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# Deep eutectic solvent route synthesis of zinc and copper vanadate n-type semiconductors – mapping oxygen vacancies and their effect on photovoltage†

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Ternary metal oxides  $M_2V_2O_{7-\delta}$  ( $M = \text{Zn}$  and  $\text{Cu}$ ) were synthesized by dissolving binary metal oxide precursors in an environmentally benign deep eutectic solvent (DES), which is a eutectic mixture of a hydrogen bond donor and acceptor, followed by annealing in an open crucible. The unique reaction environment provided by the evolved ammonia allows for stabilization of oxygen vacancies and reduced oxidation states of metal ions within an oxide matrix without the need for any post-treatment with flammable reducing agents. According to comprehensive characterization, including X-ray photoelectron spectroscopy (XPS), neutron powder diffraction, and UV-vis spectroscopy, oxygen deficiency is accompanied by reduced oxidation states of metal centers ( $\text{Cu}^+$  or  $\text{V}^{4+}$ ), resulting in oxides with mixed-valence metal oxidation states. The amount of oxygen vacancies can be tuned by changing the annealing temperature providing control over band gaps of ternary metal oxides and mid-gap states from reduced metal centers. All synthesized vanadates are n-type semiconductors based on negative photovoltages obtained from surface photovoltage spectroscopy (SPS). A decay of the photovoltage with increasing annealing temperatures is attributed to electron trapping and electron/hole recombination at  $\text{V}^{4+}$  and  $\text{Cu}^+$  mid-gap states. This work shows for the first time the impact of oxygen vacancies on the electronic structure of DES synthesized oxides for solar energy conversion applications.

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## Introduction

Since the discovery of photoelectrochemical (PEC) water oxidation with  $\text{TiO}_2$ ,<sup>1</sup> research on semiconducting metal oxide photocatalysts has been growing. Other  $d^0$  binary metal oxides containing  $\text{Zr}^{4+}$ ,  $\text{Nb}^{5+}$ ,  $\text{Ta}^{5+}$ , and  $\text{W}^{6+}$  have been studied due to their photocatalytic properties.<sup>2–13</sup> However, their wide band-gaps limit absorption of visible-light, which is a major part of the solar spectrum at the surface of the earth. Ternary metal oxides are attractive candidates for solar PEC applications because their crystal and, consequently, electronic structure can be modified by introducing an additional metal.<sup>13</sup> For example,  $\text{SnNbO}_6$  with a band gap of 2.3 eV exhibits improved photocatalytic activity for hydrogen evolution under visible light irradiation with Pt loading.<sup>14</sup>  $\text{Fe}_2\text{V}_4\text{O}_{13}$  was found to have a narrow band gap of 1.83 eV and show high efficiency in photoreduction of  $\text{CO}_2$  to hydrocarbon fuels.<sup>15</sup>

Oxygen vacancies play a central role in optical absorption, transport properties, and photocatalytic performance of oxide materials. For example,  $\text{SrTiO}_3$  photoelectrodes must be annealed in a hydrogen atmosphere at 1050 °C in order to produce the desired semiconducting behavior for water oxidation.<sup>16</sup> Similarly, hydrogenated  $\text{TiO}_2$  can be prepared by heat treatment of white  $\text{TiO}_2$  in 20 bar  $\text{H}_2$  at 200 °C for 5 days.<sup>17</sup> This oxygen-defective  $\text{TiO}_{2-\delta}$  exhibits improved UV photocatalytic activity for hydrogen evolution from aqueous solutions of sacrificial electron donors. Hydrogen-treated  $\text{WO}_3$  and  $\text{BiVO}_4$  with oxygen vacancies exhibit enhanced catalytic activity for photoelectrochemical water oxidation.<sup>18,19</sup>

Recently, copper vanadates were reported to be a potential photoanode candidate for the visible-light driven water splitting reaction.<sup>20–24</sup> In the electronic structure of  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$  the valence band maximum (VBM) and conduction band minimum (CBM) are dominated by O 2p and Cu 3d orbitals respectively.<sup>22</sup> The partially filled d-subshell of copper improves visible light absorption and photocurrent for the oxygen evolution reaction.<sup>20–23</sup> Interestingly,  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$  and  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$  belong to the family of  $M_2V_2O_7$  ( $M = \text{Zn}$  and  $\text{Cu}$ ) with similar monoclinic structures. Both compounds also resemble each other in that they have O 2p as the main contribution to the valence band

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maximum (VBM).<sup>22,25</sup> However, the conduction band of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  predominantly consists of V 3d orbitals due to the completely filled d-subshell of zinc.<sup>25</sup> To the best of our knowledge, a systematic study on  $\text{M}_2\text{V}_2\text{O}_7$  ( $\text{M} = \text{Zn}$  and  $\text{Cu}$ ) has not been reported.

In this work,  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_{7-n}$  and  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_{7-m}$  were prepared by reaction in a deep eutectic solvent (DES), which is a eutectic mixture of a hydrogen bond donor (urea) and acceptor (choline chloride) in a 2 : 1 molar ratio.<sup>26–28</sup> This synthetic route includes dissolution of binary oxides in an environmentally friendly and benign eutectic mixture followed by heat treatment in air. High solubility of binary metal oxides in the DES and fast mixing of the metal precursors are attractive features of this synthetic method, which also provide control over the size and composition of the products and concentration of oxygen vacancies. Moreover, this route does not require flammable reducing agents and/or a complex experimental set-up for post-treatment. Thermal stability and synthesis optimization of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_{7-n}$  and  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_{7-m}$  were studied, and band gap modification by oxygen vacancies was investigated. The impact of oxygen vacancies on the generation of a surface photovoltage under visible light is discussed.

