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Photocatalytic, Structural and Optical Properties of Mixed Anion Solid solutions Ba₃Sc₂₋ 1

_xIn_xO₅Cu₂S₂ and Ba₃In₂O₅Cu₂S_{2-v}Se_v 2

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Gregory J. Limburn^a, Matthew J. P. Stephens^a, Benjamin A. D. Williamson^b, Antonio Iborra-Torres^a, 3

David O. Scanlon^{c,d,e}, and Geoffrey Hyett^{a*} 4

^a Department of Chemistry, University of Southampton, Southampton, SO17 1BJ, UK 6

^b Department of Materials Science and Engineering, Norwegian University of Science and Technology

(NTNU), Trondheim 7491, Norway

^c Department of Chemistry, University College London, 20 Gordon Street, London, WC1H 0AJ, UK

^d Thomas Young Centre, University College London, Gower Street, London WC1E 6BT, United Kingdom

e Diamond Light Source Ltd., Diamond House, Harwell Science and Innovation Campus, Didcot, Oxfordshire

12 OX11 0DE, UK

*Corresponding author: <u>g.hyett@soton.ac.uk</u>

Abstract

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Nine members of two contiguous solid solutions, $Ba_3Sc_{2-x}In_xO_5Cu_2S_2$ and $Ba_3In_2O_5Cu_2S_{2-y}Se_y$ (x, y = 15 0. 0.5, 1, 1.5 and 2), were synthesised at temperatures between 800 °C and 900 °C by stoichiometric 16 combination of binary precursors. Their structures were determined by Rietveld refinement of X-ray 17 powder diffraction data and found to adopt the SmNi₃Ge₃ structure with I4/mmm symmetry. 18 19 Approximate Vegard law relationships were found within each solution between the lattice 20 parameters and composition, with an observed cell volume of 466.4 Å³ for Ba₃Sc₂O₅Cu₂S₂ <mark>⁰²,</mark> 21 increasing to 481.0 Å³ for Ba₃In₂O₅Cu₂S₂ and finally to 499.0 Å³ for Ba₃In₂O₅Cu₂Se₂. In the first solid 22 solution, this volume increase is driven by the replacement of scandium by the larger indium ion, generating increased strain in the copper chalcogenide layer. In the second solution the 23 substitution into the structure of the larger selenium drives further volume expansion, while 24 relieving the strain in the copper chalcogenide layer. Band gaps were estimated from reflectance 25 26 spectroscopy and these were determined to be 3.3 eV, 1.8 eV and 1.3 eV for the three end members Ba₃Sc₂O₅Cu₂S₂, Ba₃In₂O₅Cu₂S₂ Ba₃Sc₂In₂O₅Cu₂Se₂, respectively. For the intermediate 27 compositions a linear relationship between band gap size and composition was observed, driven in 28 the first solution by the introduction of the more electronegative indium lowering the conduction 29 band minimum and in the second solution by the substitution of the electropositive selenium 30 raising the valance band maximum. Photocatalytic activity was observed in all samples under solar 31

- 32 simulated light, based on a dye degradation test, with the exception of $Ba_3In_2O_5Cu_2Se_{1.5}S_{0.5}$. The



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Introduction 34

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Layered mixed anion materials have attracted significant attention due to the identification of a 35 number of examples with functional properties such as superconductivity,¹⁻⁴ p-type conductivity,⁵⁻⁷ 36 and photocatalysis.⁸⁻¹¹ In these mixed anion materials oxygen is paired with a heavier anion such as 37 sulphur or arsenic, leading to the adoption of structures with alternating layers due to the inability 38 of the larger and more polarisable heavy anions to share crystallographic sites with the oxide 39 anions. In phases where two or more cations are present, the layering is reinforced with separation 40 of the metal ions based on hard-hard or soft-soft interactions leading to a preference to occupy 41 either the oxide layers or heavier anion layers.¹² 42

43 There has been specific interest in the use of oxysulfides, such as Sm₂Ti₂O₅S₂, as visible light 44 photocatalysts for water splitting applications.¹³ The most widely studied metal oxide photocatalysts such as titanium dioxide have wide band gaps in excess of 3 eV,¹⁴ which can 45 therefore only be activated by UV light. Interest in metal chalcogenides has been driven by their 46 47 much smaller band gaps, which would allow more efficient use of solar radiation,¹⁵ but they have often been found to be unstable due to competing photocorrosion of the sulfide or selenide ions 48 making them unsuitable for use in aqueous environments.¹⁶ The most recent work by Wang et al,⁸, 49 50 has demonstrated overall photocatalytic water splitting from Y₂Ti₂O₅S₂ with IrO₂ and Rh/Cr₂O₃ co-51 catalysts and crucially they found this material to be stable with respect to photocorrosion of the 52 sulfide ions. This work further highlights the potential of oxychalcogenide materials as visible light responsive photocatalysts, and opens up the possibility of using the structural and compositional 53 <mark>%754</mark> diversity of the guinary oxychalcogenides to apply materials design to optimise a visible light active 55 photocatalyst.

For a single junction water splitting photocatalyst the band gap is the principal materials criteria for 56 efficacy. The ideal band gap value lies in the range 1.6 eV to 2.5 eV and is a compromise between 57 the need to maximise the solar flux that can be absorbed and the need to drive the water splitting 58 reactions with sufficient over potential.¹⁷ In addition to the principle criteria of band gap, a review 59 60 by Inoue has identified, based on empirical observation, a number of additional materials criteria required to ensure sufficient kinetics for a viable photocatalyst. The two most important 61 observations are that (1) active photocatalysts typically contain metal ions with d^0 or d^{10} 62 configurations, and (2) these cations have a distorted or asymmetric co-ordination environment.¹⁸ 63 Inoue postulated that the closed shell configuration is necessary to minimise electron-hole 64 recombination, as ions with d^1 - d^9 configurations could act as charge carrier traps. Ions with a d^{10} 65

configuration were also highlighted as being advantageous as materials containing these cations 66 ew Article Online would have conduction bands consisting of highly dispersed s and p orbitals providing high high 39/DOTA06629J 67 mobility pathways for photoexcited electrons. The apparent requirement for an asymmetric co-68 ordination environment was explained as producing a localised dipole favouring electron-hole 69 70 separation.

In this paper, we report the synthesis, structures, optical properties, and photocatalytic activity of 72 two novel solid solutions series comprising nine samples: Ba₃Sc_{2-x}In_xO₅Cu₂S₂ and Ba₃In₂O₅Cu₂S_{2-y}Se_y 73 74 (x, y = 0, 0.5, 1, 1.5, 2). We have found that all these materials meet the primary and secondary 75 prerequisite criteria required for effective solar photocatalysis outlined above. To the best of our 76 knowledge all of these are novel, with the exception of the end member Ba₃Sc₂O₅Cu₂S₂, which we have previously reported.¹⁹ These adopt the $A_3B_2O_5C_2X_2$ structure, a variant of the SmNi₃Ge₃ 77 structure type,²⁰ where A, B and C are cations and X is a heavy anion. These structures can be 78 79 considered as being composed of an $[A_3B_2O_5]^{2+}$ layer adopting a fragment of the *perovskite* 80 structure, and a $[C_2X_2]^{2-}$ layer where the third cation is in tetrahedral co-ordination to the heavy anion, and can be considered as a fragment of the *anti-litharge* structure. The $A_3B_2O_5C_2X_2$ structure 81 82 has been adopted by materials containing a number of different $[C_2X_2]^{2-}$ heavy anion layers including copper sulfide,^{21, 22} silver selenide,²³ and iron arsenide compounds,^{24, 25} and which can be 83 84 considered as part of a larger family of mixed anion compounds which have previously been reviewed.¹² In these materials, the properties are often dictated by the heavy anion layer. For 85 <mark>98-№</mark> example, in iron arsenide superconductors the heavy anion layer acts as the charge carrier layer,²⁶ and similarly in p-type conducting copper sulphides the heavy anion layer is the transport layer.²⁷ 87

