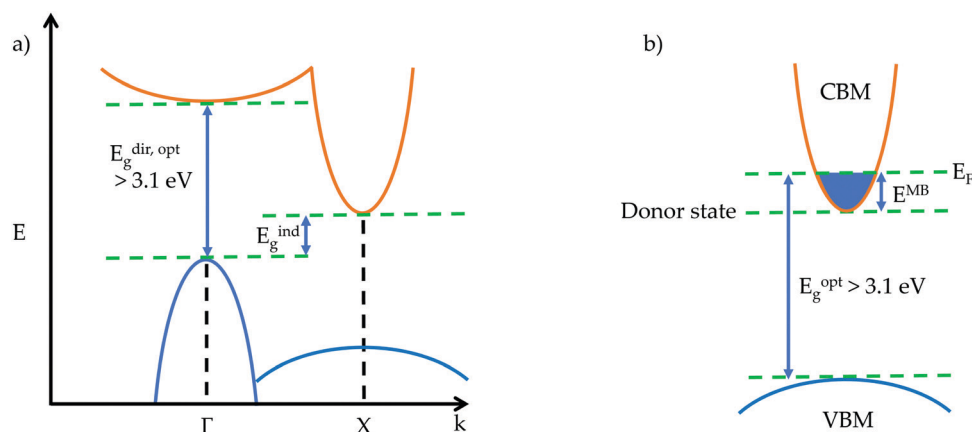


Fig. 1 Simple schematics of (a) a band structure with an indirect fundamental gap and direct optical gap, and (b) the widening of the optical gap through the Moss-Burstein shift in an n-type transparent conductor.



n-type TCM In_2O_3 , as long as the allowed transitions comprising the optical gap span an energy range greater than 3.1 eV.¹⁶ It also means that a material with a small indirect gap but larger optical gap (equal to or larger than the direct gap) can be transparent as the indirect absorption will only be weak in comparison, as is the case in the bipolar TCM SnO .¹⁸

Also important to transparency is the prevention of intra-band excitation. This means that for a p- (n-) type TCM, there should be an optical gap greater than 3.1 eV between the VBM (CBM) and the next occupied state below (unoccupied state above). This ensures that when charge carriers are added to the system in the form of defects, typically near or in the VBM (CBM), optical transitions to (from) these defect states from below (to above) still fulfil the transparency criteria.¹⁹

2.1.3 Conductivity mechanism in TCMs. Conductivity σ (S cm^{-1}) is governed by the equation:

$$\sigma = ne\mu, \quad (1)$$

where n is the charge carrier concentration (cm^{-3}), e the elementary charge (C) and μ the charge carrier mobility ($\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$). Intuitively, one would therefore aim to maximise both carrier concentration and charge carrier mobility in order to maximise conductivity. However, there is a precarious balancing act to maintain, as adding too many charge carriers (by introducing defects into the system) can reduce their overall mobility due to scattering processes such as ionised impurity scattering, carrier-carrier scattering, grain boundary scattering and both polar and non-polar electron-phonon scattering.^{20–22}

Charge carrier concentration is also intimately linked to transparency *via* the plasmon frequency ω_p^2 relationship, describing the oscillation of charge carriers within an applied field:

$$\omega_p^2 = \frac{ne^2}{m^* \epsilon_\infty \epsilon_0}, \quad (2)$$

where m^* is the charge carrier effective mass (kg), ϵ_∞ the high frequency dielectric constant and ϵ_0 the permittivity of free space. If the value of n is large enough (typically $> 5 \times 10^{21} \text{ cm}^{-3}$), the plasmon frequency will be shifted from the near infra-red (NIR) range into the visible, reducing transparency.⁶ Therefore, prioritising high charge carrier mobility over high charge carrier concentration is a viable strategy for obtaining high levels of conductivity.

Charge carrier mobility μ is controlled by the equation:

$$\mu = \frac{e\tau}{m^*}, \quad (3)$$

where τ is the average time between collisions (s). Therefore high mobility is dependent on large carrier lifetimes and having a low charge carrier effective mass. The effective mass describes the mass a charge carrier appears to have in response to the forces it experiences when moving through a lattice, and strictly is a tensor. More commonly, the effective mass is given with reference to a particular crystal direction, averaged over all three directions or with reference to the direction with the least favourable effective mass to give the “worst case scenario”.

Using a parabolic approximation to model the appropriate (valence or conduction) band edge:

$$E(k) = E_0 + \frac{\hbar^2 k^2}{2m^*}, \quad (4)$$

where E_0 is the eigenvalue of the band for that value of k and \hbar is the reduced Planck constant, the effective mass can be straightforwardly calculated from the second derivative of the band energy $E(k)$ with respect to the wave-vector k :

$$m^* = \frac{\hbar^2}{\frac{d^2 E(k)}{dk^2}}. \quad (5)$$

Qualitatively, the charge carrier effective mass can be assessed by the dispersion or curvature of the relevant band edge, with a greater curvature indicating a lower effective mass and therefore higher mobility.

The carrier lifetime is determined by scattering mechanisms concerning lattice phonons, charge carriers, grain boundaries and ionised impurities,²¹ and is therefore strongly dependent on the elements in the system, the crystal structure, and the quality of the sample, for example if it is a single crystal, thin film, pressed powder or has undergone nano-structuring.²³

2.1.4 Defects and dopability. To reach competitive levels of conductivity ($> 10^4 \text{ S cm}^{-1}$), TCMs must exhibit a high tolerance for defect formation, requiring charge carrier concentrations of at least 10^{19} cm^{-3} . Charge carriers are introduced by the formation of defects in a system, with acceptor defects generating holes at the VBM for p-type systems and donor defects providing electrons at the CBM for n-type systems. These defects can be either intrinsic, taking the form of vacancies, antisites and interstitials, or extrinsic as substitutions and interstitials. Their formation energies can be calculated under different simulated growth conditions using chemical potential limits, giving an indication of how prevalent a particular defect species will be. While several efforts have been made to screen material dopability, such as the branch-point energy (BPE) method,^{24,25} a cheap and reliable way to assess defect tolerance is yet to emerge, with the BPE method particularly poor at identifying p-type materials.²⁶ Instead, expensive, large-scale defect studies are required to gain an understanding of the defect chemistry of a potential TCM, a significant bottleneck in the search process.

The concentration of a defect n_d , and hence charge carriers, is ultimately determined by the free energy of formation, with the main component coming from enthalpy (vibrational entropy is assumed to be negligible here):

$$n_d \approx gN_{\text{site}} \exp\left(-\frac{\Delta H_f}{k_B T}\right), \quad (6)$$

where g is the degeneracy and N_{site} the number of sites in the host material, ΔH_f is the formation energy of the defect, and k_B and T are Boltzmann's constant and temperature respectively.²⁷ Therefore, the lower the formation energy of a defect, the higher its concentration in the host material. Crucially, the active charge state of the defect species should be stable either in or near the appropriate band edge. Depending on the energy



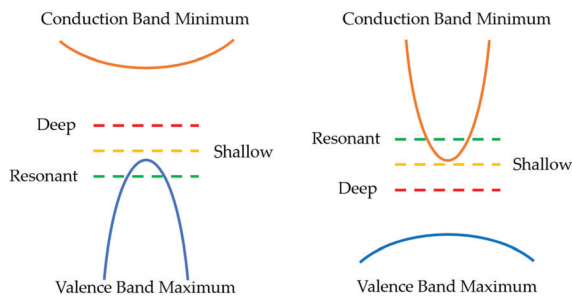


Fig. 2 Defect classifications in p- and n-type TCMs (left and right respectively).

gap between the defect state and band edge (Fig. 2), a defect can fall into one of three classes; *deep*, where the defect state lies within the band gap; *shallow*, where it is within $k_B T$ of the band edge, allowing conductivity *via* thermal activation at room temperature; and *resonant*, where the defect state lies within the relevant band edge. Many dopants, oxygen vacancies and adventitious hydrogen defects in n-type post-transition metal oxides fall into the *shallow* or *resonant* category, giving rise to high levels of conductivity,^{28–30} whereas in p-type TCMs *deep* acceptor defects dominate, severely limiting electronic performance.³¹ Engineering defect states closer to the VBM is critical to drive up the conductivity of p-type TCMs if they are to compete with their n-type counterparts.

For a more thorough discussion on the calculation of defects in crystalline systems and implementations of the necessary corrections for *ab initio* calculations, the reader is directed to the discussions by Lany and Zunger,^{32,33} Freysoldt *et al.*,³⁴ Kumagai *et al.*,^{35,36} and Walsh.²⁷

2.2 Sustainability requirements

In addition to the stringent electronic requirements of TCMs, there are other a handful of other factors to be considered when designing a new material. It will ideally consist of cheap, non-toxic, earth abundant elements to enable a secure, safe and sustainable manufacturing process. While ITO is a highly successful n-type TCM, the unstable supply and often wildly fluctuating price of raw indium is of concern,³⁷ feeding research efforts into seriously optimising existing indium-based technologies and developing indium-free alternatives.^{28,38} Furthermore, synthesis should be straightforward, easily scalable to provide incentive for industrial investment, the material will ideally integrate well into existing device infrastructures, have excellent air and moisture stability and long-term durability. A comprehensive overview of p-type TCM device design and performance, including thin film transistors and complementary metal oxide semiconductor devices, can be found in the work by Wang *et al.*⁵

3 Prototypical p-type TCMs

3.1 Chemical modulation of the valence band

The first p-type TCMs were designed around the electronic structure of the n-type transparent conducting oxides (TCOs). In n-type TCOs such as In_2O_3 , SnO_2 and ZnO , the valence band

is dominated by O 2p electronic states that display highly localised behaviour due to the large electronegativity of oxygen atoms.^{28,29,39} To engineer a p-type TCO, the localisation of the valence band needs to be modified such that holes can conduct throughout the material rather than being localised around the oxygen atoms. This can be done by introducing a closed shell cationic species with similar energy levels to O 2p states to facilitate the sharing of electron density, and for this they identified Cu^+ with electronic configuration $3d^{10}4s^0$. The crystal structure also plays an important role, with materials that enable maximum spatial overlap between Cu and O atoms achieving the greatest degree of covalency and in turn giving the most disperse valence band maxima, while maximising the distance between Cu atoms to prevent inter-band d–d transitions that could cause colourations and bring down the energy of the conduction band states.⁴⁰ These strategies were pioneered by Kawazoe *et al.*⁴¹ and have formed the basis of two decades of p-type TCM research.