## Experimental

### Materials synthesis

Urea (Alfa, 99%, crystalline),  $\text{ZnO}$  (Alfa, 99.9%),  $\text{Cu}_2\text{O}$  (Alfa, 99.9%), and  $\text{V}_2\text{O}_5$  (Alfa, 99.6% min) were used as received without any further purification. Choline chloride (Sigma-Aldrich,  $\geq 98\%$ ) was dried at 120 °C under vacuum prior to use. A deep eutectic solvent (DES) was prepared by mixing urea and choline chloride in a 2 : 1 molar ratio (54.322 g urea and 63.750 g choline chloride for  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  and 55.200 g urea and 64.800 g choline chloride for  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ ). The mixture was heated at 70 °C in a parafilm-covered glass beaker with stirring until a homogeneous liquid was obtained. Each metal precursor (0.567 g  $\text{ZnO}$  and 0.633 g  $\text{V}_2\text{O}_5$  for  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  and 0.506 g  $\text{Cu}_2\text{O}$  and 0.644 g  $\text{V}_2\text{O}_5$  for  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ ) was dissolved in a separate beaker containing the DES under vigorous stirring at 70 °C. The concentration of  $\text{ZnO}$ ,  $\text{Cu}_2\text{O}$ , and  $\text{V}_2\text{O}_5$  was 0.06986 M, 0.03493 M, and 0.03493 M, respectively. The solutions with the dissolved precursors were further mixed ( $\text{ZnO}$  and  $\text{V}_2\text{O}_5$  for  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  and  $\text{Cu}_2\text{O}$  and  $\text{V}_2\text{O}_5$  for  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ ) and left stirring at 70 °C overnight with a parafilm cover. Afterwards, the solutions (light green for  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  and dark blue for  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ ) were transferred to porcelain crucibles for calcination (7 mL of solution in 30 mL crucibles). A heat treatment was performed in open crucibles, in air using a muffle furnace (box-type). A heating rate of 10 °C  $\text{min}^{-1}$  was employed. For  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$ , the precursor solution was heated at 230 °C for 2 hours and then calcined at 500 °C, 600 °C, or 700 °C for a span of 10.5 hours (500 °C) or 12 hours (600 °C and 700 °C). For  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ , the precursor solution was heated at 300 °C for 7 hours and then calcination was carried out at 400 °C or 500 °C for 7 hours. About 27 mg of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  (76% yield) and 32 mg of  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  (72% yield) powdered products were obtained. Different colors were observed for samples annealed

at different temperatures: pale yellow (500 °C)/grey (600 °C)/light pink (700 °C) for  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  and orange (400 °C)/dark red (500 °C) for  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ .

### Characterization

**Powder X-ray diffraction.** Samples were characterized by powder X-ray diffraction (PXRD) using a Rigaku Miniflex 600 diffractometer with Cu K $\alpha$  radiation ( $\lambda = 1.54051$  Å). Diffraction scans were collected from 5 to 90° 2 $\theta$  on a zero-background plate at room temperature in air. Phase identification was performed with the PDF-2 database using PDXL software.<sup>29</sup>

**In situ synchrotron powder X-ray diffraction.** Variable temperature synchrotron powder X-ray diffraction data were collected at the synchrotron beamline 17-BM at the Advanced Photon Source (APS), Argonne National Laboratory (ANL). Pre-calcined samples were loaded into silica capillaries (0.5 mm inner diameter and 0.7 mm outer diameter), which were placed inside a secondary shield capillary in a flow furnace. A thermocouple was set as close as possible to the sealed end of the inner silica capillary, and 20% oxygen gas in helium gas was flowed into the open end of the inner silica capillary during the measurement. The data were collected with  $\lambda = 0.24128$  Å in a temperature range from 23 °C to 800 °C with a heating rate of 20 °C  $\text{min}^{-1}$ . More details regarding the experimental setup can be found elsewhere.<sup>30</sup> Temperature calibration was applied by calibration using the melting points of elemental Sn, Sb, and Ge.

**Scanning electron microscopy and energy dispersive spectroscopy.** Scanning electron microscopy (SEM) was performed using a FEI Quanta 250 field emission SEM at 15 kV. Energy dispersive X-ray spectroscopy (EDS) was performed using an Oxford X-Max 80 detector for elemental composition analysis. Powdered samples were deposited on a SEM sample holder using carbon tape, followed by coating with 5 nm of iridium metal.

**Diffuse reflectance UV-vis spectroscopy.** Diffuse reflectance UV-vis spectra were collected using a BLACK-Comet C-SR-100 spectrometer equipped with a SL1 tungsten halogen lamp (vis-IR) and a SL3 deuterium lamp (UV). Compacted samples were prepared on glass slides by flattening the solids with a metal spatula. Indirect bandgap values for samples were estimated by extrapolating the linear slope of Tauc plots by plotting  $(\alpha \times h\nu)^{1/2}$  vs.  $h\nu$ , where  $\alpha$  is absorption and  $h\nu$  is the excitation energy in eV.

**Differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA).** DSC/TGA measurements were taken using a Netzsch STA449 F1 Jupiter coupled with a Netzsch quadrupole mass spectrometer 403 D Aeolos and a Bruker Tensor 37 FTIR spectrometer. Zinc vanadate and copper vanadate samples for DSC/TG were previously calcined at 500 °C and 400 °C, respectively, using the temperature profile described in the Materials synthesis section. Approximately 5 mg of sample was loaded into an alumina crucible with an alumina cover. The sample was heated from 40 °C to 950 °C and cooled from 950 °C to 310 °C with a ramp/cooling rate of 10 °C  $\text{min}^{-1}$ . The experiments were performed under a constant gas flow of 20%  $\text{O}_2$  and 80%  $\text{N}_2$ .