For the solid solutions $Ba_3Sc_{2-x}In_xO_5Cu_2S_2$ and $Ba_3In_2O_5Cu_2S_{2-y}Se_y$ we will show that the band gap 88 can be controlled as a function of composition, spanning the useful range for photocatalysis of 89 1.32 eV to 3.27 eV. They also contain the d^{10} indium (III) ion in an asymmetric co-ordination 90 environment, which can favour electron-hole separation, while the presence of copper 91 chalcogenide layers are anticipated to act as effective hole conduction pathways.²⁷ Finally, the 92 photocatalytic ability of the materials under solar simulated radiation has been directly confirmed 93 using a dye degradation test. 94

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96 Experimental Methods

Solid-state synthesis. Samples in the solid solutions Ba₃Sc_{2-x}In_xO₅Cu₂S₂ and Ba₃In₂O₅Cu₂S_{2-x}Se_x. 97 98 with x and y = 0, 0.5, 1, 1.5 and 2 were prepared on a 0.5 g scale by combining appropriate stoichiometric ratios of binary sulfide, selenide and oxide precursors in a nitrogen filled glove box. 99 ខ្លា<mark>់</mark>100 Samples were pelletised and placed in alumina crucibles, before being sealed under vacuum in 101 gg silica ampoules. The alumina cubicles prevented the samples from contacting the silica tubes with which they could react at elevated temperatures. The sealed pellets of $Ba_3Sc_xIn_{2-x}O_5Cu_2S_2$ (series 1) <u>...</u>103 were heated for 12 hours at 800 °C while samples of Ba₃In₂O₅Cu₂S_xSe_{2-x} (series 2) were heated for j 104 24 hours at 900 °C. The precursors used were Sc₂O₃, In₂O₃, Cu₂S, Cu₂O, BaS, BaO, and BaSe. Sc₂O₃ ŭ 105 (Sigma-Aldrich, 99.9%), In₂O₃ (Sigma-Aldrich, 99.99%) Cu₂S (Sigma-Aldrich, 99.99%) and Cu₂O (Alfa .⁵106 Aesar, 99.9%) were purchased and used as provided. BaS was synthesized by the reaction of BaCO₃ ¥107 (Alfa Aesar 99.99%) with CS₂ vapour (Fisher, 99.8%) in argon (BOC Pureshield), at 900 °C for 8 801^{mo} 109 109 hours.²⁸ BaO was synthesized by decomposition of BaCO₃ under vacuum at 1000 °C for 14 hours. BaSe was produced in a two-stage process. In the first step, water insoluble barium selenite was ບໍ່ັ110 prepared by precipitation upon mixing of two 200 ml aqueous solutions; one of 4.3 g of Na₂SeO₄ **111** (Sigma-Aldrich, BioXtra) and the other of 6.0 g Ba(NO₃)₂ (Sigma-Aldrich, 99%), chilled to 0 °C over §112 ice. This produced a white precipitate of BaSeO₄ that was filtered and dried in air at 70 °C. The ^{.s} 113 BaSeO₄ was then reduced under an atmosphere of 5% H₂ in N₂ (BOC) at 500 °C for 4 hours, to yield 114 This artic BaSe. The purity of all synthesized precursors was confirmed by X-ray diffraction. All precursor storage and handling was carried out in a nitrogen-filled glove box. Once prepared, samples of 115 94.16 N-Ag 17 Ba₃Sc_{2-x}In_xO₅Cu₂S₂ and Ba₃In₂O₅Cu₂S_{2-v}Se_v were found to be air stable and stored under standard atmosphere.

Materials characterisation. The layered copper chalcogenide samples were characterised using 118 powder X-ray diffraction patterns collected on a Bruker D2 diffractometer, equipped with a copper 119 K_{α} X-ray source. Patterns were collected in the range 10° < 2 θ <100° two theta, with a step size of 120 0.02° and a collection time of at least 3 hours. Patterns were analysed by the Rietveld refinement 121 method carried out using the GSAS-II software suite.²⁹ An additional diffraction pattern was 122 collected on a sample LaB₆ and this was used as the crystalline standard to provide a pseudo-voigt 123 peak profile function to account for instrumental broadening, and used in each of the subsequent 124 refinements. The compounds were also analysed using EDX spectroscopy, with data collected using 125 a JOEL JSM5910 scanning electron microscope with a 10 keV acceleration voltage, and the Oxford 126 127 INCA 300 EDS system.

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128 Diffuse reflectance spectra were collected on the powder samples using a Perkin Elmer Lambda

129 750S spectrometer, equipped with an integrating sphere. Data were collected in the range of 300⁽⁰⁶⁶²⁹⁾

nm to 2500 nm, and the Kubelka-Munk relationship used to estimate sample absorption.³⁰ Band

131 gaps for the samples were determined using the Tauc method from this spectroscopic data.³¹

Photocatalysis tests. An assessment was made of the photocatalytic potential of the materials. For this, the samples were decorated with a cobalt oxide co-catalyst, using a procedure modified from the literature.³² 25 mg of each chalcogenide powder sample was impregnated with 2 ml of a 8.59 × 10^{-4} mol dm⁻³ solution of Co(NO₃)₂ \square 6H₂O (Sigma Aldrich, 98+%) in acetone, and then left exposed to air to allow the solvent to evaporate overnight. The resulting powder samples were heated to 700 °C under nitrogen (flow rate: 250 ml min⁻¹) for 60 min, decomposing the cobalt nitrate to presumed metallic cobalt nanoparticles. The active CoO_x co-catalyst was obtained on the powder surface by mild oxidation of the cobalt loaded powders in air at 200 °C for 1h.

The photocatalytic test used was the degradation of a blue dye, dichloroindophenol (DCIP, Acros 98+%), in the presence of a glycerol as a sacrificial oxidant (Fisher, reagent grade).³³ The test solution was made up in dichloromethane (DCM, Fisher, reagent grade) with a concentration of DCIP of 5.5×10^{-5} mol dm⁻³ and a concentration of glycerol of 5.5×10^{-3} mol dm⁻³. In each experiment, a 5 mg sample of the cobalt loaded oxychalcogenide powder was placed in the test solution and agitated by a magnetic stirrer. The slurry was exposed to a 5-sun solar simulator (LS0104 150 W Xenon lamp) with a measured intensity of 509 mW cm⁻² for 3 hours, and then a centrifuge was used to separate the powder from the dye solution. The solution was then transferred to a cuvette and a visible absorption spectrum recorded in the range 400 nm to 800 nm, using a PE750S spectrometer. Changes in dye concentration caused by photocatalytic degradation were determined assuming a Beer-Lambert law relationship between absorption and concentration.