3.2 Delafossites

The delafossite CuAlO_2 is the prototypical p-type TCO, highlighted as an exemplar material in the design strategies outlined by Kawazoe *et al.* It has an optical band gap of 3.5 eV and p-type conductivity of around 1 S cm^{-1} when nominally undoped.⁴¹ The crystal structure, shown in Fig. 3, consists of linear Cu–O–Al–O–Cu linkages, facilitating excellent spatial overlap between Cu and O atoms. Density functional theory (DFT) calculations using the hybrid functional HSE06 reveal an indirect band gap of 3.52 eV, matching well with the optical gaps reported in experiment, and good dispersion in the $\Gamma \rightarrow F$ direction. Defect calculations have shown that the lowest energy p-type defects in CuAlO_2 are Cu on Al substitutions (Cu_{Al}) and copper vacancies (V_{Cu}), which both contribute to the modest conductivity measured at room temperature,³¹ although both of these defect states are

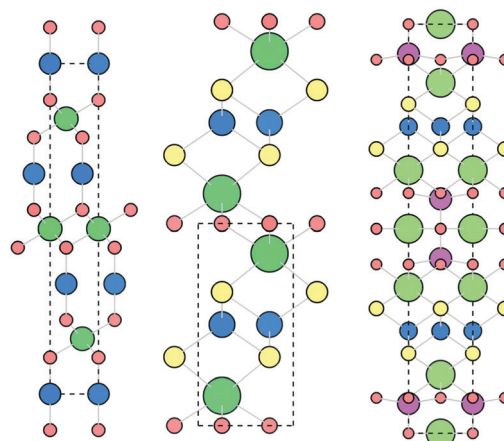


Fig. 3 The rhombohedral delafossite (left, CuMO_2 , Cu = blue, M = green, O = red), quaternary oxychalcogenide (centre, LnCuOCh , Ln = green, Cu = blue, O = red, Ch = yellow) and quinary oxychalcogenide (right, $(\text{Cu}_2\text{S}_2)(\text{A}_3\text{M}_2\text{O}_5)$, Cu = blue, Ch = yellow, A = green, M = pink, O = red, see Section 4.6) crystal structures. Viewed along *c* axis, visualised using VESTA.⁵⁶



deep acceptors. The holes generated by V_{Cu} are localised on the 6 surrounding Cu atoms, while there is a small distortion of the O–Cu–O dumbbell, indicative of a “reasonably localised polaron”. A small activation energy is required for the holes to overcome this interaction and migrate through the system, which places an intrinsic limit on the conductivity.³¹ While CuAlO_2 itself is not a high performance p-type TCM, it is the *first* TCM and embodies the design principles set out by Kawazoe *et al.*, kick-starting the research drive for Cu-based transparent conductors.

Substitution of the M^{3+} cation in the delafossite structure has been widely investigated in order to further reduce the localisation of the valence band and enable greater p-type conductivity. Cr,^{42–46} Sc,⁴⁷ Y,⁴⁷ Ga,^{48,49} and In have all been studied,⁵⁰ with CuCrO_2 achieving the best performance – p-type conductivity of 278 S cm^{-1} and an optical gap of 3.52 eV when co-doped as Mg:N-CuCrO_2 .⁴⁵ This is due to the large degree of covalency that is achieved in Cu–O–Cr–O–Cu linkages compared to the other M^{3+} cations, demonstrated through Bader charge analysis and partial electronic density of states calculations. Indeed, the charge transfer decreases in the order Cr, Sc, Y, Al, correlating with the reported experimental conductivities.⁵¹ This increased charge transfer has been observed separately using DFT in a series of $\text{CuAl}_{1-x}\text{Cr}_x\text{O}_2$ compounds, and distinct changes in the valence band are observed in X-ray photoelectron spectroscopy measurements of CuAlO_2 and CuCrO_2 .^{42,52}

Defect studies of CuCrO_2 have shown that the formation energy of substitutional Mg_{Cr} is around 0.9 eV lower than V_{Cu} , explaining the improved conductivity upon introduction of Mg.⁴³ The interstitial oxygen defect O_i is also 1.5 eV lower in energy in CuCrO_2 compared to CuAlO_2 , as the larger ionic radius of Cr generates more space in the structure, improving conductivity. It is suggested that N_{O} incorporation also generates charge carriers,⁴⁴ lowering the Fermi level and widening the band gap through a p-type Moss–Burstein shift, explaining the increased transparency compared to undoped films.^{45,46} The optical band gaps of CuGaO_2 and CuInO_2 are larger still, at 3.60 eV and 3.90 eV respectively.^{48–50} Due to its favourable band edge positions, CuInO_2 is capable of bipolar doping,^{49,50} but conductivity in both of these compounds still remains several orders of magnitude behind CuCrO_2 .

A study by Huda *et al.* rationalises the band gap trends for the group III delafossites.⁴⁹ For CuAlO_2 , CuGaO_2 and CuInO_2 (group III^a delafossites crystallising in the rhombohedral structure shown in Fig. 3), the fundamental band gaps are indirect and decrease down the group, but the optical gaps increase. This can be explained by the nature of the conduction band states at Γ , which are comprised of M s states – moving from Al to Ga to In, these states lower in energy, decreasing the fundamental band gap. Meanwhile, the conduction band states that correspond to the *direct* transition (metal p states) are, relatively, shallower, causing a slight rise in energy and explaining the increase in optical gaps. In the group III^b delafossites (Sc, Y and La, which crystallise in the slightly different hexagonal delafossite structure), the metal s states at the Γ point in the conduction band are not nearly as deep, and instead the partially filled, lower

energy d orbitals control the behaviour of the conduction band. Moving from Sc to Y to La sees a smaller relative change in the energy of the d states than moving from Al to Ga to In does in their s states, hence the difference in the fundamental and optical gaps is less pronounced in the group III^b delafossites.⁴⁹

SrCu_2O_2 crystallises in a delafossite related structure (tetragonal *I41/amd*, with trademark linear O–Cu–O dumbbell bonds among distorted SrO_6 octahedra), allowing good spatial overlap between Cu and O atoms. It has an optical gap of 3.3 eV from experiment due to direct transitions, in qualitative agreement with GGA calculations (a direct gap but with a smaller value of 2.1 eV due to the band gap under-estimation inherent in GGA DFT).⁵³ This is in contrast with most delafossites which have an *indirect* band gap. Conductivity originates from a thermally activated polaronic hopping mechanism, as in CuAlO_2 and CuCrO_2 , with cationic vacancies V_{Cu} and V_{Sr} having formation energies of around 1 eV each. Conductivity fails to surpass $10^{-2} \text{ S cm}^{-1}$, even when extrinsically doped with Ca or K,^{53–55} precluding SrCu_2O_2 from becoming a high performance material.

The delafossites are an important family of materials, and certainly deserve their place in the archives as the “original blueprint” for p-type TCM design. They benefit from relatively straightforward deposition and wide optical gaps, but are inherently flawed due to the deep acceptor levels present and subsequent polaronic conductivity mechanisms.

3.3 Oxychalcogenides

With the general formula LnCuOCh ($\text{Ln} = \text{La, Y, Pr, Nd, etc.}$ and $\text{Ch} = \text{S, Se, Te}$), Cu-oxychalcogenides possess a structural motif that enables p-type conductivity and optical transparency (Fig. 3). With alternating layers of $[\text{Ln}_2\text{O}_2]^{2+}$ and $[\text{Cu}_2\text{Ch}_2]^{2-}$, oxychalcogenides combine the wide band gap properties of metal oxides with the excellent orbital mixing of Cu d and Ch p states, while maintaining a spatial separation between the two sub-structures. This allows for high hole mobility, as the oxygen 2p states are spatially and energetically separated from the valence band orbitals of the $[\text{Cu}_2\text{Ch}_2]^{2-}$ layers.