**X-ray photoelectron spectroscopy (XPS).** XPS measurements were performed using a Kratos Amicus/ESCA 3400 instrument.



The sample was irradiated with 240 W non-monochromated Mg K $\alpha$  X-rays, and photoelectrons emitted at 0° from the surface normal were energy analyzed using a DuPont type analyzer. The pass energy was set at 150 eV. CasaXPS was used to process raw data files and either a Shirley or linear baseline was applied to the spectra. The adventitious C 1s set to a binding energy of 284.8 eV was used for charge correction of XPS spectra. The % concentration of “oxygen defects”, V<sup>4+</sup> and Cu<sup>+</sup> was calculated as the ratio of the peak areas: the peak with the binding energy, corresponding to the species in question (e.g. O defects, V<sup>4+</sup>, and Cu<sup>+</sup>) over the sum of the peak areas for “oxygen defects” and lattice oxygen, or V<sup>4+</sup> and V<sup>5+</sup>, or Cu<sup>2+</sup> and Cu<sup>+</sup>.

**High-resolution synchrotron powder X-ray diffraction (HR SPD).** High-resolution synchrotron powder X-ray diffraction data were collected at 295 K using beamline 11-BM at the Advanced Photon Source (APS), Argonne National Laboratory using an average wavelength of 0.412749 Å. Discrete detectors covering an angular range from −6 to 16° 2 $\theta$  are scanned over a 34° 2 $\theta$  range, with data points collected every 0.001° 2 $\theta$  at a scan speed of 0.01° s<sup>−1</sup>. Rietveld refinement was performed using GSAS software.<sup>31</sup>

**Neutron powder diffraction (NPD).** Time-of-flight NPD was performed on a POWGEN diffractometer at the Spallation Neutron Source in Oak Ridge National Laboratory. The diffraction data were collected at 298 K using neutron beams with wavelengths of 0.8 Å and 2.665 Å for *d*-spacing ranges of 0.40–5.66 Å and 1.08–5.57 Å, respectively. 495 mg sample was loaded into a vanadium can (6 mm inner diameter). Rietveld refinement was performed using GSAS software.<sup>31</sup>

**Surface photovoltage spectroscopy (SPS) measurements.** SPS data were obtained using the vibrating Kelvin probe method. Here a gold mesh reference probe was mounted approximately 1.0 mm above the sample film and controlled with a Kelvin control 07 unit (Delta PHI Besocke, Germany). The samples were illuminated with monochromatic light from a 150 W Xe lamp filtered through an Oriel Cornerstone 130 monochromator (0.1–0.3 mW cm<sup>−2</sup>). The spectra were corrected for drift effects by subtracting a dark scan.

**Sample preparation for SPS.** Fluorine-doped tin oxide-coated glass (FTO) substrates for surface photovoltage measurements were sonicated in acetone and water for 3 hours each to remove any surface impurities. Sample powders were suspended in H<sub>2</sub>O at a concentration of 5 mg mL<sup>−1</sup> and sonicated for 15 minutes. Then 0.1 mL solution of each sample was deposited dropwise onto a 1 cm<sup>2</sup> area on a cleaned substrate and dried uncovered in air. The  $\beta$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub> films were annealed at 400 °C for 2 hours in air with a slow ramp of 2 °C min<sup>−1</sup>. To remove possible V<sub>2</sub>O<sub>5</sub> impurities after annealing, the films were dipped in 0.1 M NaOH solution at pH = 13 for 30 minutes with careful stirring, rinsed in H<sub>2</sub>O, and dried in air. SPS measurements were then performed in air.  $\alpha$ -Zn<sub>2</sub>VO<sub>2</sub>O<sub>7</sub> films were pre-annealed on a hotplate at 250 °C for 3 minutes in air and then slowly (2 °C min<sup>−1</sup>) heated up to 500 °C in air and the temperature was maintained for 2 hours. SPS measurements of these samples were performed under vacuum (2.5 × 10<sup>−4</sup> mbar).

## Results and discussion

The monoclinic structures of M<sub>2</sub>V<sub>2</sub>O<sub>7</sub> vanadates are similar for both metals, M = Cu and Zn, and are based on corner-sharing VO<sub>4</sub> tetrahedra and MO<sub>5</sub> units with two distinct coordination environments for Cu and Zn (Fig. 1a and c).<sup>32,33</sup> Each Zn ion in  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> is coordinated with five oxygen atoms to form a ZnO<sub>5</sub> distorted trigonal bipyramid, while the coordination environment of Cu ions in  $\beta$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub> is a distorted square pyramid.<sup>32,33</sup> In turn, in the high temperature polymorphs of M<sub>2</sub>V<sub>2</sub>O<sub>7</sub> (Fig. 1b and d), Zn and Cu ions have a distorted tetrahedral coordination environment.<sup>34,35</sup> Monoclinic  $\beta$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> has a V ion occupying the distorted VO<sub>5</sub> trigonal bipyramid, while VO<sub>4</sub> tetrahedra are found in orthorhombic  $\alpha$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub>.<sup>34,35</sup>