Computational methodology. Computational calculations were carried out using density 152 functional theory (DFT) within the Vienna ab-initio Simulation Package (VASP).³⁴⁻³⁷ The hybrid 153 functional HSE06³⁸ (α = 25%) was used to accurately describe the nature of the electronic structure 154 of each end member compound relative to experiment and has been shown to accurately describe 155 Sr₃Sc₂O₅Cu₂S₂ and Ba3Sc2O₅Cu₂S₂ in previous work.^{19, 27}To describe the interaction between the 156 core and valence electrons, the projector augmented wave (PAW) method was employed.³⁹ The 157 computational workflow involved a geometric optimisation of each compound, Ba₃Sc₂O₅Cu₂S₂, 158 Ba₃In₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂Se₂ using a Γ -centred 6×6×2 k-point grid and a plane-wave energy 159 6

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cut-off of 500 eV where convergence was deemed complete when the maximum force on any 160

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162 Density of states (DOS), band structure and optical absorption calculations were carried using the relaxed cells and plotted using the open source python package: sumo.⁴⁰ The optical absorption 163 spectra were calculated using the real and imaginary parts of the dielectric constant calculated ยู่164 165 gg through a Kramers-Kronig transformation and a summation over the unoccupied bands respectively using a method by Furthmüller and co-workers.⁴¹ Within this formalism however, intracial 3.0 band and indirect absorptions are ignored and only the direct valence to conduction band transitions are considered.41,42

The effective masses of charge carriers (m^*_{ij}) can be calculated using the relationship:

$$\frac{1}{m_{ii}^*} = \frac{\delta^2 E(k) \, 1}{\delta k_i \delta k_j \, \hbar^2}$$

Where E(k) is the eigenvalue of the band at a specific k.

atom was less than 0.01 eV Å⁻¹.

Calculation of the phonon spectra was performed using the finite-displacement method within the Phonopy⁴³ code using the PBEsol functional^{44, 45} (with a +U value of 5.17 eV applied to the copper atoms) known for accurately describing the structural parameters and bond lengths necessary for accurate phonon calculations.²⁷ Initially each compound was relaxed to a tight electronic convergence criterion of 1×10⁻⁸ eV Å⁻¹ and force convergence of 1×10⁻³ eV Å⁻¹ such that the residual forces on all the atoms was minimal.

879 **Results and discussion**

All targeted members of the two solid solutions, Ba₃Sc_{2-x}In_xO₅Cu₂S₂ (series 1) and Ba₃In₂O₅Cu₂S_{2-y}Se_y 180 (series 2), were successfully made with a minimum 95% purity, and an average 97.7% purity as 181 determined by XRD and supported by EDX measurements. The powder X-ray diffraction patterns of 182 the target compounds were refined from a starting model based on the previously reported 183 Sr₃Sc₂O₅Cu₂S₂ structure,⁶ but with appropriate ion substitutions to match the assumed composition. 184 In each refinement the background, lattice parameters and atomic positions were refined. Uniaxial 185 size and strain parameters were refined to account for peak broadening beyond the instrumental 186 broadening determined from modelling data collected on a LaB_6 high crystallinity sample. 187 188 Constraints were applied to make the thermal ellipsoids of identical atoms equal. The positions of

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the Sc/In mixed cation sites in series 1 and the S/Se mixed anion sites in series 2 were also 189 /iew Article Online constrained to be identical. Initially the isotropic thermal ellipsoid parameters of the atoms were 190 191 refined, but these were found to be too strongly correlated to the atomic site positions, preventing both from being be refined simultaneously while still producing meaningful results. To overcome 192 this, fixed values for the thermal displacement for each atom type were used (for example 0.02 Å² for all barium atoms, full list in ESI Tables S1 and S2) allowing the atomic site positions to be refined so that meaningful comparisons of bond distances and angles could be made between samples. The values for these fixed thermal displacements were derived by taking the average values from the nine initial refinements. Additional peaks were also observed in each diffraction pattern, which could not be assigned to the main phase. These were due to impurity phases, variously BaCO₃, BaCu₂S₂ and Sc₂O₃. These were modelled using the standard structures from the ICSD and literature,⁴⁶⁻⁴⁸ allowing for the phase fractions to be refined. Lattice parameters and particle size broadening parameters were also refined for these impurity phases where sufficient peaks could be observed.

Using this methodology, good fits to the data were found for all models based on the expected compositions, with wR_p values ranging from 3.16% to 4.55%, confirming the successful formation of the target phases. The diffraction patterns and their model Rietveld fits can be found in the ESI figures S1 to S3, with an example given in figure 1 of the refined fit to the diffraction data collected on Ba₃In₂O₅Cu₂S₂. The purity of the samples was found to be high, with the principal phase comprising between 95.6 wt% to 98.7 wt% in each case. A summary of the refinement results for the series 1 samples can be found in table 1, and the series 2 samples in table 2. Full details of the structural refinements can be found in the ESI in table S1 and S2, including the refined atomic site positions, thermal ellipsoid values used, and impurity concentrations.

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	Ba ₃ Sc ₂ O ₅ Cu ₂ S	$Ba_3Sc_{1.5}In_{0.5}O_5Cu_2$	Ba ₃ ScInO ₅ Cu ₂ S	$Ba_3Sc_{0.5}In_{1.5}O_5Cu_2S$	Ba ₃ In ₂ O ₅ Cu ₂ S ₂
	2	S	2		DOI: 10.1039/D0TA06629J
Lattice	4.1458(2)	4.1577(2)	4.1680(3)	4.1772(2)	4.1862(1)
parameter a / Å	4.15				4.19
Lattice	27.136(2)	27.218(2)	27.317(4)	27.401(2)	27.444(1)
parameter c / Å	27.38				27.73
Volume / Å ³	466.42(6)	470.50(5)	474.55(8)	478.12(6)	480.95(2)
	471.48				487.91
Data Points	4444	4444	4444	4444	4345
Reflections	100	101	101	102	98
(main phase)					
Parameters	34	29	24	30	23
Purity	95.6%	96.1%	98.7%	97.9%	97.5%
wRp	3.72%	3.43%	3.99%	3.16%	4.46%
RF ²	3.32%	2.18%	3.48%	2.28%	3.12%
Chi2	2.04	1.98	2.38	1.96	1.93

Table 1. Summary Rietveld refinement for series 1 compounds $Ba_3Sc_{2-x}In_xO_5Cu_2S_2$. All structures were refined in *I4/mmm*, and errors are two standard deviations. Full site refinement results and CIF files are available in the ESI. Values of *a*, *c* and volume given in italics are the result of the computational modelling based on HSE06 functional.



Figure 1. Rietveld refinement of 14/mmm structure of $Ba_3In_2O_5Cu_2S_2$ to diffraction data, with lattice parameters a = 4.1862 Å and c = 27.444 Å. Upper section: diffraction data is in blue, model is in green and background in red. Inset shows the high angle region from 60° to 100° 2 θ . Lower section shows the difference curve (cyan) and tick marks indicating the positions of the peaks of, in descending order: $Ba_3In_2O_5Cu_2S_2$ (green), $BaCu_2S_2$ (red) and $BaCO_3$ (cyan).