The first oxychalcogenide p-type transparent conductor was LaCuOS ,⁵⁷ with an optical band gap of 3.1 eV and conductivity of around 20 S cm^{-1} upon Sr-doping.^{58,59} The highly electro-positive La raises the energy of the conduction states, giving a wide band gap, while holes generated by Cu vacancies are confined to the conduction pathways in the Cu–S layers.⁵⁷

Oxychalcogenides offer a rich search space for new TCs as they are open to a large variation in chemical composition. By varying the Ln site from La to Pr to Nd (in LnCuOS), the optical band gap changes from 3.14 eV to 3.03 eV to 2.98 eV. This is due to the lanthanide contraction, decreasing bond lengths in the *a*-axis, altering the Cu–Cu bond distance, increasing Cu 4s anti-bonding state interaction in the conduction band and lowering the band gap.⁶⁰ Altering the Ch anion from S to Se to Te (across LaCuOCh) also sees a decrease in the band gap from 3.14 eV to 2.80 eV to 2.40 eV, but not because of the change in ionic radii. Instead, the relative energy of the Ch orbital at the valence band increases, closing the band gap. The overlap between Cu 3d and



Ch np improves as the Ch gets larger, resulting in improved hole mobility down the group.^{60,61}

Of these variations, LaCuOSe has been widely studied due to its high p-type conductivity, achieving 910 S cm^{-1} at carrier concentrations of 10^{21} cm^{-3} when doped with Mg.⁶² This level of conductivity is competitive with n-type oxides, but with an optical band gap of 2.8 eV LaCuOSe is not fully transparent. Hybrid DFT calculations place the formation energy of V_{Cu} at around 1.5 eV under Cu-poor growth conditions, and the transition level occurs around 0.1 eV above the VBM hence the high conductivity. The Mg_{La} defect occurs at a slightly higher formation energy of 2.4 eV, but it is proposed that Sr and Ca can act as low energy acceptors that could improve the conductivity even further.⁶³ However, optical transitions from the V_{Cu} defect prevent a Moss–Burstein shift that would open up the band gap, so the transparency of LaCuOSe is permanently compromised.

The mixed cation system $(\text{La}_{1-x}\text{Y}_x\text{CuOS})$ (with $x = 0 \rightarrow 0.25$) holds the record for the highest experimental conductivity of an oxychalcogenide with a band gap greater than 3 eV – 89.3 S cm^{-1} and 3.06 eV respectively.⁶⁴ XPS and ICP (inductive coupled plasma) compositional analysis reveals a 14% Cu deficiency, suggesting that V_{Cu} is responsible for the conductivity. This is likely true in light of the defect landscape of pure LaCuOS, where V_{Cu} formation is favourable. The best films are highly crystalline, but are still orders of magnitude smaller than the required levels of conductivity for high performance TCs.

3.4 Oxides

As well as ternary and quaternary oxides, a handful of binary oxides have influenced the direction of p-type TC research, and deserve an honourable mention. Cuprous oxide, Cu_2O , has an exemplar valence band for p-type TCMs, where the Cu 3d and O 2p orbitals interact to such an extent that hole mobility of $256 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ has been observed experimentally.⁶⁵ However, Cu atoms are closer together in Cu_2O than in the CuAlO_2 crystal structure, giving greater overlap between Cu 4s states in the conduction band, bringing the energy down and ultimately reducing the fundamental band gap to 2.17 eV.⁶⁶ A disallowed transition from VBM to CBM results in an optical gap of 2.5 eV, but this still falls short of the transparency requirement, and deep acceptor levels are also a disadvantage, imposing an upper limit on achievable conductivity.⁶⁶ Nonetheless, Cu_2O has seen reasonable success, certainly as a proof-of-concept, as a p-type thin-film transistor (TFT).⁶⁷

Cr_2O_3 is an antiferromagnetic insulator with an optical band gap of around 3.4 eV.⁶⁸ Strong electron correlation in Cr_2O_3 results in a conduction band of 3d character, while the valence band is made up of Cr 3d and O 2p orbitals. This electronic structure allows p-type doping, capable of reaching conductivity of 28 S cm^{-1} when doped with Ni.⁶⁹ DFT calculations show that Mg-doping is also possible with relatively low formation energy (0.9 eV), however the acceptor level is quite deep, so conductivity levels are unlikely to surpass that of Ni-doped Cr_2O_3 .⁷⁰

SnO is a curious transparent conductor with a fundamental indirect band gap of 0.7 eV but an optical band gap of 2.6 eV

(Fig. 4), resulting in thin films that display a yellow hue.¹⁸ Films deposited by magnetron sputtering are capable of reaching a maximum hole mobility of $18.7 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ at carrier concentrations of $2.2 \times 10^{22} \text{ cm}^{-3}$.⁷¹ However, the most interesting feature of SnO is that it can act as a bipolar transparent conductor and form a homogeneous p–n junction.⁷² This is precisely due to the nature of the band structure and the forbidden fundamental band gap, with the relatively high energy valence band states resulting in a small ionisation potential that is conducive to p-type conductivity, and the relatively low energy conduction band states resulting in a large electron affinity that allows n-type conductivity, all while remaining (semi-)transparent.¹⁸ SnO remains an interesting candidate for bipolar transparent semiconductors, but the yellowish tint, as well as the undesirable tendency to oxidise to SnO_2 , pose significant obstacles to its widespread application.

NiO is a popular p-type semiconductor with a large optical band gap, often reported in excess of 3.4 eV, and with transmissivity in the range of 60–90% depending on film deposition method.^{6,73–75} Conductivity is modest at best, with several experimental reports failing to synthesise films surpassing 10 S cm^{-1} ,^{idref="cit74 cit75">74,75} although hole mobility of around $28 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ has been achieved through oxygen assisted sputtering (at the cost of transparency). Doping the Ni site with various alkali metals, in particular Li, has been reported to improve conductivity.^{76,77} However, a consistent theoretical model has yet to be agreed upon, with further work required to understand the nature of the hole localisation in both doped and non-doped NiO.^{11,78} While the electronic properties of NiO are not ground-breaking, it is an incredibly attractive TCM due to its simple rock-salt structure.⁷⁹ This allows it to interface straightforwardly with other cubic materials, such as successful n-type transparent conductors like Sn-doped In_2O_3 , leading to major efficiency gains in perovskite solar devices as an interfacial anode layer,⁷³ and as a result is one of the most widely implemented commercial p-type transparent electrodes.⁶

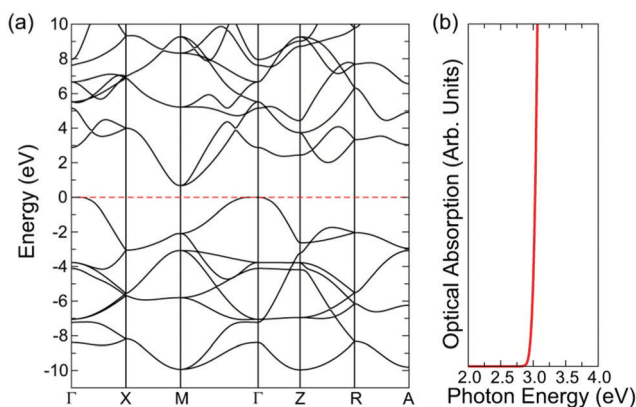


Fig. 4 Band structure and optical absorption spectrum of SnO, calculated with the PBE0–vdW functional. Reprinted with permission from Quackenbush *et al.*¹⁸ Copyright 2021 American Chemical Society.



4 Emerging p-type TCMs

4.1 Materials screening

High throughput computational screening has been an invaluable tool in the search for new p-type TCMs. Databases containing thousands of crystal structures and electronic structure data (for example the Materials Project)⁹⁹ can be queried with sophisticated search parameters, assessing swathes of materials for potential p-type TCM properties. Brunin *et al.* provide a thorough overview and explanation of the processes involved, discussing each stage of the search funnel,²⁶ and an assessment of how well high throughput descriptors actually perform can be found by Woods-Robinson *et al.*¹⁰⁰ The quality of the data in such databases is ultimately the limiting factor in the efficacy of the screening process – for example, older entries in crystallographic databases may not have been verified for any number of decades. For electronic structure databases, it is simply not possible to assess every single entry at a high level of theory – in fact, most entries are computed with GGA exchange correlation functionals, well known to underestimate band gaps and perform poorly for certain systems, so results from screening will often need to be validated by further independent research, and screening parameters themselves will often have to be compromised to include results that would otherwise be screened out (for example, if searching by GGA band gap, one might choose a minimum of 1 eV rather than 3 eV as GGA band gaps are underestimated by around 50%). The focus of this section will be on innovative directions in p-type TCM research, with many of the materials featured having been discovered in high throughput screening studies.

4.2 d⁶ Spinel oxides

The electronic configuration of the Cu(I) ion, 3d¹⁰, is well suited for mixing with oxygen 2p states and increasing the dispersion of the valence band, but there are other “quasi-closed shell” configurations that can also act as a solid foundation for p-type oxides. The d⁶ spinel oxides are excellent examples of this, where for low spin systems in an octahedral field they are essentially closed shell (the three t_{2g} orbitals are fully occupied). The filled metal t_{2g} orbitals interact with the O 2p states at the VBM, while the unoccupied e_g states contribute to the conduction band, similar to the splitting of the d orbitals in antiferromagnetic Cr₂O₃.⁶⁸ ZnM₂O₄ (M = Co, Rh, Ir) have indirect band gaps of 3.96 eV, 2.88 eV and 2.37 eV respectively, calculated with the HSE06 functional,⁸⁴ and relatively low formation energy of the Zn_M defect is reported for each system. The Co variant is the most studied experimentally, achieving conductivity levels of 21 S cm⁻¹, and can also be doped n-type to deposit a p–n heterojunction.^{6,101} While an intuitive example of materials design, it is unlikely that the d⁶ spinels will achieve record performances as the valence bands remain quite flat, and conductivity originates from a polaronic hopping mechanism. Furthermore, there have been conflicting experimental reports regarding the trends in conductivity and band gaps in these systems,^{6,83} indicating that further understanding is required.

4.3 p-Type perovskites – the next generation?

Sr-doped LaCrO₃ is another take on the quasi-closed shell configuration design strategy, where half-filled t_{2g} orbitals comprise the valence band. A forbidden transition from t_{2g}³ → e_g⁰ (2.8 eV) opens up the next available transition as the optical gap at 4.6 eV, resulting in excellent levels of transparency.⁸⁵ Doping to 50% Sr on the La site, arguably now forming a mixed cation perovskite, results in conductivity of 50 S cm⁻¹, but at the cost of a 50% loss in transmission. Despite this, Sr-doped LaCrO₃ remains of interest due to its perovskite structure, which could interface well with other perovskite based TCMs, such as n-type La-doped BaSnO₃.