## Synthesis

A deep eutectic solvent is a eutectic mixture of a hydrogen-bond donor and a hydrogen-bond acceptor, typically quaternary ammonium salt choline chloride C<sub>5</sub>H<sub>14</sub>ClNO ((2-hydroxyethyl) trimethylammonium chloride, melting point of 302 °C).<sup>26–28</sup> Due to hydrogen bonding between choline chloride and the hydrogen-bond donor, the melting point of the mixture is lowered as compared to that of its individual components.<sup>26–28</sup> In this work, urea (mp 132–135 °C) was used as the hydrogen-bond donor, which when combined with choline chloride in a molar ratio of 2 : 1 forms a eutectic with a melting point of 30 °C, as determined by establishing a solid–liquid binary phase diagram of choline chloride–urea.<sup>36</sup> Binary transition metal oxides, such as ZnO, Cu<sub>2</sub>O, and V<sub>2</sub>O<sub>5</sub>, exhibit high solubility in the urea : choline eutectic mixture: 1.31 M for ZnO at 70 °C, 0.189 M for Cu<sub>2</sub>O at 70 °C, and 0.030 M for V<sub>2</sub>O<sub>5</sub> at 50 °C.<sup>37</sup> Notably, these solubilities are comparable to that in hydrochloric acid,<sup>37</sup> thus the deep eutectic solvent can be considered as an environmentally sustainable replacement for corrosive acids in the synthesis of materials. Cu<sub>2</sub>O was utilized for the synthesis of  $\beta$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub>, as opposed to CuO, because the latter has low solubility (0.00349 M).<sup>37</sup> For the synthesis, metal oxide precursors ZnO, Cu<sub>2</sub>O, and V<sub>2</sub>O<sub>5</sub> were fully dissolved in the urea : choline chloride mixture at 70 °C. According to Abbott *et al.*, [ZnClO·urea]<sup>−</sup> and [VO<sub>2</sub>Cl<sub>2</sub>]<sup>−</sup> species are present in the eutectic urea : choline chloride mixture upon dissolution of ZnO and V<sub>2</sub>O<sub>5</sub> oxides.<sup>37</sup> We observed that the resulting solutions have distinct colors, probably due to various oxidation states of the metal in the metal complex. The [ZnClO·urea]<sup>−</sup> complex produces a colorless liquid, while the [VO<sub>2</sub>Cl<sub>2</sub>]<sup>−</sup> complex is green, suggesting increased *d* orbital splitting of V<sup>5+</sup> causing a blue-shift of the optical absorption. The colors of CuO and Cu<sub>2</sub>O dissolved in the urea : choline chloride mixture are light blue and dark blue, respectively, indicating that the oxidation state of copper ions might be the same in the resulting copper oxide solutions. The solutions were then combined and calcined in open crucibles, allowing for the removal of the deep eutectic solvent in the form of gaseous NH<sub>3</sub>, HCl, and CO<sub>2</sub> followed by crystallization of ternary oxides. Samples were first heated at 230 °C for  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> and 300 °C for  $\beta$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub> to remove NH<sub>3</sub> (g) and HCl (g) and to avoid



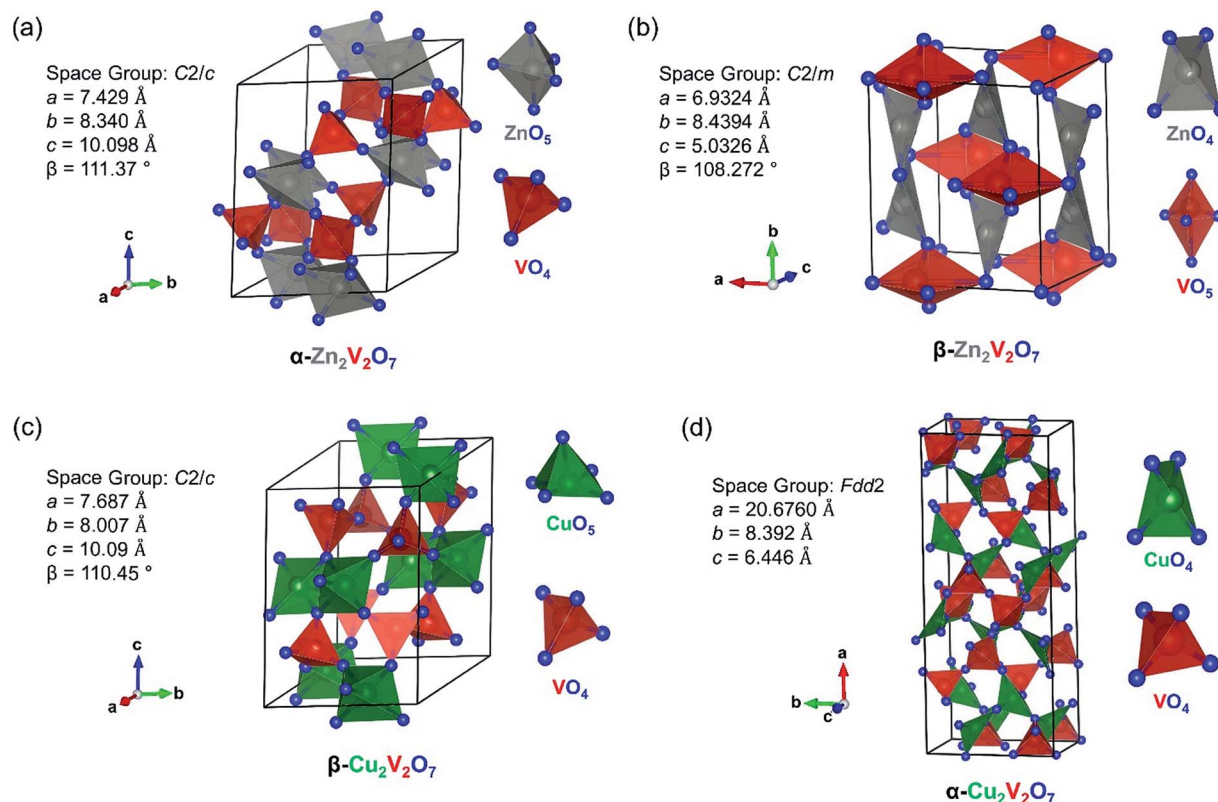


Fig. 1 Structure of (a)  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  (ICSD no. 2886), (b)  $\beta$ - $\text{Zn}_2\text{V}_2\text{O}_7$  (ICSD no. 250002), (c)  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  (ICSD no. 23479) and (d)  $\alpha$ - $\text{Cu}_2\text{V}_2\text{O}_7$  (ICSD no. 40973). Zn: grey, Cu: green, V: red, and O: blue.