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	$Ba_3In_2O_5Cu_2S_2^*$	$Ba_3In_2O_5Cu_2S_{1.5}Se_{0.}$	Ba ₃ In ₂ O ₅ Cu ₂ SSe	$Ba_3In_2O_5CuS_{0.5}Se_{1.5}$	$Ba_3In_2O_5Cu_2Se$
		5			2
Lattice	4.1862(1)	4.1958(2)	4.2060(2)	4.2138(2)	4.2225(1)
parameter a /	4.19				4.22
Å					
Lattice	27.444(1)	27.576(2)	27.716(2)	27.841(2)	27.985(1)
parameter c /	27.73				28.03
Å					
Volume / Å ³	480.95(2)	485.47(5)	490.31(5)	494.35(6)	498.95(4)
	487.91				499.67
Data Points	4345	4444	4444	4148	4444
Reflections	98	106	108	98	111
(main phase)					
Parameters	23	30	30	30	32
Purity	97.5%	98.7%	98.6%	98.5%	97.8%

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wRp	4.46%	3.87%	4.11%	4.55%	4.28%	
RF ²	3.12%	3.55%	3.85%	4.74%	3.30% View #	rticle Online
Chi2	1.93	1.72	1.93	3.50	2.19	JTA06629J

Table 2. Rietveld refinement results for series 2 compounds, $Ba_3In_2O_5Cu_2S_{2-y}Se_y$. All structures were refined in *I4/mmm*, and errors are two standard deviations. Full site refinement and CIF files are available in the ESI. Values of *a*, *c* and volume given in italics are the result of the computational modelling based on HSE06 functional. *Also the end member of Series 1 data repeated her for convenience.

The results of the analysis of the diffraction data were supported by energy dispersive X-ray spectroscopy (EDX). For each sample, EDX data were collected on 5 different particles chosen at random, and the elemental composition quantified based on the expected heavy elements, with oxygen excluded from the analysis due to presence of environmental contaminants containing oxygen preventing reliable estimation of the amount of this element in the sample. The average elemental composition for each sample resulting from this analysis is shown graphically in figure 2, alongside the values expected from the targeted stoichiometry as a guide. These show a good match between the experimental and expected values, so that we can conclude that both the XRD and EDX data confirm that the target phases have been made as intended. The end member Ba₃Sc₂O₅Cu₂S₂ has been previously reported,¹⁹ however to the best of our knowledge the solid solution end members Ba₃In₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂Se₂, alongside all the intermediate phases, are reported in this work for the first time.



Figure 2. Plot of elemental composition as determined by EDX excluding oxygen, for the two solid solutions: $Ba_3Sc_{2-x}In_xO_5Cu_2S_2$ and B $a_3In_2O_5Cu_2S_{2-y}Se_y$. Thin solid lines represent the percentage of each element expected based on the target composition.

Lattice parameters for the layered oxychalcogenides derived from the Rietveld refined structural
models, as a function of composition, are displayed in figure 3. These show a good approximation
to Vegard's law for both solid solutions. In series 1 the replacement of the smaller scandium ion
(radius of 0.75 Å) with the larger indium ion (0.80 Å) leads to a steady increase in both the *a* and *c*lattice parameters, with an increase of approximately 1% in both *a* and *c* once full replacement by

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 $Ba_3In_2O_5Cu_2S_2$, we observe a similar increase in both a and c lattice parameters. Coincidently, the 240

241 increase in a in series 2 occurs at a similar rate as a function of the fraction of substitution as in

series 1, although that of the c parameter is greater. Overall in series 2 there is an increase of 0.9% 242 in a in series 2 for the full substitution of selenium, but an expansion of approximately 2% in the c .<mark>243</mark> a Creative Commons Attribution-NonCommercial 3.0 Unported Licer 5 9 5 9 parameter.



Figure 3. Plot of lattice parameters for the $Ba_3Sc_{2-x}In_xO_5Cu_2S_2$ and $Ba_3In_2O_5Cu_2S_{2-y}Se_y$ solid solutions (x,y = 0, 0.5, 1, 1.5 2) as a function of composition. The a lattice parameter scale is to the left and values are shown as black squares. The c lattice parameter scale is to the right, and values are shown with hollow triangles. Error bars are smaller than the symbols used, and the dashed lines are guides for the eye, and indicative of the Vegard law relationship between lattice dimensions and composition.

^{.s}250 The results of the refinement of the atomic site positions within the unit cell models provides bond ^{oj}251 lengths and angles, and therefore insight into the changes in the structures across the solid solutions that drive the overall expansion of the unit cells. The limitations of laboratory X-ray 252 53 diffraction equipment mean that the position of the light oxygen atoms can only be ascertained 54 with limited confidence. However, it is still possible to draw conclusions regarding the general trends, and with greater confidence for the heavier atoms. Figure 4 shows variations in key bond 255 256 lengths and angles across the structures as a function of composition. These values can be found tabulated in the ESI in table S3. 257

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Figure 4. Left: Schematic diagram of the unit cell of the $Ba_3MO_5Cu_2Ch_2$ structures, where M = Sc, In and Ch = S, Se. Centre and Right: Plots of the key angles (triangles) and bond lengths (squares) of the oxide and chalcogenide blocks respectively as a function of composition, including the variation in overall block height for each (circles). Where they are not shown, error bars are smaller than the symbols used.

The general unit cell diagram for these samples is shown in figure 4, and highlights that the structure can be considered as layered blocks of $[Ba_3M_2O_5]^{2+}$ and $[Cu_2Ch_2]^{2-}$, where M = Sc or In and Ch = S or Se. The nature of this layering means that any change in one of the blocks that expands it parallel to the [100] direction must be matched by an expansion in that direction in the other block in order to maintain a commensurate *a* lattice parameter. For example, an expansion of the oxide block in the [100] direction due to a substitution would inevitably force a change in the bonding or geometry to the chalcogenide layer even if it remains compositionally unchanged. In contrast changes to the dimensions of a block parallel to the [001] direction can occur independently, without necessarily affecting the structure of the other layer. Thus changes to the structure that lead to changes in the *a* lattice parameter are dependent upon both oxide and chalcogenide blocks, whereas changes in the *c* direction can occur due to either variations in the height of the oxide block, the chalcogenide layer, or the interlayer spacing, and these can be independent of each other.

In series 1 there is steady overall increase in the value of both the *a* and *c* lattice parameters of approximately 1% from $Ba_3Sc_2O_5Cu_2S_2$ to $Ba_3In_2O_5Cu_2S_2$. The key structural change is the substitution of the larger indium into the oxide block which drives the 1% expansion of the cell in the *a* direction. The geometry of the Sc/In site is pyramidal, and the equatorial metal oxygen bond increases in length by ~0.7% across the series, accounting for the majority of the expansion in the *a* lattice parameter, the rest of the expansion coming from a small change in the geometry with a decrease in the O_{ax} -*M*- O_{eq} (*M* = Sc/In) bond angle from 97.8(5)° to 96.7(4)° as the average metal site 12 Page 13 of 23

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position moves towards the sulfur. The most significant bond length change is the increase in the 283 /iew Article Online axial M-O bond length from 1.98(1) Å to 2.062(3) Å, an increase of ~4%. It is this increase which the addition of the second sec 284 drives the overall expansion of the cell in the *c* direction. The response of the compositionally 285 constant copper sulphide layer to the expansion in the *a* lattice parameter is a reduction in the 286 block height parallel to the [001] direction, counter to the overall expansion of the c lattice <mark>.</mark>:287 .288 parameter. As might be expected the Cu-S bond length remains approximately constant, with the 289 change in the height of the copper containing layer instead being driven by a distortion in the <u>ୂ</u>290 copper geometry, observed by the increase in the S-Cu-S angle from 113.7(5)° to ~115.8(4)°. 165^{rel} Therefore, the decrease in the sulfide block height with expansion in the [001] direction is driven by ^b292 changes in the oxide block, but the overall size of the unit cell increases in the *c* direction because ^N-293 of the larger increase in the height of the $[Ba_3M_2O_5]^{2+}$ layer.