The rhombohedral double perovskite Ba₂BiTaO₆ was identified through a high throughput screening study as a wide band gap oxide with good hole mobility.^{26,86} The filled Bi 6s² valence shell hybridises with the O 2p states at the VBM, while the Ta forms high energy conduction band states to ensure a large band gap of around 3.6 eV calculated with hybrid DFT (3.8 eV with GW theory), although from experiment this is even larger at 4.5 eV.⁸⁶ When doped with K on the Ba site, Ba₂BiTaO₆ displayed hole mobility of 30 cm² V⁻¹ s⁻¹, but extremely low carrier concentration (10¹⁴ cm⁻³) and therefore low conductivity. This is due to the charge compensating n-type defects that form, such as V_O and Ta_{Bi},⁸⁷ which ultimately prevent Ba₂BiTaO₆ from achieving competitive levels of conductivity.

Doping with K onto Ba sites has also been reported in the traditionally n-type material BaSnO₃.⁸⁸ A heterojunction was created with p-type K-doped and n-type La-doped BaSnO₃, demonstrating stability after high bias and thermal cycling. The p-type films displayed mobility of 0.30 cm² V⁻¹ s⁻¹ at a carrier concentration of 10¹³ cm⁻³, giving low overall conductivity. An activation energy for K dopants is reported to be 0.5 eV, but there is no support from DFT calculations or theory otherwise at the time of writing.⁸⁸ Given the low carrier concentration, it is likely that the intrinsic n-type defects in BaSnO₃, such as oxygen vacancies, or even adventitious hydrogen, will dominate in this system even under cation-poor conditions,⁸⁹ and high performance p-type activity is unfavourable.

Caesium based halide double perovskites, with the general formula Cs₄M²⁺B₂³⁺X₁₂^{VII} (M = Mg, Ca, Sr, Zn, Cd, and Sn(II), B = Sb, In and Bi, X = Cl, Br and I), have gained much interest in recent years as potential p-type TCMs, after a screening study identified 7 from a possible 54 elemental combinations that displayed thermodynamic stability, a band gap greater than 2.8 eV, a hole effective mass lighter than 1.2m₀ and good intrinsic p-type conductivity.⁹⁰ Cs₄CdSb₂Cl₁₂ is chosen as the most promising candidate, with delocalised Sb 5s orbitals mixing with Cl p states at the VBM suggesting good p-type performance, and defect calculations predicting intrinsic conductivity as a result of shallow Cd_{Sb} acceptor levels with low formation energy. However, Cs₄CdSb₂Cl₁₂ and Cs₄CdBi₂Cl₁₂ have been shown to be the only two of the predicted seven that are thermodynamically stable and synthesisable from a solvothermal method, and that they are in fact transparent insulators rather than conductors.⁹¹ Hybrid DFT calculations show that the band gap is around 3.3 eV, and that the valence band actually lies lower in energy than the initial GGA



calculations from the previous study suggested. The initial GGA calculations therefore predicted unrealistically facile p-type doping. Furthermore, upon inclusion of all stable competing phases of $\text{Cs}_4\text{CdSb}_2\text{Cl}_{12}$, it is found that the chemical potential limits alter the defect landscape significantly, to the point where Cd_{sb} is fully charge compensated by various native n-type defects, resulting in a hole concentration that is too far below the limit measurable for Hall techniques (Fig. 5).⁹¹ This quite firmly rules out caesium based halide double perovskites from becoming successful p-type TCMs, and demonstrates the importance of rigorously calculating chemical potential limits for defect analysis, as well as capturing an accurate description of band edges and gaps.

A separate high throughput study highlighted another set of double halide perovskites, starting with over 16 000 Cs- and Rb-based possible compounds, searching for thermodynamic stability, a GGA band gap greater than 1.8 eV and hole effective mass less than $1m_0$.¹⁰² They report 17 double perovskites, and one ternary perovskite CsPbF_3 , with (meta) stability and band gaps predicted to be greater than 3 eV with the hybrid functional HSE06, although 7 of these contain heavy and or toxic elements. This narrows the band gap upon consideration of relativistic effects (using spin orbit coupling calculations) and also renders them unsuitable for every day use. The relatively low hole effective masses in the remaining candidate materials, such as $\text{Rb}_2\text{AgBiCl}_6$ and $\text{Cs}_2\text{InLaBr}_6$ (0.49 and $0.59m_0$ respectively), are a result of halogen p and metal s states mixing well at the valence band. However the authors recognise that the ultimate obstacle for these materials is their p-type dopability, which in this case has not been studied computationally, and there is a call to arms for experimental investigation.¹⁰² Hu *et al.* have already demonstrated that it is not possible to generate sufficient p-type carriers in $\text{Cs}_4\text{CdSb}_2\text{Cl}_{12}$ due to the spontaneous formation of native n-type defects,⁹¹ which could be the downfall of these Cs and Rb double halide perovskites too.

4.4 Binary oxides

Achieving p-type doping in materials such as In_2O_3 , SnO_2 and ZnO has been a goal of materials researchers for decades, as this would enable an efficient homogeneous p-n junction and seamless integration with existing device architecture. Indeed, there have been occasional experimental reports of p-type conductivity in native n-type materials,^{88,103–107} but these results are often transient and difficult to reproduce. This is because these post-transition metal oxides are natively n-type, with “hole-killing” defects forming under all synthesis conditions. Multiple hybrid DFT studies have gone on to prove this quantitatively, showing that the formation of acceptor defects in these materials is thermodynamically unstable.^{28,30,89,108–111} One possible strategy for achieving p-type conductivity in such systems is to significantly alter the make-up of the valence band, for example through mixed cation alloying.

With an ultra-wide band gap of 4.9 eV,^{112–114} Ga_2O_3 is a prospective material for the power electronics industry due to its large breakdown voltage and improved band gap over rival materials GaN and SiC. Significant alloying of Ga_2O_3 with Bi_2O_3 is a promising route to engineering p-type behaviour in this natively n-type system.¹¹⁵ The formation of an intermediary valence band of Bi s character at around 1.6 eV above the valence band of the host material allows mixing with oxygen 2p states to improve band dispersion, while maintaining optical transparency.¹¹⁵ Mg, Zn and Cu are investigated as potential extrinsic dopants in both Ga_2O_3 and the Bi- Ga_2O_3 alloy, with Mg_{Ga} and Zn_{Ga} calculated to have relatively low formation energies, but deep transition levels. This is due to the wavefunction of the acceptor state coupling to the localised hole state on an oxygen in the valence band, regardless of the position of the valence band. The Cu_{Ga} defect does not couple to a valence band oxygen, and so even though it is a deep acceptor in Ga_2O_3 , it is shallow in the alloyed system now that the valence band has shifted up in energy, and the holes are localised on the Cu impurity site.¹¹⁵ This is an intriguing new



Fig. 5 Schematic of HSE and GGA band gap calculations and effects of miscalculated chemical potentials (left), transition level diagram calculated with HSE under the most favourable p-type conditions, showing n-type charge compensation (right). Reprinted with permission from Hu *et al.*⁹¹ Copyright 2021 Advanced Functional Materials, John Wiley and Sons.



approach to engineering dispersion and acceptor levels, and such alloying could be applied to other materials with strongly correlated valence (or indeed conduction) bands. However, the localisation of the holes around the Cu defect site suggests that mobility may still not reach the levels required for use on a commercial scale.

β -TeO₂ has been recently proposed as a wide band gap semiconductor due to its direct band gap of 3.70 eV in monolayer form.^{92,116} Hybrid DFT calculations show that the valence band is comprised mainly of Te 5p and O 2p, with smaller contributions from Te 5s orbitals, and that there is high, anisotropic mobility of both electrons and holes, with the latter predicted to reach mobility of 9100 cm² V⁻¹ s⁻¹ in the $\Gamma \rightarrow X$ direction for the monolayer system.¹¹⁶ It was calculated to have a low cleavage energy similar to black phosphorus, so exfoliation is expected to be relatively straightforward. Indeed, synthesis of β -TeO₂ bilayers has been successful from a eutectic melt (Te:Se 5%:95%) droplet that is rolled on a silicon wafer using a van der Waals printing process.⁹² Both EELS and XPS measurement of the valence band confirmed a band gap of 3.7 eV, the hole effective mass was determined to be 0.51 m_0 from scanning tunnelling spectroscopy data, in good agreement with the theoretical value, and the bilayers performed well when deposited in field effect transistors. DFT calculations on the *bilayer* reveal hole mobility values of 436 cm² and 7690 V⁻¹ s⁻¹ in the *a* and *b* directions respectively, due to the different conduction pathways on these axes. Hall measurements of the mobility gave an average value of 141 cm² V⁻¹ s⁻¹.⁹² The p-type dopability remains to be tested, with the possibility of charge compensating native defects posing the biggest challenge, and there is of course the problem of Te abundance, but if a large hole concentration can be engineered in β -TeO₂ then it could be an exciting direction for future research.