extensive foaming. Formation of crystalline  $\text{NH}_4\text{Cl}$  in the ventilation system supports the hypothesis. The gas generated from this first heating step made a pH paper turn dark green and then the liquid sample became a tar-like product, suggesting that a basic mixture of gaseous  $\text{NH}_3$  and  $\text{HCl}$  escaped first and the leftover hydrocarbon was eliminated upon further heating.  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  was synthesized at three different calcination temperatures, 500 °C, 600 °C and 700 °C, while calcination at 400 °C results in an amorphous product. Powder X-ray diffraction peaks are indexed in accordance with  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  (ICSD no. 2886) with a minor amount of impurity ( $\sim 2$  wt%) of  $\text{ZnV}_2\text{O}_6$  as shown in Fig. 2a.  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  was synthesized by a similar route, but lower calcination temperatures (400 °C and 500 °C) were employed. Powder X-ray diffraction peaks are indexed to  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  (ICSD no. 23479) as shown in Fig. 2b. It is notable that  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  was obtained even at a low temperature of 400 °C, at which crystalline  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  was not observed. Moreover, the mild synthetic conditions yield a phase-pure crystalline  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  unlike a recent report on a solution combustion synthesis, in which  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  was synthesized at 500 °C with impurities (3.0%  $\alpha$ - $\text{Cu}_2\text{V}_2\text{O}_7$  and 6.5%  $\alpha$ - $\text{CuV}_2\text{O}_6$ ).<sup>38</sup>

The recent work by Söldner *et al.* reports on a DES-assisted synthesis of spinel-type ferrites  $\text{MFe}_2\text{O}_4$  ( $\text{M} = \text{Mg}, \text{Co}$  and  $\text{Ni}$ ) from stable binary metal oxide precursors,<sup>39</sup> which normally would require a high annealing temperature for the solid-state reaction to form  $\text{MFe}_2\text{O}_4$ . Using a choline chloride and maleic acid mixture as a solvent, ternary ferrites were synthesized at

a lower temperature (500 °C and 600 °C) as compared to the solid-state reaction.<sup>39</sup> Our results demonstrate that the synthesis route employing the deep eutectic solvent as a reaction medium can be extended to  $\text{M}_2\text{V}_2\text{O}_7$  ( $\text{M} = \text{Zn}$  and  $\text{Cu}$ ) systems.

Further insight into thermal stability and optimized synthesis conditions of zinc and copper vanadates was obtained from *in situ* high-temperature powder X-ray diffraction (HT PXRD) and DSC/TGA. It is crucial to remove the urea : choline mixture before the thermal stability study, so the transformation of oxides is not hindered by the thermal decomposition of the deep eutectic solvent. Thus, the pre-calcined powdered samples were used in these experiments (calcined at 500 °C for zinc vanadates and 400 °C for copper vanadates). Fig. 3 shows the evolution of powder X-ray diffraction patterns of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  powders upon heating followed by cooling (with a minor impurity of  $\text{ZnV}_2\text{O}_6$ ,  $\sim 2$  wt%). Upon heating to 615 °C, no change in PXRD patterns was observed except the shift of Bragg peaks because of the thermal expansion.  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  transformed into  $\beta$ - $\text{Zn}_2\text{V}_2\text{O}_7$  above 615 °C and remained the same until 914 °C, above which the crystalline phase disappeared due to melting. Upon cooling the melt from 955 °C,  $\beta$ - $\text{Zn}_2\text{V}_2\text{O}_7$  appeared as a main phase below 718 °C and transformed back to  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  at 590 °C. DSC/TGA (Fig. 4) of the same sample corroborated these findings but allowed more accurate determination of the transition temperatures.

Upon heating, the first sharp endothermic peak was observed at 602 °C, which corresponds to the  $\alpha \rightarrow \beta$  transformation of





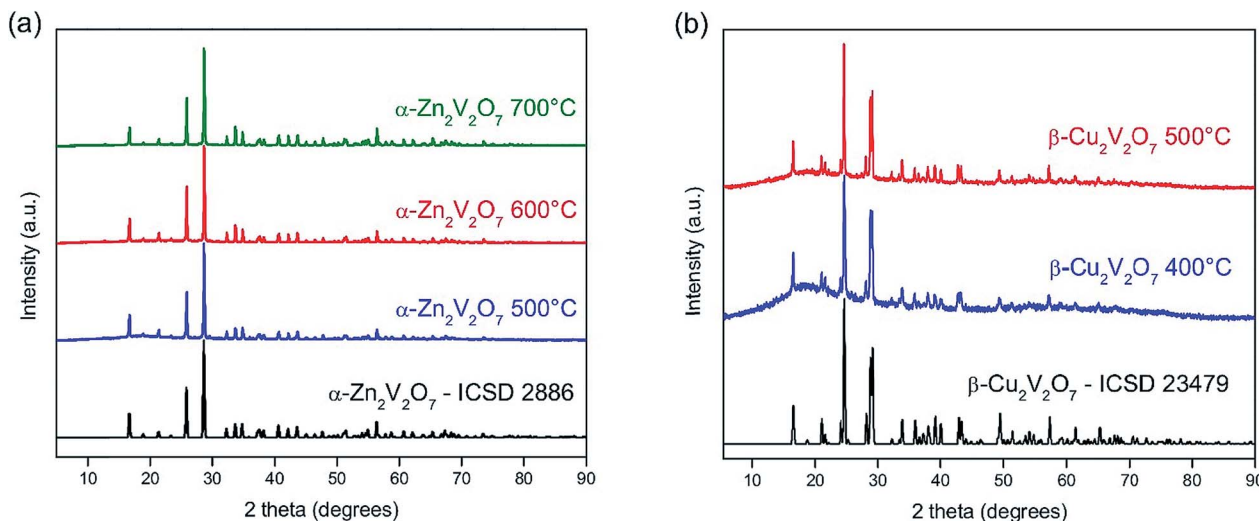


Fig. 2 PXRD patterns of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  synthesized at 500, 600 and 700 °C (a) and  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  synthesized at 400 and 500 °C (b) together with calculated patterns using experimentally determined crystal structures.<sup>32,33</sup>