nquuy 294 This accordion-like scissoring of the copper sulfide block with an inverse relationship between the suo 295 block height and the a lattice parameter due to distortions in the copper sulfur bond angles follows the trend observed in the oxysulfides, and accounts for the compatibly of the copper chalcogenide §296 ບັ້297 layers with a diversity of metal oxide layers of differing dimensions. Prior works on related materials ⁹298 have found examples of copper sulfide layers that can accommodate values of a lattice parameter §299 as low as 3.83 Å in Ca₄Fe₂O₆Cu₂S₂ and as high as 4.08 Å in Sr₃Sc₂O₅Cu₂S₂,^{22, 49} or up to 4.12 Å if the fluoride layered BaFCuS is considered.⁵⁰ The values of *a* lattice parameter we have identified for the 108^{arti} Ba₃Sc_{2-x}In_xO₅Cu₂S_{2 solid} solution of 4.146 Å to 4.186 Å significantly extends the range of feasible dimensions for layered copper sulfides, and to our knowledge are the largest values yet reported. 102 원 03 The end member Ba₃In₂O₅Cu₂S₂ may well be reaching the limit of what is possible for a stable [Cu₂S₂]²⁻ layer, with a S-Cu-S angle of approximately 116° showing significantly higher distortion 04 **1** 305 away from tetrahedral geometry than any prior example.

In series 2, Ba₃In₂O₅Cu₂S_{2-v}Se_v, the expansion of the lattice parameters is driven by changes in the 306 307 chalcogenide layer, in contrast to series 1 where it occurs in the oxide layer. With the substitution of the larger selenium for sulphur we see the expected increase in the copper chalcogenide bond 308 length, of 2.7% across the series. Simultaneously we observe a relaxation of the copper 309 environment back towards a more ideal tetrahedral geometry, with the Ch-Cu-Ch bond angle 310 decreasing to 112.5(1)° indicating that the larger copper selenide layer is a better, less strained 311 match for the [Ba₃In₂O₅]²⁺ layer than the copper sulfide layer. The combined effect of these changes 312 is an expansion of the *a* lattice parameter of ~0.9%, and also a very significant increase in the 313 314 height of the chalcogenide block in the [001] direction across the series of 7%. The response of the

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now compositionally unchanging [Ba₃In₂O₅]²⁺ layer to the increase in the *a* lattice parameter is a compensatory decrease in height parallel to the [001] direction, although the effect is Smaller than^{6629J} the similar compensation effect observed in the chalcogenide layer in series 1. The equatorial In-O bond length increases slightly, by 0.5% across the series, while the apical In-O length remains approximately constant. The key effect of the changing dimensions of the oxide block is a change in the geometry of the indium ion with the O-In-O bond angle decreasing in order to accommodate the increase in the [100] direction, and this is the cause of the overall decrease in the height of the oxide block.

The results of the computational modelling of the structures of the end member compositions using the HSE06 functional and PBEsol functional are provided the ESI in tables S4 and S5, including the lattice parameters, bond lengths and bond angles. Each of the end member compounds displayed the same I4/mmm symmetry after geometric relaxation to within 0.01 Å⁻¹. Each of the calculated lattice parameters are within 2% of the experimental values owing to the efficacy of both HSE06 and PBEsol in describing the structural properties of this family of layered oxychalcogenide..^{19, 27}As finite temperatures are typically neglected from DFT, these results indicate that there is minimal thermal expansion and change to the structural properties between 0K and RTP and thus it is unlikely that the calculated electronic properties will deviate from those observed through experiment As expected, the substitution of S for Se increases the a and c parameters and increases all of the bond lengths respectively whilst the Cu-Ch-Cu (Ch = S, Se) bond angle decreases. This results in a contraction of the $[Cu_2Ch_2]^{2-}$ layers shown by the Ba-Ba interlayer distances, or 'block height' also **1**34 seen in the experimental values. In terms of bond lengths, the layers typically remain independent, 36 i.e. between Ba₃Sc₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂S₂ the Cu-Ch and Cu-Cu distances show minimal changes, yet within the $[Ba_3M_2O_5]^{2+}$ layer, the Ba-O and M-O distances undergo a larger change than 338 between Ba₃In₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂Se₂. As mentioned in previous work, the influence of structure as well as composition plays a distinct role in the tunability of these layered 339 oxychalcogenide semiconductors.¹⁹ 340

In order to estimate the band gaps for the materials, diffuse reflection spectra were recorded on each of the powder samples and the reflectance data were modelled using the Kubelka-Munk function, f(R). Tauc plots of $[f(R).hv]^2$ against photon energy (hv) were used to estimate the direct band gap by extrapolation of a tangent from the point of inflection to the abscissa, and can be found in ESI figure S4. This analysis gave band gap values of 3.27(5) eV for Ba₃Sc₂O₅Cu₂S₂, 1.77(5) eV for Ba₃In₃O₅Cu₂S₂, and 1.32(5) eV for Ba₃In₃O₅Cu₂Se₂, the end members of the solid solutions.

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- For the intermediate compositions, there is an approximately linear relationship between the 347
- composition and the experimentally determined band gap, as shown in figure 5(a), although With a shown in fi 348
- 349 slight deviation for the series 1 solution, $Ba_3Sc_{2-x}In_xO_5Cu_2S_2$.



Figure 5 (a): Experimentally determined band gap values as a function of composition for the solid solutions $Ba_3Sc_{2x}In_xO_5Cu_2S_2$ and Ba₃In₂O₅Cu₂S_{2-y}Se_y. Dotted lines represent linear relationship between end members, and are shown as a guide for the eye. (b-d): Band structures for $Ba_3Sc_2O_5Cu_2S_2$, and $Ba_3Sc_2O_5Cu_2S_2$, and $Ba_3Sc_2O_5Cu_2S_2$ respectively. In each panel the valence and conduction bands are coloured blue and orange respectively and the valance band maximum is set to 0 eV. (e-q): Calculated density of states diagrams for the three end member compounds. In each panel the valence band maximum (VBM) is set to 0 eV. For the two Incontaining compounds, (f) and (g), a scaled view of the CBM is inset.

⁻ E 357 The band structures of the three end members were computationally modelled, to help understand 58 the changes in band gap, and can be seen in figures 5(b)-(d), with the density of states diagrams in 59 figure 5(e)-(g). For each compound the valence band maxima (VBM) is composed approximately of B 60 50:50 Cu 3d:Ch p states meaning that the [Cu₂Ch₂]²⁻ layer dominates the p-type conductivity in these systems and has been noted previously.^{19, 27} The full break down of states at the VBM and 361 362 conduction band minimum (CBM) are given in the ESI in table S6. The majority of the O 2p states lie around -6 eV to -2 eV for each compound, owing to the electronegativity of oxygen and the 363 reason why typically in metal oxide photocatalysts such as TiO₂ the VBM lies deep compared to the 364 redox potentials of water.⁵¹ It is expected that a better bulk band alignment will exist for each of 365 these compounds due to the VBM moving towards the vacuum due to the overlap between Cu 3d 366 and Ch p. This has the added effect of increasing dispersion at the VBM giving rise to lower 367 effective masses. For Ba₃Sc₂O₅Cu₂S₂, Ba₃In₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂Se₂, the direct fundamental 368 band gaps are modelled as 3.04 eV, 1.37 eV and 1.09 eV respectively and are direct (at Γ) with the 369 370 exception of $Ba_3Sc_2O_5Cu_2S_2$ where the VBM occurs at Γ and the CBM at the X point. The energy 15