4.5 Beyond oxides for valence band dispersion

An alternative approach to engineering a dispersive valence band is to move away from oxides and instead encourage cation orbital mixing with the 3p states of phosphorus. With a higher principal quantum number, phosphorus valence orbitals are more diffuse than those of oxygen. Ternary phosphides MCuP where M = Mg, Ca, Sr and Ba have been investigated using hybrid DFT, with excellent mixing of Cu 3d and P 3p states reported for each compound (42:54 for MgCuP and 38(9):48 for CaCuP, SrCuP and BaCuP) resulting in extremely low hole effective masses in the *a* and *b* directions for the trigonal planar coordinated Cu–P systems CaCuP, SrCuP and BaCuP.⁹³ This is another example of how the synergistic crystal-electronic structure relationship is key for engineering p-type TCMs – the smaller ionic radius of Mg forces the Cu in MgCuP to adopt a tetrahedral coordination environment, which has poorer overlap with the P 3p orbitals resulting in a less disperse VBM, compared to the trigonal planar arrangement in the other three systems due to their larger group II cations. Conductivity of pressed powder samples of CaCuP indicate intrinsic p-type conductivity, and optical measurements indicate a strong absorption onset at around 2.8 eV, in good

agreement with theory^{–93} thin films or single crystals are likely to display higher levels of conductivity and mobility, with Cu vacancies likely responsible for the intrinsic conductivity. While not completely transparent, the ternary phosphides certainly teach us valuable design principles for engineering valence band dispersion.

Boron phosphide (BP) is the product of another high throughput screening process, where the search space was expanded from oxides to include sulfides, nitrides and phosphides. Indeed, the authors found that of the 30 000 compounds screened, phosphides had on average the lowest hole effective mass, but also the lowest band gaps.⁹⁴ However, for BP this is not an issue, as even though the indirect gap is quite small (2.26 eV with the HSE06 functional and 32% HF exchange) it possesses a much larger direct gap of around 4 eV, easily surpassing the transparency requirement. It is found that AlP also displays this behaviour, but the hole effective mass in BP is significantly lower and is the most promising candidate that emerged from this screening process. A separate study, using electron-phonon coupling calculations, predicts hole mobility as high as 900 cm² V⁻¹ s⁻¹.⁹⁵ The good overlap of B 2p and P 3p orbitals gives rise to the disperse valence bands, while the relative energies of the group III cation s states control the magnitude (and type) of the gap – in BP the s antibonding states are much higher in energy, resulting in the direct transition at Γ occurring between p bonding and antibonding states, whereas with Al the s antibonding states are slightly lower, causing a smaller direct band gap, and in Ga and In the s antibonding states are actually lower in energy than the p antibonding states, causing the fundamental gap to become direct and transparency unachievable.⁹⁴ There are no low energy “hole killer” defects in BP, and it is suggested that substitutional Be_B would be an excellent acceptor defect, with low formation energy and no self compensation. Experimental reports of BP are relatively sparse, with the best conductivity and hole concentration measured in the 1970s (2800 S cm⁻¹ and 1 × 10¹⁹ cm⁻³),⁹⁶ deposited in a hydrogen rich atmosphere and subsequently annealed. In ZnO, it is reported that sample growth in hydrogenic conditions can facilitate the formation of low energy hydrogen-defect complexes – subsequent annealing removes the hydrogen, but the low energy defects remain, giving excellent conductivity.¹¹⁷ It is possible that a similar mechanism is possible in BP, not accounted for in the defect study by Varley *et al.*⁹⁴ BP has also been extensively researched as a hardening coating for military aircraft windshields, evidence that it is sufficiently transparent and relatively straightforward to synthesise on a large scale.¹¹⁸ However, there have been few reports of high quality single crystals grown in recent years, indicating that synthesis challenges may still remain for p-type TCM applications.

Cuprous iodide was first documented as a transparent conductor over 100 years ago, when Bädeker iodised metallic copper films in iodine vapour, forming layers of transparent CuI.¹¹⁹ Today, film synthesis is still straightforward, with the direct evaporation of CuI powder onto glass substrates a popular deposition technique.¹⁰ With a direct band gap of around 3.1 eV, it is comfortably transparent, and has relatively



disperse valence bands formed of I 5p and Cu 3d states giving rise to a hole effective mass of roughly $0.30m_0$, indicating high mobility and conductivity.⁹⁷ The best mobility of thin films of CuI is $43.9 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$,¹²⁰ but growing epitaxial single crystals of CuI has been challenging despite the long history of the material. Recently, molecular beam epitaxy growth of single crystal CuI on sapphire and Si substrates has been reported, displaying average hole concentration and mobility of $5.47 \times 10^{17} \text{ cm}^{-3}$ and $45 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$.⁸⁰ Photoluminescence spectra indicate that CuI can emit light at an intensity one order of magnitude greater than GaN, finding possible use in LED applications, and temperature dependent measurements reveal what is suggested to be a Cu-vacancy related band, confirming previous DFT predictions that Cu vacancies are responsible for intrinsic conductivity.^{80,97} Generating a greater number of charge carriers is key to driving up conductivity in this material, but perhaps a more concerning issue is its poor air and moisture stability, especially at elevated temperatures, due to the ease with which Cu migration can occur.¹⁰

Non-oxide materials have been of great interest to researchers in recent years, as they lend themselves well to improving valence band dispersion. However, significant issues remain which question their suitability as the p-type TCMs of tomorrow. Air and moisture sensitivity are obvious issues that could prevent their widespread use, as well as the inclusion or release of toxic chemicals upon device breakdown (such as phosphorus), but that does not rule out non-oxides as useful materials for more niche applications where humans are unlikely to come into contact with the device. Despite the challenges that stability and toxicity may pose, great insight into future materials design has been gained from understanding the bonding in these systems.

4.6 Next generation quinary oxychalcogenides

The first generation of oxychalcogenides identified for p-type TCM applications had the general formula LnCuOCh – by switching out the binary oxide layer for a ternary oxide, an even greater variation in chemical composition is accessible (Fig. 6 shows this schematically). The first successful quinary oxychalcogenide was $(\text{Cu}_2\text{S}_2)(\text{Sr}_3\text{Sc}_2\text{O}_5)$, with a direct optical band gap of 3.1 eV, room temperature conductivity of 2.8 S cm^{-1} , and an impressive hole mobility of $150 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$,⁹⁸ competitive with electron mobility in the highest performing n-type TCOs. Similar to the quaternary oxychalcogenides, the hole conductivity pathway is spatially separated from the oxygen containing layers, so the valence band is not made of O 2p character but instead between Cu 3d and S 3p states, enabling high hole mobility.¹²¹

A recent screening study of 24 variations to the $(\text{Cu}_2\text{S}_2)(\text{A}_3\text{M}_2\text{O}_5)$ structure where A = Sr, Ca, Ba, and Mg and M = Sc, Al, Ga, In, Y and La has identified three novel compounds with dynamic stability and optical transparency. $(\text{Cu}_2\text{S}_2)(\text{Ca}_3\text{Al}_2\text{O}_5)$ and $(\text{Cu}_2\text{S}_2)(\text{Ba}_3\text{Sc}_2\text{O}_5)$ have optical gaps of 3.17 eV and 3.24 eV respectively, suggesting complete optical transparency, while $(\text{Cu}_2\text{S}_2)(\text{Sr}_3\text{Sc}_2\text{O}_5)$ has an optical gap of 3.06 eV, on the border of transparency.⁴⁰ Focussing on the CaAl and BaSc compounds, they possess light hole effective masses in the $\Gamma \rightarrow N$ direction of $0.37m_0$ and $0.43m_0$



Fig. 6 Exploring the chemical variations to the quinary oxychalcogenide structure. Reprinted with permission from Williamson *et al.*⁴⁰ Copyright 2021, Elsevier Inc.

respectively, indicative of extremely high hole mobility considering that $(\text{Cu}_2\text{S}_2)(\text{Sr}_3\text{Sc}_2\text{O}_5)$ displays mobility of $150 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ with a hole effective mass of $0.45m_0$. Predicted conductivities of the CaAl and BaSc compounds (from Boltzmann transport theory, assuming a hole concentration of 10^{21} cm^{-3}) are 1767 and 2058 S cm^{-1} , both exceeding the calculated value at this hole concentration for the parent compound $(\text{Cu}_2\text{S}_2)(\text{Sr}_3\text{Sc}_2\text{O}_5)$ of 1673 S cm^{-1} , but these calculations assume high levels of p-type dopability. At a lower hole concentration of 10^{18} cm^{-3} , the calculated conductivity for $(\text{Cu}_2\text{S}_2)(\text{Sr}_3\text{Sc}_2\text{O}_5)$ is 2.03 S cm^{-1} , in reasonably good agreement with the experimental value of 2.8 S cm^{-1} .⁹⁸ Powder samples of the BaSc compound confirm the direct absorption onset at around 3.24 eV, although there is an absorption of lower intensity at around 2.26 eV, which could be from impurities or unwanted transitions (such as from defect states or Cu–Cu d transitions).⁴⁰ Clearly these quinary oxysulfides show potential as p-type TCMs, but more extensive work is required – deposition as a thin film would allow for Hall measurements to be taken, quantifying the levels of conductivity and mobility, and a defect analysis would give an indication of the p-type dopability of these materials. However, one downside of these materials is their inherent structural complexity both for computationalists and experimentalists – the cost of performing a hybrid DFT defect study on a quinary system is extremely high, and the stability window of these systems may not be particularly large, especially when trying to grow under certain metal-poor conditions to encourage acceptor defect formation, such that phase pure films could be difficult to deposit.