$\text{Zn}_2\text{V}_2\text{O}_7$ . The onset temperature of the second endothermic peak at 850 °C indicates gradual melting of  $\beta$ - $\text{Zn}_2\text{V}_2\text{O}_7$ . Upon cooling, the first exothermic peak was observed at 795 °C, which is due to the formation of crystalline  $\beta$ - $\text{Zn}_2\text{V}_2\text{O}_7$  from the melt. The second exothermic peak at 566 °C is indicative of the  $\beta$  to  $\alpha$  phase transformation of  $\text{Zn}_2\text{V}_2\text{O}_7$ . The bump from 309 to 533 °C in the DSC curve is due to the sample deposited at the bottom of the crucible, which is in contact with a thermocouple. The deviation of temperatures at which thermal events occurred during the *in situ* PXRD and DSC/TGA could be due to different heating/cooling rates used during the experiment. We believe that the DSC/TGA curves provide a more accurate temperature of thermal events due to slower heating and cooling rates as well as because the bulk effect is being registered as opposing to HT PXRD results. The phase transformation temperature determined here is in accordance with the phase diagram established

by Kurzawa *et al.*: 590 °C for the  $\alpha \rightarrow \beta$  transformation and above 855 °C for the melting of  $\beta$ - $\text{Zn}_2\text{V}_2\text{O}_7$ .<sup>40</sup> Additionally, there was no variation in the TG signal upon heating/cooling of the sample, suggesting that the sample calcined at 500 °C does not contain any carbon-based impurities from the urea : choline chloride

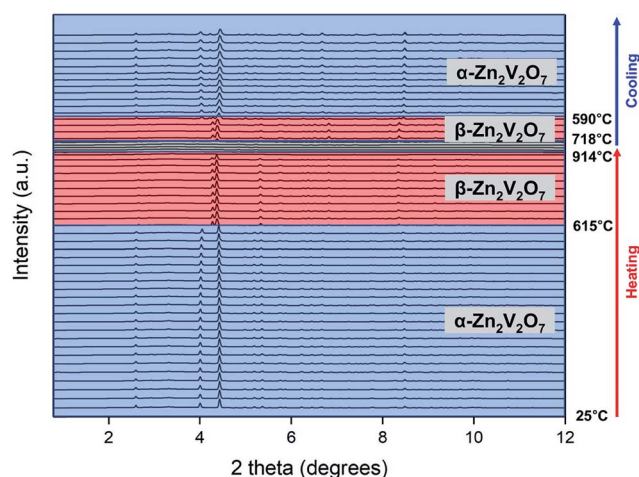


Fig. 3 *In situ* high-temperature powder X-ray diffraction (HT PXRD) patterns of  $\text{Zn}_2\text{V}_2\text{O}_7$ .

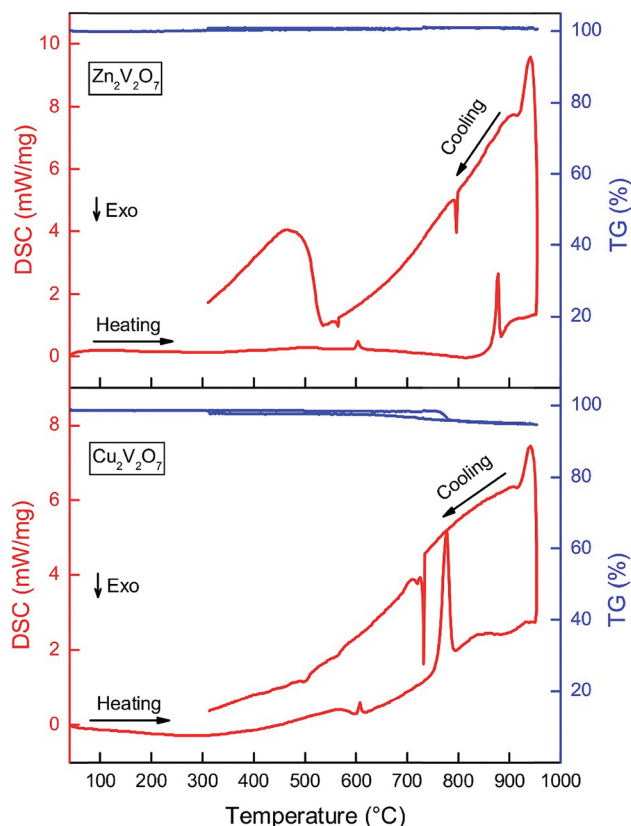
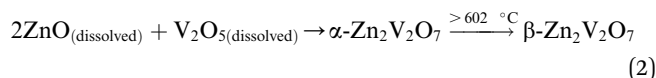
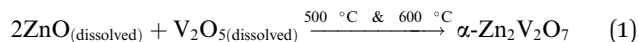


Fig. 4 DSC/TGA of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  (top) and  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  (bottom) samples measured up to 900 °C.