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difference between the indirect and direct band gap (at Γ) is < 2 meV, however it is well known that 371 this type of behaviour gives rise to enhanced exciton lifetimes such as seen in anatase^CTiO₂¹⁰ BIOX^{629J} 372 (X = CI, Br, I) photocatalysts.^{52, 53} In all three end member compounds there exists two degenerate 373 bands at the VBM resulting in heavy and light hole effective masses. The full results of the effective 374 mass modelling are given in the ESI in Table S7, however herein the light effective masses will be _.375 documented, as the path of least resistance will likely be taken. The hole effective masses in the $[Cu_2Ch_2]^{2-}$ plane can be found at $\Gamma - N$ and $\Gamma - Z$ whilst the direction corresponding to the interplanar direction is given by $\Gamma - X$. The $\Gamma - X$ direction results in highly flat bands as expected with (light) hole effective masses of ~42 m_{e_1} ~84 m_e and ~41 m_e respectively. For Ba₃Sc₂O₅Cu₂S₂, Ba₃In₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂Se₂ the light effective masses in the $\Gamma - N$ and $\Gamma - Z$ direction are 0.44 m_e and 0.65 m_e, 0.38 m_e and 0.51 m_e, and 0.26 m_e and 0.36 m_e respectively. It is clear then, that both the increase in a lattice parameter as well as the exchange of sulfur for selenium results in a lowering of the valence band effective mass.

The composition of the CBM is largely determined by the $[Ba_3M_2O_5]^{2+}$ layer. In the case of Ba₃In₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂S₂, the CBM is dominated by In 5s states mixed with O 2s states with negligible contributions from Cu 4s, Ch ns states and Ba 6s states. This is in contrast with Ba₃Sc₂O₅Cu₂S₂ where scandium states do not play a role and instead the CBM is made up of Ba 5d states mixed with Cu 3s+d and negligible O 2s+p states. The reasoning for this is due to the lower lying In s states compared to the Ba d states which results in lower band gaps despite the larger a lattice parameter from Ba₃Sc₂O₅Cu₂S₂ to Ba₃In₂O₅Cu₂S₂. The CBM dispersion difference between 190 Ba₃Sc₂O₅Cu₂S₂ and the In-containing compounds is distinctly noticeable in the band structures in Figure 5, whereby the In 5s states form highly dispersive bands with effective masses of 0.95 m_e, 92 393 0.21 m_e and 0.19 m_e respectively. Parallels can be made with the n-type conducting flagship transparent conducting oxide: In_2O_3 which also show the same low CBM effective masses of ~0.22 394 395 me.⁵⁴ In practice this means that Ba₃In₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂Se₂ will possess high electron mobility which will aid transport of photoexcited electrons to the surface of the material to perform 396 catalysis.⁵⁵ The In-based systems may also be bipolar semiconductors, a property which has been 397 shown to exist in delafossite CuInO2.56,57 398

The modelled band structures provide insight into the observed experimental trends in the band gap. In Series 1 the change in the composition of the CBM moves it lower in energy towards the valance band with increasing indium content, while the VBM itself remains undisturbed. The other key effect of the exchange of indium for scandium in the first series, as observed in analysis of the

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X-ray diffraction data, is in the expansion of the lattice parameters and an increase in the Cu-Cu 403 distance. This has been previously shown to lead to a small increase in band gap due to decreased 6629J 404 405 dispersion of the valence and conduction bands under the influence of the Cu-Cu interactions.¹² The effect is expected to be approximately only +0.04 eV for the increase in Cu-Cu distance in the 406 .407 Ba₃Sc_{2-x}In_xO₅Cu₂S₂ solution, however it may account for the small deviation from trend observed for .⁵⁰/2408 the latter members of the solution, as shown in the first half of figure 5(a). In the second solid pated 409 solution the exchange of sulphur in $Ba_3In_2O_5Cu_2S_2$ leads to a shift in the valance band maximum õe 410 towards the vacuum and giving an overall decrease in the band gap. The effect across the whole of nercial 11 the solution is a decrease of the band gap by 0.45 eV. This is comparable to similar observed ^b2412 differences in band gap energy for pairs of layered oxychalcogenides where both the oxysulfide and oxyselenide are known, for example, $Sr_3Sc_2O_5Cu_2Ch_2$ with band gaps of 3.10 eV for Ch = S and ^{nq}1414 2.90 eV when $Ch = Se_{1,58}^{6,58}$ and LaOCuCh with band gaps 3.14 eV (Ch = S) and 2.82 eV ($Ch = Se_{1,59}^{59}$

The optical absorption coefficients were also simulated and can be found in the ESI in figure S5, as a function of photon energy. The optical band gaps for Ba₃Sc₂O₅Cu₂S₂, Ba₃In₂O₅Cu₂S₂ and ບັ້417 Ba₃In₂O₅Cu₂Se₂, are calculated to be 3.24 eV, 1.84 eV and 1.48 eV respectively. This corresponds well **e**418 with those gained in experiment (3.27 eV, 1.77 eV and 1.32 eV respectively). A strong (steep) ₅419 absorption corresponds to an increased likelihood of photoexcited charge carriers and therefore a ^{...} <u>9</u>420 greater potential for photocatalytic activity. Each end member compound has a relatively steep 125 This artic absorption, particularly Ba₃Sc₂O₅Cu₂S₂ however for the series 2 compounds the onset is more gradual. For Ba₃In₂O₅Cu₂S₂ and Ba₃In₂O₅Cu₂Se₂ the difference in optical band gaps is not as large as 122 ¥.23 compared to Ba₃In₂O₅Cu₂S₂ and Ba₃Sc₂O₅Cu₂S₂. Thus solid solutions of Ba₃In₂O₅Cu₂S_{2-v}Se_v will not largely affect the optical properties but will affect the electronic properties by raising the VBM and 24 425 increasing VBM curvature for more mobile holes evident from the band structures and effective 426 masses.

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	Ba ₃ Sc _{2-x} In _x O ₅ Cu ₂ S ₂				Ba ₃ In ₂ O ₅ Cu ₂ S _{2-y} Se _y					
x/y	0	0.5	1	1.5	2	0	0.5	1	1.5	2
Band Gap / eV	3.27(5	2.80(5	2.41(5	2.09(5)	1.77(5	1.77(5	1.65(5)	1.55(5	1.44(5	1.32(5)
)))))))	

DCIP Degradation 27.0% 22.9% 14.8% 15.4% 14.5% 14.5% 5.9% 3.8% 0.3% 5.5% Ba₃In₂O₅Cu₂S_{2-y}Se_y. Dye degradation is the percentage of the dye degraded after 3 hours exposure to a solar simulator lamp. Error on DCIP measurements +2/5%/2simulator lamp. Error on DCIP measurements ±2/5%/