5 An outlook on applications

A rich variety of inorganic materials have been presented in this review, covering many of the areas at the forefront of current research. This section will briefly discuss some of the current applications relying upon p-type TCMs, how these can be improved, and some of the challenges surrounding future



device integration. For thorough and dedicated reviews on applications and devices the reader is directed to the works of Shi *et al.* and Wang *et al.*^{5,11}

For industrial applications such as laptop, phone and television displays, n-type transparent conductor LCD technology continues to dominate.¹¹ With users desiring ever-improving resolution, electronic properties such as mobility and on-off ratio are absolutely vital, and p-type technology is several orders of magnitude behind systems such as amorphous indium-gallium-zinc-oxide (IGZO) thin-film transistors (TFTs).¹²² However, OLED displays are becoming increasingly popular, due to their improved contrast, lower power consumption, thinner display and wider viewing angle, and for these p-type TFTs are the preferential technology as they can simplify device architecture and improve efficiency.^{11,123}

Of the materials discussed in this review, the binary oxides Cu₂O, SnO and NiO have been most widely studied for p-type TFT applications. Despite being “the first” p-type TCO, CuAlO₂ was only first deposited as a TFT in 2012, 15 years after its conception, with device mobility < 1 cm² V⁻¹ s⁻¹ and an on-off switching ratio of around 800 (several orders of magnitude behind IGZO).^{5,124} Cu₂O can achieve the highest hole mobility, up to 256 cm² V⁻¹ s⁻¹,⁶⁵ but when implemented as a TFT the mobility fails to surpass 1 cm² V⁻¹ s⁻¹ due to poor interfacing and grain boundaries.⁶⁷ SnO also suffers from deposition problems, with SnO₂ formation severely impacting the performance in TFT and complementary metal oxide semiconductor (CMOS) applications. NiO has the better chemical stability of the three, but suffers from low device mobility when implemented as a TFT, likely due to the localised holes generated by acceptor defects.^{11,78} NiO has however been met with success when integrated into perovskite solar cells as an interfacial anode layer, improving power conversion efficiency by acting as a hole-transport layer.⁷³ Establishing improved deposition processes to nullify or contain the effects of grain boundary scattering and chemical instability is necessary to drive these oxides forward.

Implementation of materials with greater valence band dispersion is the next logical step in p-type TFT performance testing (for example CuI)¹²⁵ in order to improve device mobility, but the greatest challenges are successful deposition and chemical stability. The quasi-two-dimensional layered materials discussed in this review, such as β-TeO₂ and the oxychalcogenides for example, are perhaps less attractive than the three-dimensional or cubic crystal structures belonging to materials such as BP and NiO, due to charge transport being restricted to only two directions – this will create extra challenges for device integration. In the meantime, continued use of oxide p-type TCMs as hole transport (and electron blocking) layers to improve power conversion efficiency and overall performance is their most reliable application, while a ground-breaking, n-type matching material is awaiting discovery.

6 Conclusions

In this review, the fundamental design principles for p-type transparent conductors were outlined, the chief stratum of

which is to delocalise electron density away from single elemental species and encourage orbital mixing to form a disperse valence band. This is in line with the notion of “chemical modulation of the valence band” first suggested for p-type *oxides* by Kawazoe *et al.*,⁴¹ but in recent years has been extended to include anions such as phosphides and halides.^{93,120} The discovery of new materials with potential p-type TCM properties has been turbo-charged with the advent of large scale database screening, an approach that is likely to spearhead efforts in discovering p-type TCMs in the years to come. Indeed, an increasing number of studies concerning the discovery of new battery and solar cell materials are making use of wide-scale database screening.^{126–128} Searching beyond known compounds in databases, noted by Walsh and Park,¹²⁹ and into uncharted territory may reveal new directions for p-type TCM research – success has already been seen in the photovoltaics field, where machine-learning models have been able to predict chemical compositions that will produce band gaps suitable for solar energy applications using only earth abundant materials.¹³⁰ However, the results from such large-scale screening studies will only ever be as reliable as the databases upon which they are built, and also often return complicated quaternary and quinary crystal structures, which pose significant challenges for theorists and experimentalists alike.

Of the most interest in recent years are the significant developments made in the non-oxide field, with excellent valence band dispersion achievable in phosphides, selenides, tellurides and halides.^{63,80,92,94,116,120} Many of these materials have 3-dimensional crystal structures, in the sense that their conduction pathways exist in all three cartesian directions, such as BP due to its zinc blende structure, which is a significant advantage over the quasi-2-dimensional structures of delafossites and oxychalcogenides when implementing them into a heterojunction device. Overcoming stability, deposition and safety concerns in these simple non-oxide systems is a major goal in the coming years, as the allure of straightforward interfacing and potentially cheap synthesis of these basic crystal structures is attractive to industry – indeed, CuI has already generated interest as a p-type TFT.¹²⁵ For applications where cost is irrelevant, more complex or non-earth abundant materials are likely to dominate. Significant challenges remain for most of the systems discussed in Section 4, with the majority only in their infancy, but these emerging materials mean the future looks bright for p-type TCM research.

Author contributions

J. W. conceptualised, wrote and prepared the manuscript with guidance from DOS.

Conflicts of interest

There are no conflicts to declare.



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References

- C. G. Granqvist and A. Hultåker, *Thin Solid Films*, 2002, **411**, 1–5.
- N. Noor and I. P. Parkin, *J. Mater. Chem. C*, 2013, **1**, 984–996.
- Transparent Electronics Market – Growth, Trends and Forecast (2020–2025), 2020, <https://www.mordorintelligence.com/industry-reports/transparent-electronics-market>.
- J.-Y. Kwon, D.-J. Lee and K.-B. Kim, *Electron. Mater. Lett.*, 2011, **7**, 1–11.
- Z. Wang, P. K. Nayak, J. A. Caraveo-Frescas and H. N. Alshareef, *Adv. Mater.*, 2016, **28**, 3831–3892.
- K. H. L. Zhang, K. Xi, M. G. Blamire and R. G. Egdell, *J. Phys.: Condens. Matter*, 2016, **28**, 383002.
- N. Zhang, J. Sun and H. Gong, *Coatings*, 2019, **9**, 137.
- R. Cao, H.-X. Deng and J.-W. Luo, *ACS Appl. Mater. Interfaces*, 2019, **11**, 24837–24849.
- L. Hu, R. H. Wei, X. W. Tang, W. J. Lu, X. B. Zhu and Y. P. Sun, *J. Appl. Phys.*, 2020, **128**, 140902.
- A. N. Fioretti and M. Morales-Masis, *J. Photonics Energy*, 2020, **10**, 1.
- J. Shi, J. Zhang, L. Yang, M. Qu, D.-C. Qi and K. H. L. Zhang, *Adv. Mater.*, 2021, 2006230.
- O. Ostroverkhova, *Chem. Rev.*, 2016, **116**, 13279–13412.
- H. B. Lee, W.-Y. Jin, M. M. Ovhal, N. Kumar and J.-W. Kang, *J. Mater. Chem. C*, 2019, **7**, 1087–1110.
- J. Tauc, *Mater. Res. Bull.*, 1968, **3**, 37–46.
- A. Dolgonos, T. O. Mason and K. R. Poepplmeier, *J. Solid State Chem.*, 2016, **240**, 43–48.
- A. Walsh, J. L. F. Da Silva, S.-H. Wei, C. Körber, A. Klein, L. F. J. Piper, A. DeMasi, K. E. Smith, G. Panaccione, P. Torelli, D. J. Payne, A. Bourlange and R. G. Egdell, *Phys. Rev. Lett.*, 2008, **100**, 167402.
- M. Burbano, D. O. Scanlon and G. W. Watson, *J. Am. Chem. Soc.*, 2011, **133**, 15065–15072.
- N. F. Quackenbush, J. P. Allen, D. O. Scanlon, S. Sallis, J. A. Hewlett, A. S. Nandur, B. Chen, K. E. Smith, C. Weiland, D. A. Fischer, J. C. Woicik, B. E. White, G. W. Watson and L. F. J. Piper, *Chem. Mater.*, 2013, **25**, 3114–3123.
- Ç. Kılıç and A. Zunger, *Phys. Rev. Lett.*, 2002, **88**, 095501.
- N. Preissler, O. Bierwagen, A. T. Ramu and J. S. Speck, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **88**, 085303.
- A. M. Ganose, J. Park, A. Faghaninia, R. Woods-Robinson, K. A. Persson and A. Jain, *Nat. Commun.*, 2021, **12**, 2222.
- S. Majdi, V. Djurberg, N. Suntornwipat, M. Gabrysch and J. Isberg, *Adv. Theory Simul.*, 2020, **4**, 2000103.
- G. S. Nolas, J. Sharp and H. J. Goldsmid, *Thermoelectrics*, Springer Berlin Heidelberg, 2001, pp. 177–207.
- W. Mönch, *Appl. Surf. Sci.*, 1997, **117–118**, 380–387.
- A. Schleife, F. Fuchs, C. Rödl, J. Furthmüller and F. Bechstedt, *Appl. Phys. Lett.*, 2009, **94**, 012104.
- G. Brunin, F. Ricci, V.-A. Ha, G.-M. Rignanese and G. Hautier, *npj Comput. Mater.*, 2019, **5**, 63.
- A. Walsh, *npj Comput. Mater.*, 2021, **7**, 72.
- J. E. N. Swallow, B. A. D. Williamson, S. Sathasivam, M. Birkett, T. J. Featherstone, P. A. E. Murgatroyd, H. J. Edwards, Z. W. Lebens-Higgins, D. A. Duncan, M. Farnworth, P. Warren, N. Peng, T.-L. Lee, L. F. J. Piper, A. Regoutz, C. J. Carmalt, I. P. Parkin, V. R. Dhanak, D. O. Scanlon and T. D. Veal, *Mater. Horiz.*, 2020, **7**, 236–243.
- B. A. D. Williamson, T. J. Featherstone, S. S. Sathasivam, J. E. N. Swallow, H. Shiel, L. A. H. Jones, M. J. Smiles, A. Regoutz, T.-L. Lee, X. Xia, C. Blackman, P. K. Thakur, C. J. Carmalt, I. P. Parkin, T. D. Veal and D. O. Scanlon, *Chem. Mater.*, 2020, **32**, 1964–1973.
- D. O. Scanlon and G. W. Watson, *J. Mater. Chem.*, 2012, **22**, 25236.
- D. O. Scanlon and G. W. Watson, *J. Phys. Chem. Lett.*, 2010, **1**, 3195–3199.
- S. Lany and A. Zunger, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, **78**, 235104.
- S. Lany and A. Zunger, *Modell. Simul. Mater. Sci. Eng.*, 2009, **17**, 084002.
- C. Freysoldt, J. Neugebauer and C. G. V. de Walle, *Phys. Rev. Lett.*, 2009, **102**, 016402.
- Y. Kumagai and F. Oba, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2014, **89**, 195205.
- T. Gake, Y. Kumagai, C. Freysoldt and F. Oba, *Phys. Rev. B*, 2020, **101**, 020102(R).
- U.S. Department of the Interior, U.S. Geological Survey, Mineral Commodity Summaries 2019, 2019.
- L. Wang, D. W. Matson, E. Polikarpov, J. S. Swensen, C. C. Bonham, L. Cosimbescu, J. J. Berry, D. S. Ginley, D. J. Gaspar and A. B. Padmaperuma, *J. Appl. Phys.*, 2010, **107**, 043103.
- S. C. Dixon, S. Sathasivam, B. A. D. Williamson, D. O. Scanlon, C. J. Carmalt and I. P. Parkin, *J. Mater. Chem. C*, 2017, **5**, 7585–7597.
- B. A. D. Williamson, G. J. Limburn, G. W. Watson, G. Hyett and D. O. Scanlon, *Mater.*, 2020, **3**, 759–781.
- H. Kawazoe, M. Yasukawa, H. Hyodo, M. Kurita, H. Yanagi and H. Hosono, *Nature*, 1997, **389**, 939–942.
- T. Arnold, D. J. Payne, A. Bourlange, J. P. Hu, R. G. Egdell, L. F. J. Piper, L. Colakerol, A. De Masi, P.-A. Glans, T. Learmonth, K. E. Smith, J. Guo, D. O. Scanlon,