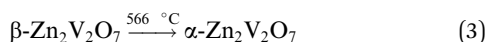


mixture. Samples calcined at a lower temperature (350 °C) have about 87% mass loss and several exothermic peaks between 400 °C and 700 °C corresponding to the elimination of the carbon-based solvent by a combustion reaction. Thus, the suggested reaction process is as follows:

Heating:



Cooling:



Upon heating, the dissolved oxide precursors react to form  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$ , which further transforms into  $\beta\text{-Zn}_2\text{V}_2\text{O}_7$ , at  $\sim 602$  °C. The  $\beta$ -form is present at a calcination temperature of 700 °C (2). The  $\beta \rightarrow \alpha$  transformation upon natural cooling resulted in  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$  (3). A quenching experiment was performed to stabilize  $\beta\text{-Zn}_2\text{V}_2\text{O}_7$ ; however, the phase present in the product was  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$ .

$\text{Cu}_2\text{V}_2\text{O}_7$  goes through a more complicated phase transformation and is less thermally stable than  $\text{ZnV}_2\text{O}_7$ , as determined by HT PXRD (Fig. S1†) and DSC/TGA (Fig. 4). The first endothermic peak of the DSC data was observed at 600 °C, which was attributed to the  $\beta \rightarrow \alpha + \gamma$  transformation together with partial decomposition of the  $\beta$ -phase into  $\text{Cu}_{1.5}\text{V}_{12}\text{O}_{29}$  and presumably amorphous copper oxide. The weak signal (onset temperature of 712 °C in the DSC data) is due to melting of the  $\alpha$ -phase, which can be understood by the low enthalpy value from the small amount of the  $\alpha$ -phase.<sup>41</sup> The third endothermic peak at 750 °C corresponds to the melting of the  $\gamma$ -phase and  $\text{Cu}_{1.5}\text{V}_{12}\text{O}_{29}$ , accompanied by 3% mass loss as evident from TGA data. Since no additional signals were observed in mass spectrum/FTIR data in this temperature range, the mass loss was attributed to oxygen evolution upon melting. Upon cooling, the first exothermic peak at 735 °C is due to partial crystallization of  $\gamma\text{-Cu}_2\text{V}_2\text{O}_7$ ,<sup>41</sup> followed by a weak exothermic peak at 720 °C from the  $\gamma \rightarrow \alpha$  transformation. Interestingly, the sample mass was regained during the crystallization process upon cooling, suggesting that oxygen atoms were restored in the crystal structure. The cooling process is complicated with multiple weak DSC signals between 400 °C and 600 °C because  $\gamma\text{-Cu}_2\text{V}_2\text{O}_7$  melts peritectically and low-melting  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$ ,  $\text{CuVO}_3$  and other vanadates containing mono- and divalent copper crystallize.<sup>41</sup>

Scanning electron microscopy (SEM) was used to characterize the morphology and microstructure of  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$  and  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$  samples obtained by calcination at various temperatures (Fig. 5).  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$  powders have pebble-shaped morphologies with sizes in the ranges 1–3  $\mu\text{m}$ , 2–20  $\mu\text{m}$ , and 4–20  $\mu\text{m}$  for the samples synthesized at 500 °C, 600 °C, and 700 °C, respectively.  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$  has a similar particle shape with a size of 0.1–0.5  $\mu\text{m}$  and 1–7  $\mu\text{m}$  for the samples synthesized at 400 °C and 500 °C, respectively. The average particle sizes were

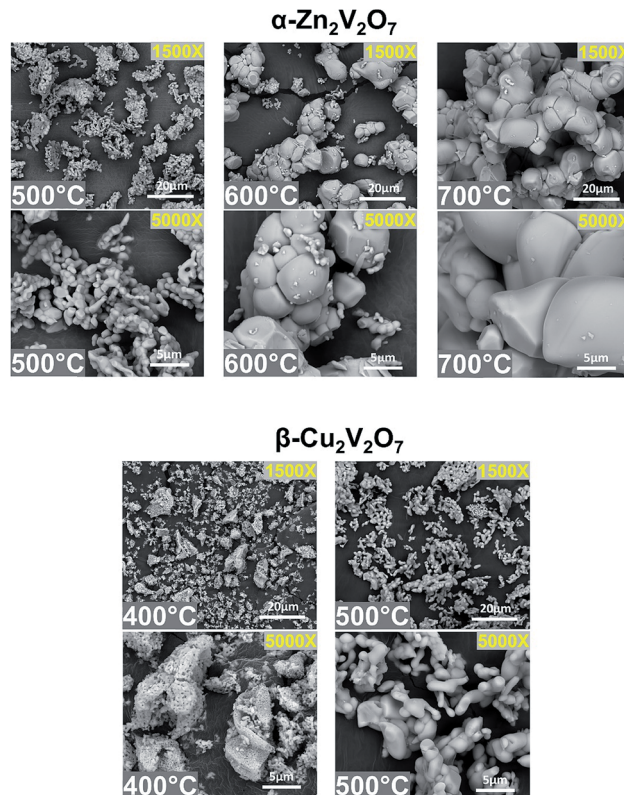


Fig. 5 SEM images of  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$  calcined at 500 °C, 600 °C, and 700 °C (from left to right) and  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$  calcined at 400 °C and 500 °C (from left to right).

estimated using the Debye-Scherrer equation, using the full width at half-maximum (FWHM) values of the three most intense diffraction peaks (Table S1†). Both methods reveal that the particle size increases as the annealing temperature increases from 500 °C to 600 °C for  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$  and from 400 °C to 500 °C for  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$ . Energy dispersive X-ray spectroscopy (EDS) confirms the absence of metals other than Cu, V, and Zn, and additionally the Zn/V and Cu/V ratio was determined to agree with the stoichiometric ratio 2 : 2 (Table S1†). Thus, higher temperature annealing leads to larger particles without altering the composition.