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Photocatalytic testing was carried using degradation of the blue dye DCIP in the presence of 432 8 433 glycerol as a sacrificial oxidant, using an unfiltered solar simulator as the light source. 5 mg samples <u>5</u>434 of oxychalcogenide powders were loaded with cobalt oxide oxidation co-catalyst, at approximately [™] 1435 0.4 wt% in order to enhance the observable rates.^{60, 61} The powder samples were continuously 0. 1930 1930 stirred to ensure homogeneous dispersion in the test solution across the 3 hour test time, after 437 which the solution was separated from the powder using a centrifuge and the change in dye 2438 concentration determined. A control experiment where the dye solution was exposed to the solar ب 10 439 simulator for 3 hours without any powder sample present found that no degradation occurs within ¹¹/₂440 the error of the measurement. The majority of the co-catalyst loaded oxychalcogenide samples <u></u>441 were found to be active photocatalysts, except for Ba₃In₂O₅Cu₂S_{0.5}Se_{1.5}. This data can be seen in Table 3. The most active samples were the two with the largest band gaps, Ba₃Sc₂O₅Cu₂S₂ and చ్ ¤443 Ba₃Sc_{1.5}In_{0.5}Cu₂S₂ where 27% and 23% of the dye was found to have degraded, respectively. The under 1900 percentages of the dye degraded for all the samples are shown in figure 6, where the remaining <u></u> <u>5</u>445 active samples have degradation fractions over 3 hours of between 5% and 15%. Across both solid .s 9446 solutions there is an inverse correlation between the extent of photocatalytic dye degradation and .8447 the band gap size. DCIP dye degradation is an indicative test, and the greater rates observed for the materials with the larger band gaps is likely to be due to the increased over potential for the 48 49 photocatalytic reaction. However, the results of dye degradation tests need to be considered with -50 caution, as although they can indicate photocatalytic activity, they cannot be used as a guarantee that the catalyst will be active for the more challenging water splitting reaction. Dye degradation 451 can also depend on the surface area, the spectral absorption overlap between the dye, and the 452 photocatalyst surface charge.⁶² For our solid solutions the samples are sufficiently similar that these 453 factors should not prevent a relative comparison of photocatalytic activity, but further research will 454 be necessary to confirm if they are capable of total organic carbon removal, or water splitting. 455

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DCIP dye degradation in 3 hours /



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Ba3In2O5Cu2S28

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Conclusions

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We have demonstrated the synthesis of two new solid solutions, Ba₃Sc_{2-x}In_xO₅Cu₂S₂ and Ba₃In₂O₅-Cu₂S_{2-v}Se_v, and shown that almost all of these display photocatalytic activity. The shared end member $Ba_{1}In_{2}O_{5}Cu_{2}S_{2}$ has a basal lattice parameter of 4.1861 Å, which is the largest yet reported for a layered oxysulfide containing anti-litharge structured copper sulphide layer. This therefore expands the upper range for the use of the [Cu₂S₂]²⁻ structural unit when considering a building block approach to materials design.

Spectroscopic analysis has shown that the with the two solid solutions it is possible to select, 67 BY-NC 68 through appropriate choice of composition, a photocatalytic material with any band gap ranging ¥69 from 3.30 eV to 1.32 eV. This spans the visible spectrum from the near UV to the near IR, although the driving force behind the change in band gap differs for the two solid solutions. From 470 Ba₃Sc₂O₅Cu₂S₂ to Ba₃In₂O₅Cu₂S₂ the conduction band minimum decreases, while from 471 Ba₃In₂O₅Cu₂S₂ to Ba₃In₂O₅Cu₂S₂ the valence band maximum increases. The ability to tune the band 472 473 gap and band edge positions through composition in the two solid solutions is a key feature that could be used in the future in the optimisation of the absorber in a water splitting photocatalyst 474 475 system.

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481 Supporting Information

Electronic Supplementary Information (ESI) available. Figures S1, S2 and S3: Rietveld refinements of X-ray diffraction patterns. Figure S4: Tauc plots derived from experimental spectra, Figure S5: calculated optical absorption data and Figure S6: calculated phonon dispersion data. The ESI also includes tables S1 and S2: details of the refined structural model and table S3: the refined bond lengths and angles. Tables S4 and S5: structural details of the computational derived models, Table S6: data on the calculated VBM and CBM composition, and table S7: the calculated effective electron and hole masses. Figure S7 Model and recorded IR spectra, S8 Position of Principle IR peaks. Table S8: tabulated IR data. See DOI: 10.1039/x0xx00000x

All data supporting this study are openly available from the University of Southampton repository at https://doi.org/10.5258/SOTON/D1457

References

- 1. D. Johrendt and R. Pottgen, *Angewandte Chemie-International Edition*, 2008, **47**, 4782-4784.
- 2. X. Zhu, F. Han, G. Mu, P. Cheng, B. Shen, B. Zeng and H.-H. Wen, *Physical Review B*, 2009, **79**, 220512.
- H. Ogino, Y. Matsumura, Y. Katsura, K. Ushiyama, S. Horii, K. Kishio and J.-i. Shimoyama, *Supercond. Sci. Technol.*, 2009, **22**, 075008.
- 4. X. Zhu, F. Han, G. Mu, B. Zeng, P. Cheng, B. Shen and H.-H. Wen, *Physical Review B*, 2009, **79**, 024516.
- B. H. Hiramatsu, K. Ueda, H. Ohta, M. Hirano, M. Kikuchi, H. Yanagi, T. Kamiya and H. Hosono, Appl.
 Phys. Lett., 2007, 91, 012104.
- 02 6. M.-L. Liu, L.-B. Wu, F.-Q. Huang, L.-D. Chen and I.-W. Chen, J. Appl. Phys., 2007, **102**, 116108.
 - S. Jin, X. Chen, J. Guo, M. Lei, J. Lin, J. Xi, W. Wang and W. Wang, *Inorg. Chem.*, 2012, **51**, 10185-10192.
- 5058.Q. Wang, M. Nakabayashi, T. Hisatomi, S. Sun, S. Akiyama, Z. Wang, Z. Pan, X. Xiao, T. Watanabe, T.506Yamada, N. Shibata, T. Takata and K. Domen, Nat. Mater., 2019, **18**, 827-832.
- 5079.M. Yashima, K. Ogisu and K. Domen, Acta Crystallographica Section B-Structural Science, 2008, 64,508291-298.
- A. Ishikawa, T. Takata, T. Matsumura, J. N. Kondo, M. Hara, H. Kobayashi and K. Domen, *J. Phys. Chem. B*, 2004, **108**, 2637-2642.
- 511 11. A. Ishikawa, T. Takata, J. N. Kondo, M. Hara, H. Kobayashi and K. Domen, *J. Am. Chem. Soc.*, 2002,
 512 124, 13547-13553.
- S. J. Clarke, P. Adamson, S. J. C. Herkelrath, O. J. Rutt, D. R. Parker, M. J. Pitcher and C. F. Smura,
 Inorg. Chem., 2008, **47**, 8473-8486.
- 51513.A. Ishikawa, Y. Yamada, T. Takata, J. N. Kondo, M. Hara, H. Kobayashi and K. Domen, Chem. Mater.,5162003, 15, 4442-4446.
- 517 14. F. E. Osterloh, *Chem. Mater.*, 2008, **20**, 35-54.
- 518 15. K. Maeda and K. Domen, *J. Phys. Chem. C.*, 2007, **111**, 7851-7861.
- 519 16. A. B. Ellis, S. W. Kaiser, J. M. Bolts and M. S. Wrighton, *J. Am. Chem. Soc.*, 1977, **99**, 2839-2848.

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520	17.	J. R. Bolton, S. J. Strickler and J. S. Connolly, <i>Nature</i> , 1985, 316 , 495-500.