- A. Walsh, B. J. Morgan and G. W. Watson, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2009, **79**, 075102.
- 43 D. O. Scanlon and G. W. Watson, *J. Mater. Chem.*, 2011, **21**, 3655–3663.
- 44 Y. Xu, G.-Z. Nie, D. Zou, J.-W. Tang and Z. Ao, *Phys. Lett. A*, 2016, **380**, 3861–3865.
- 45 M. Ahmadi, M. Asemi and M. Ghanaatshoar, *Appl. Phys. Lett.*, 2018, **113**, 242101.
- 46 M. Ahmadi, M. Asemi and M. Ghanaatshoar, *Appl. Phys. A*, 2018, **124**, 529.
- 47 R. Nagarajan, N. Duan, M. Jayaraj, J. Li, K. Vanaja, A. Yokochi, A. Draeseke, J. Tate and A. Sleight, *Int. J. Inorg. Mater.*, 2001, **3**, 265–270.
- 48 K. Ueda, T. Hase, H. Yanagi, H. Kawazoe, H. Hosono, H. Ohta, M. Orita and M. Hirano, *J. Appl. Phys.*, 2001, **89**, 1790–1793.
- 49 M. N. Huda, Y. Yan, A. Walsh, S.-H. Wei and M. M. Al-Jassim, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2009, **80**, 035205.
- 50 H. Yanagi, T. Hase, S. Ibuki, K. Ueda and H. Hosono, *Appl. Phys. Lett.*, 2001, **78**, 1583–1585.
- 51 D. O. Scanlon, K. G. Godinho, B. J. Morgan and G. W. Watson, *J. Chem. Phys.*, 2010, **132**, 024707.
- 52 D. O. Scanlon, A. Walsh, B. J. Morgan, G. W. Watson, D. J. Payne and R. G. Egdell, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2009, **79**, 035101.
- 53 K. G. Godinho, J. J. Carey, B. J. Morgan, D. O. Scanlon and G. W. Watson, *J. Mater. Chem.*, 2010, **20**, 1086–1096.
- 54 C. W. Teplin, T. Kaydanova, D. L. Young, J. D. Perkins, D. S. Ginley, A. Ode and D. W. Readey, *Appl. Phys. Lett.*, 2004, **85**, 3789–3791.
- 55 S. Sheng, G. Fang, C. Li, Z. Chen, S. Ma, L. Fang and X. Zhao, *Semicond. Sci. Technol.*, 2006, **21**, 586–590.
- 56 K. Momma and F. Izumi, *J. Appl. Crystallogr.*, 2008, **41**, 653–658.
- 57 K. Ueda, S. Inoue, S. Hirose, H. Kawazoe and H. Hosono, *Appl. Phys. Lett.*, 2000, **77**, 2701–2703.
- 58 H. Hiramatsu, K. Ueda, H. Ohta, M. Orita, M. Hirano and H. Hosono, *Thin Solid Films*, 2002, **411**, 125–128.
- 59 H. Hiramatsu, M. Orita, M. Hirano, K. Ueda and H. Hosono, *J. Appl. Phys.*, 2002, **91**, 9177–9181.
- 60 H. Hiramatsu, K. Ueda, K. Takafuji, H. Ohta, M. Hirano, T. Kamiya and H. Hosono, *J. Appl. Phys.*, 2003, **94**, 5805–5808.
- 61 K. Ueda, K. Takafuji, H. Yanagi, T. Kamiya, H. Hosono, H. Hiramatsu, M. Hirano and N. Hamada, *J. Appl. Phys.*, 2007, **102**, 113714.
- 62 H. Hiramatsu, K. Ueda, H. Ohta, M. Hirano, M. Kikuchi, H. Yanagi, T. Kamiya and H. Hosono, *Appl. Phys. Lett.*, 2007, **91**, 012104.
- 63 D. O. Scanlon, J. Buckeridge, C. R. A. Catlow and G. W. Watson, *J. Mater. Chem. C*, 2014, **2**, 3429–3438.
- 64 N. Zhang, X. Liu, D. B. K. Lim and H. Gong, *ACS Appl. Mater. Interfaces*, 2020, **12**, 6090–6096.
- 65 B. Li, K. Akimoto and A. Shen, *J. Cryst. Growth*, 2009, **311**, 1102–1105.
- 66 D. O. Scanlon, B. J. Morgan, G. W. Watson and A. Walsh, *Phys. Rev. Lett.*, 2009, **103**, 096405.
- 67 W. K. Min, S. P. Park, H. J. Kim, J. H. Lee, K. Park, D. Kim, K. W. Kim and H. J. Kim, *ACS Appl. Mater. Interfaces*, 2020, **12**, 24929–24939.
- 68 J. A. Crawford and R. W. Vest, *J. Appl. Phys.*, 1964, **35**, 2413–2418.
- 69 E. Arca, A. B. Kehoe, T. D. Veal, A. Shmeliov, D. O. Scanlon, C. Downing, D. Daly, D. Mullarkey, I. V. Shvets, V. Nicolosi and G. W. Watson, *J. Mater. Chem. C*, 2017, **5**, 12610–12618.
- 70 A. B. Kehoe, E. Arca, D. O. Scanlon, I. V. Shvets and G. W. Watson, *J. Phys.: Condens. Matter*, 2016, **28**, 125501.
- 71 J. A. Caraveo-Frescas, P. K. Nayak, H. A. Al-Jawhari, D. B. Granato, U. Schwingenschlögl and H. N. Alshareef, *ACS Nano*, 2013, **7**, 5160–5167.
- 72 H. Hosono, Y. Ogo, H. Yanagi and T. Kamiya, *Electrochem. Solid-State Lett.*, 2011, **14**, H13.
- 73 M. D. Irwin, D. B. Buchholz, A. W. Hains, R. P. H. Chang and T. J. Marks, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 2783–2787.
- 74 P. Zhai, Q. Yi, J. Jian, H. Wang, P. Song, C. Dong, X. Lu, Y. Sun, J. Zhao, X. Dai, Y. Lou, H. Yang and G. Zou, *Chem. Commun.*, 2014, **50**, 1854.
- 75 S. Chen, C. Wen, T. Kuo, W. Peng and H. Lin, *Thin Solid Films*, 2014, **572**, 51–55.
- 76 P. Puspharajah, S. Radhakrishna and A. K. Arof, *J. Mater. Sci.*, 1997, **32**, 3001–3006.
- 77 J. Arunodaya and T. Sahoo, *Materials Research Express*, 2019, **7**, 016405.
- 78 H. Chen and J. H. Harding, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2012, **85**, 115127.
- 79 S. Sasaki, K. Fujino and Y. Takeuchi, *Proc. Jpn. Acad., Ser. B*, 1979, **55**, 43–48.
- 80 D. Ahn, J. D. Song, S. S. Kang, J. Y. Lim, S. H. Yang, S. Ko, S. H. Park, S. J. Park, D. S. Kim, H. J. Chang and J. Chang, *Sci. Rep.*, 2020, **10**, 3993.
- 81 A. Kudo, H. Yanagi, H. Hosono and H. Kawazoe, *Appl. Phys. Lett.*, 1998, **73**, 220–222.
- 82 E. Papadopoulou, Z. Viskadourakis, A. Pennos, G. Huyberegts and E. Aperathitis, *Thin Solid Films*, 2008, **516**, 1449–1452.
- 83 M. Dekkers, G. Rijnders and D. H. A. Blank, *Appl. Phys. Lett.*, 2007, **90**, 021903.
- 84 M. N. Amini, H. Dixit, R. Saniz, D. Lamoen and B. Partoens, *Phys. Chem. Chem. Phys.*, 2014, **16**, 2588.
- 85 K. H. L. Zhang, Y. Du, A. Papadogianni, O. Bierwagen, S. Sallis, L. F. J. Piper, M. E. Bowden, V. Shutthanandan, P. V. Sushko and S. A. Chambers, *Adv. Mater.*, 2015, **27**, 5191–5195.
- 86 A. Bhatia, G. Hautier, T. Nilgianskul, A. Miglio, J. Sun, H. J. Kim, K. H. Kim, S. Chen, G.-M. Rignanese, X. Gonze and J. Suntivich, *Chem. Mater.*, 2015, **28**, 30–34.
- 87 D. Dahliah, G.-M. Rignanese and G. Hautier, *J. Mater. Chem. C*, 2020, **8**, 9352–9357.
- 88 H. M. Kim, U. Kim, C. Park, H. Kwon and K. Char, *APL Mater.*, 2016, **4**, 056105.



- 89 D. O. Scanlon, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **87**, 161201(R).
- 90 J. Xu, J.-B. Liu, J. Wang, B.-X. Liu and B. Huang, *Adv. Funct. Mater.*, 2018, **28**, 1800332.
- 91 S. Hu, B. Xia, Y.-P. Lin, T. Katase, J. Fujioka, T. Kamiya, H. Hosono, K.-Z. Du and Z. Xiao, *Adv. Funct. Mater.*, 2020, **30**, 1909906.
- 92 A. Zavabeti, P. Aukarasereenont, H. Tuohey, N. Syed, A. Jannat, A. Elbourne, K. A. Messalea, B. Y. Zhang, B. J. Murdoch, J. G. Partridge, M. Wurdack, D. L. Creedon, J. van Embden, K. Kalantar-Zadeh, S. P. Russo, C. F. McConville and T. Daeneke, *Nat. Electron.*, 2021, **4**, 277–283.
- 93 B. A. D. Williamson, J. Buckeridge, J. Brown, S. Ansbros, R. G. Palgrave and D. O. Scanlon, *Chem. Mater.*, 2017, **29**, 2402–2413.
- 94 J. B. Varley, A. Miglio, V.-A. Ha, M. J. van Setten, G.-M. Rignanese and G. Hautier, *Chem. Mater.*, 2017, **29**, 2568–2573.
- 95 V.-A. Ha, B. Karasulu, R. Maezono, G. Brunin, J. B. Varley, G.-M. Rignanese, B. Monserrat and G. Hautier, *Phys. Rev. Mater.*, 2020, **4**, 065401.
- 96 K. Shohno, M. Takigawa and T. Nakada, *J. Cryst. Growth*, 1974, **24–25**, 193–196.
- 97 J. Wang, J. Li and S.-S. Li, *J. Appl. Phys.*, 2011, **110**, 054907.
- 98 M.-L. Liu, L.-B. Wu, F.-Q. Huang, L.-D. Chen and I.-W. Chen, *J. Appl. Phys.*, 2007, **102**, 116108.
- 99 A. Jain, S. P. Ong, G. Hautier, W. Chen, W. D. Richards, S. Dacek, S. Cholia, D. Gunter, D. Skinner, G. Ceder and K. A. Persson, *APL Mater.*, 2013, **1**, 011002.
- 100 R. Woods-Robinson, D. Broberg, A. Faghaninia, A. Jain, S. S. Dwaraknath and K. A. Persson, *Chem. Mater.*, 2018, **30**, 8375–8389.
- 101 S. Kim, J. A. Cianfrone, P. Sadik, K.-W. Kim, M. Ivill and D. P. Norton, *J. Appl. Phys.*, 2010, **107**, 103538.
- 102 H.-C. Wang, P. Pistor, M. A. L. Marques and S. Botti, *J. Mater. Chem. A*, 2019, **7**, 14705–14711.
- 103 P. Qin, G. Fang, N. Sun, X. Fan, Q. Zheng, F. Cheng, J. Wan and X. Zhao, *Thin Solid Films*, 2012, **520**, 3118–3124.
- 104 Y. S. Liu, C. I. Hsieh, Y. J. Wu, Y. S. Wei, P. M. Lee and C. Y. Liu, *Appl. Phys. Lett.*, 2012, **101**, 122107.
- 105 J. G. Lu, Z. Z. Ye, F. Zhuge, Y. J. Zeng, B. H. Zhao and L. P. Zhu, *Appl. Phys. Lett.*, 2004, **85**, 3134–3135.
- 106 C. X. Shan, J. S. Liu, Y. J. Lu, B. H. Li, F. C. C. Ling and D. Z. Shen, *Opt. Lett.*, 2015, **40**, 3041.
- 107 F. Sun, C. X. Shan, B. H. Li, Z. Z. Zhang, D. Z. Shen, Z. Y. Zhang and D. Fan, *Opt. Lett.*, 2011, **36**, 499.
- 108 I. Chatratin, F. P. Sabino, P. Reunchan, S. Limpijumnong, J. B. Varley, C. G. V. de Walle and A. Janotti, *Phys. Rev. Mater.*, 2019, **3**, 074604.
- 109 J. E. N. Swallow, B. A. D. Williamson, T. J. Whittles, M. Birkett, T. J. Featherstone, N. Peng, A. Abbott, M. Farnworth, K. J. Cheetham, P. Warren, D. O. Scanlon, V. R. Dhanak and T. D. Veal, *Adv. Funct. Mater.*, 2017, **28**, 1701900.
- 110 C. R. A. Catlow, A. A. Sokol and A. Walsh, *Chem. Commun.*, 2011, **47**, 3386.
- 111 J. L. Lyons, A. Janotti and C. G. V. de Walle, *Appl. Phys. Lett.*, 2009, **95**, 252105.
- 112 J. B. Varley, J. R. Weber, A. Janotti and C. G. V. de Walle, *Appl. Phys. Lett.*, 2010, **97**, 142106.
- 113 H. Peelaers and C. G. V. de Walle, *Phys. Status Solidi B*, 2015, **252**, 828–832.
- 114 J. E. N. Swallow, C. Vorwerk, P. Mazzolini, P. Vogt, O. Bierwagen, A. Karg, M. Eickhoff, J. Schörmann, M. R. Wagner, J. W. Roberts, P. R. Chalker, M. J. Smiles, P. Murgatroyd, S. A. Razek, Z. W. Lebens-Higgins, L. F. J. Piper, L. A. H. Jones, P. K. Thakur, T.-L. Lee, J. B. Varley, J. Furthmüller, C. Draxl, T. D. Veal and A. Regoutz, *Chem. Mater.*, 2020, **32**, 8460–8470.
- 115 X. Cai, F. P. Sabino, A. Janotti and S.-H. Wei, *Phys. Rev. B*, 2021, **103**, 115205.
- 116 S. Guo, Z. Zhu, X. Hu, W. Zhou, X. Song, S. Zhang, K. Zhang and H. Zeng, *Nanoscale*, 2018, **10**, 8397–8403.
- 117 K. Senthilkumar, M. Tokunaga, H. Okamoto, O. Senthilkumar and Y. Fujita, *Appl. Phys. Lett.*, 2010, **97**, 091907.
- 118 B. C. Monaghan, A. D. Morrison, E. M. Waddell, D. R. Gibson, S. A. D. Wilson and K. L. Lewis, *Infrared Thin Films: A Critical Review*, 1992, pp. 110–142.
- 119 K. Bädeker, *Ann. Phys.*, 1907, **327**, 749–766.
- 120 M. Grundmann, F.-L. Schein, M. Lorenz, T. Böntgen, J. Lenzner and H. von Wenckstern, *Phys. Status Solidi A*, 2013, **210**, 1671–1703.
- 121 D. O. Scanlon and G. W. Watson, *Chem. Mater.*, 2009, **21**, 5435–5442.
- 122 Y. Hara, T. Kikuchi, H. Kitagawa, J. Morinaga, H. Ohgami, H. Imai, T. Daitoh and T. Matsuo, *J. Soc. Inf. Disp.*, 2018, **26**, 169–177.
- 123 J. F. Wager, *Information Display*, 2014, **30**, 26–29.
- 124 Z. Q. Yao, B. He, L. Zhang, C. Q. Zhuang, T. W. Ng, S. L. Liu, M. Vogel, A. Kumar, W. J. Zhang, C. S. Lee, S. T. Lee and X. Jiang, *Appl. Phys. Lett.*, 2012, **100**, 062102.
- 125 A. Liu, H. Zhu, W.-T. Park, S.-J. Kang, Y. Xu, M.-G. Kim and Y.-Y. Noh, *Adv. Mater.*, 2018, **30**, 1802379.
- 126 F. T. Bülle, A. Bhowmik, T. Vegge, J. M. G. Lastra and I. E. Castelli, *Batteries Supercaps*, 2021, **4**, 1–10.
- 127 D. W. Davies, K. T. Butler, J. M. Skelton, C. Xie, A. R. Oganov and A. Walsh, *Chem. Sci.*, 2018, **9**, 1022–1030.
- 128 D. F. Dahlliah, G. Brunin, J. George, V.-A. Ha, G.-M. Rignanese and G. Hautier, *Energy Environ. Sci.*, 2021, DOI: 10.1039/d1ee00801c.
- 129 A. Walsh and J.-S. Park, *Matter*, 2020, **3**, 604–606.
- 130 D. W. Davies, K. T. Butler and A. Walsh, *Chem. Mater.*, 2019, **31**, 7221–7230.