- 521 18. Y. Inoue, Energy & Env. Sci., 2009, 2, 364-386.
- 10.1039/D0TA06629J DOI B. A. D. Williamson, G. J. Limburn, G. W. Watson, G. Hyett and D. O. Scanlon, Matter, 2020, DOI: 522 19. 523 https://doi.org/10.1016/j.matt.2020.05.020.
- 524 20. O. Y. Mruz, V. K. Pecharsky, A. N. Sobolev and O. I. Bodak, Kristallografiya, 1990, 35, 202-204.
- 525 21. W. J. Zhu and P. H. Hor, J. Solid State Chem., 1997, 134, 128-131.
- 526 22. K. Otzschi, H. Ogino, J. Shimoyama and K. Kishio, J. Low Temp. Phys., 1999, 117, 729-733.
- g527 23. H. Ogino, Y. Katagi, J.-i. Shimoyama, K. Yamanoi, M. Tsuboi, T. Shimizu, N. Sarukura and K. Kishio, ⁵528 Optical Materials, 2014, 36, 1978-1981.
- ^{__} 529 F. Hummel, M. Tegel, B. Gerke, R. Pöttgen and D. Johrendt, Zeitschrift für Naturforschung B, 2015, 24. ž530 70, 671-676.
- ວັ້531 G. F. Chen, T. L. Xia, H. X. Yang, J. Q. Li, P. Zheng, J. L. Luo and N. L. Wang, Supercond. Sci. Technol., 25. 2531 532 532 533 533 533 534 2009, **22**, 072001.
- 26. D. Johrendt, H. Hosono, R. D. Hoffmann and R. Pottgen, Z. Krist.-Cryst. Mater., 2011, 226, 435-446.
- D. O. Scanlon and G. W. Watson, Chem. Mater., 2009, 21, 5435-5442. 27.
- ల్డ్535 28. G. Hyett, Z. A. Gal, C. F. Smura and S. J. Clarke, Chem. Mater., 2008, 20, 559-566.
- **2536** 29. B. H. Toby and R. B. Von Dreele, J. Appl. Crystallogr., 2013, 46, 544-549.
- .5<u>5</u>37 30. P. Kubelka, J. Opt. Soc. Am., 1948, 38, 448-457.
- [.] [[]=538 J. Tauc, Mater. Res. Bull., 1968, 3, 37-46. 31.
- Ž539 32. A. E. Maegli, S. Pokrant, T. Hisatomi, M. Trottmann, K. Domen and A. Weidenkaff, The Journal of 2539 2540 *Physical Chemistry C*, 2014, **118**, 16344-16351.
- <u></u>541 33. A. Mills and M. McGrady, J. Photochem. Photobiol., A, 2008, 193, 228-236.
- §542 34. G. Kresse and J. Furthmuller, Computational Materials Science, 1996, 6, 15-50.
- 35. G. Kresse and J. Furthmuller, *Physical Review B*, 1996, **54**, 11169-11186.
- ⊸544 G. Kresse and J. Hafner, *Physical Review B*, 1993, 47, 558-561. 36.
- ⁹2545 37. G. Kresse and J. Hafner, *Physical Review B*, 1994, **49**, 14251-14269.
- จู้546 A. V. Krukau, O. A. Vydrov, A. F. Izmaylov and G. E. Scuseria, J. Chem. Phys., 2006, 125. 38.
- ہ547 39. P. E. Blochl, *Physical Review B*, 1994, **50**, 17953-17979.
- <u>_</u>548 A. M. Ganose, A. J. Jackson and D. O. Scanlon, Journal of Open Source Software, 2018, 3, 717-719. 40.
- ಕ್ಷೆ549 B. Adolph, J. Furthmuller and F. Bechstedt, *Physical Review B*, 2001, 63, 125108. 41.
- ^{ta}550 42. M. Gajdos, K. Hummer, G. Kresse, J. Furthmuller and F. Bechstedt, *Physical Review B*, 2006, 73, Ë551 045112.
- 52 A. Togo and I. Tanaka, Scr. Mater., 2015, 108, 1-5. 43.
- G. I. Csonka, J. P. Perdew, A. Ruzsinszky, P. H. T. Philipsen, S. Lebegue, J. Paier, O. A. Vydrov and J. 53 44. BY-NC 54 G. Angyan, *Physical Review B*, 2009, **79**, 155107.
- 55 J. P. Perdew, A. Ruzsinszky, G. I. Csonka, O. A. Vydrov, G. E. Scuseria, L. A. Constantin, X. L. Zhou and 45. 356 K. Burke, Phys. Rev. Lett., 2008, 100, 136406.
- 557 J. P. Devilliers, Am. Mineral., 1971, 56, 758-767. 46.
- 558 47. J. E. Iglesias, K. E. Pachali and H. Steinfink, J. Solid State Chem., 1974, 9, 6-14.
- W. O. Milligan, L. W. Vernon, H. A. Levy and S. W. Peterson, The Journal of Physical Chemistry, 559 48. 560 1953, 57, 535-537.
- 561 49. D. O. Charkin, A. V. Sadakov, O. E. Omel'yanovskii and S. M. Kazakov, Mater. Res. Bull., 2010, 45, 562 2012-2016.
- 563 50. H. Yanagi, S. Park, A. D. Draeseke, D. A. Keszler and J. Tate, J. Solid State Chem., 2003, 175, 34-38.
- 564 51. B. A. D. Williamson, J. Buckeridge, N. P. Chadwick, S. Sathasivam, C. J. Carmalt, I. P. Parkin and D. O. 565 Scanlon, Chem. Mater., 2019, 31, 2577-2589.
- 566 52. A. M. Ganose, M. Cuff, K. T. Butler, A. Walsh and D. O. Scanlon, *Chem. Mater.*, 2016, **28**, 1980-1984.
- 567 A. M. Alotaibi, B. A. D. Williamson, S. Sathasivam, A. Kafizas, M. Alqahtani, C. Sotelo-Vazquez, J. 53. 568 Buckeridge, J. Wu, S. P. Nair, D. O. Scanlon and I. P. Parkin, Acs Applied Materials & Interfaces, 569 2020, 12, 15348-15361.
- 570 54. J. E. N. Swallow, B. A. D. Williamson, S. Sathasivam, M. Birkett, T. J. Featherstone, P. A. E. 571 Murgatroyd, H. J. Edwards, Z. W. Lebens-Higgins, D. A. Duncan, M. Farnworth, P. Warren, N. H.

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- 572 Peng, T. L. Lee, L. F. J. Piper, A. Regoutz, C. J. Carmalt, I. P. Parkin, V. R. Dhanak, D. O. Scanlon and T. View Article Online DOI: 10.1039/D0TA06629J 573 D. Veal, Materials Horizons, 2020, 7, 236-243.

- 574 55. X. Zhou and H. Dong, Chemcatchem, 2019, 11, 3686-3713.
- 575 56. K. G. Godinho, B. J. Morgan, J. P. Allen, D. O. Scanlon and G. W. Watson, Journal of Physics-576 Condensed Matter, 2011, 23, 334201.
- 577 57. M. Sasaki and M. Shimode, J. Phys. Chem. Solids, 2003, 64, 1675-1679.
- 578 58. Y. Iwasa, H. Ogino, D. Song, K. Yamanoi, T. Shimizu, J. Ueda, S. Tanabe and N. Sarukura, Optical g579 Materials, 2018, 84, 205-208.
- ^{ਡੋ}580 59. K. Ueda and H. Hosono, J. Appl. Phys., 2002, 91, 4768-4770.
- 581 F. Oehler and S. G. Ebbinghaus, Solid State Sci., 2016, 54, 43-48. 60.
- 582 F. Oehler, R. Naumann, R. Köferstein, D. Hesse and S. G. Ebbinghaus, Mater. Res. Bull., 2016, 73, 61. ⊃̃583 276-283.
 - 62. S. Bae, S. Kim, S. Lee and W. Choi, Catal. Today, 2014, 224, 21-28.



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