X-ray photoelectron spectroscopy (XPS) was carried out to understand the chemical composition and the chemical state of metals and oxygen on the surface of  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$  and  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$ . Fig. 6 shows the O 1s and V 2p core level spectra of  $\alpha\text{-Zn}_2\text{V}_2\text{O}_7$ . The O 1s region is composed of two components, with a narrow peak at 529.7 eV attributed to the lattice oxygen and a broader peak at 531.5 eV due to oxygen defects.<sup>42</sup> Interestingly, the concentration of oxygen defects gradually increases as the annealing temperature increases from 500 °C to 700 °C (Table 1), suggesting that high temperature annealing under basic conditions with evolution of ammonia promotes oxygen vacancy formation. The two vanadium peaks originate from spin orbit-splitting of V 2p<sub>3/2</sub> and V 2p<sub>1/2</sub>,<sup>43</sup> while each peak is a superposition of two peaks corresponding to V<sup>5+</sup> and V<sup>4+</sup>. Two fitted peaks at 517.1 eV (V 2p<sub>3/2</sub>) and 524.5 eV (V 2p<sub>1/2</sub>)<sup>43</sup> are from



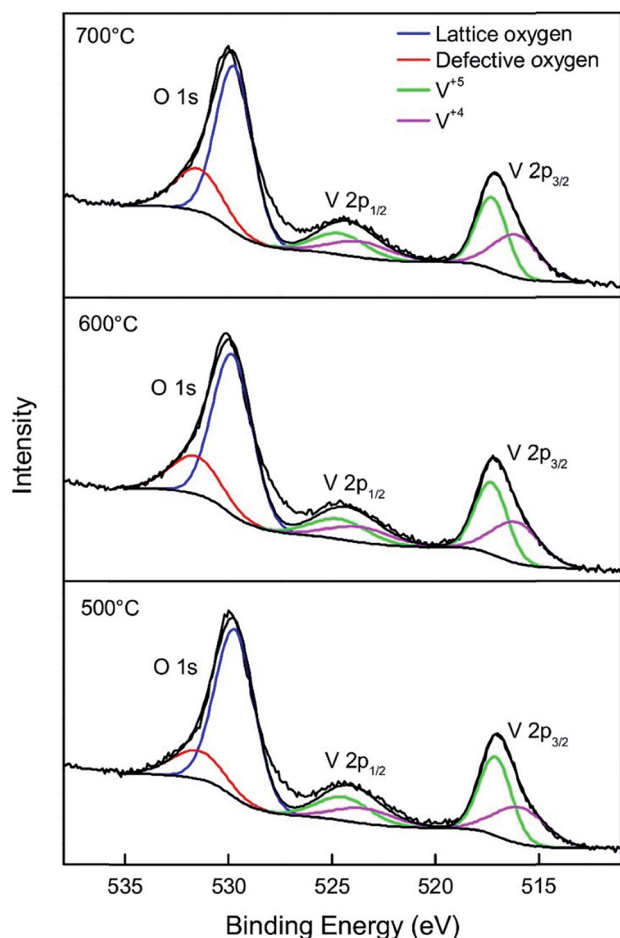


Fig. 6 XPS data of the O 1s and V 2p regions of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$ .

$\text{V}^{5+}$ , while peaks located at 515.9 eV ( $\text{V } 2p_{3/2}$ ) and 523.6 eV ( $\text{V } 2p_{1/2}$ )<sup>43</sup> are attributed to  $\text{V}^{4+}$ , which needs to be present to compensate the charge imbalance from oxygen vacancies. Unsurprisingly, the concentration of  $\text{V}^{4+}$  was the highest in  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  annealed at 700 °C (Table 1). Yan *et al.* and Li *et al.* also prepared  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  and performed XPS analysis, however, only  $\text{V}^{5+}$  was identified.<sup>25,44</sup> Thus, we believe that  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  is stable

under ultra-high vacuum conditions, and the partial reduction of  $\text{V}^{5+}$  stems from the oxygen vacancy created by the synthesis method utilized. Peak positions of Zn  $2p_{3/2}$  and  $2p_{1/2}$  are in agreement with those of ZnO (Fig. S2†),<sup>43</sup> indicating that the divalent oxidation state of zinc remains unchanged due to the completely filled d-subshell.

XPS spectra (Fig. 7) show the chemical states of  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ . The O 1s core level spectra (Fig. 7a) are identified as the superposition of two peaks located at 529.6 eV and 531.5 eV. The narrow peak at 529.6 eV is attributed to lattice oxygen, and the broad peak at 531.5 eV originates from oxygen defects.<sup>42</sup> It was noticed that the concentration of oxygen defects in  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  shows a similar trend to that in  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$ : a greater amount of oxygen vacancies was observed in the sample synthesized at a higher temperature (Table 1). The shake-up satellite peaks of Cu 2p (939–944 eV and 958–965 eV) (Fig. 7b), in agreement with the CuO spectrum,<sup>43</sup> indicate the presence of  $\text{Cu}^{2+}$ . Both  $2p_{3/2}$  and  $2p_{1/2}$  peaks of Cu have a shoulder peak at a lower binding energy (931.6 eV and 951.3 eV, respectively), which is due to  $\text{Cu}^{1+}$  formed to compensate oxygen vacancies.<sup>43</sup> The concentration of  $\text{Cu}^{1+}$  increases from  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  synthesized at 400 °C to 500 °C due to the greater charge imbalance created by the oxygen defects. Two peaks located at 516.3 eV and 523.8 eV are attributed to  $\text{V } 2p_{3/2}$  and  $\text{V } 2p_{1/2}$  of  $\text{V}^{5+}$ , respectively (Fig. S3†).<sup>43</sup> Peak splitting between  $\text{V } 2p_{3/2}$  and  $\text{V } 2p_{1/2}$  is about 7.3 eV, which agrees with the reported value.<sup>43</sup> Kim *et al.* recently prepared  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  by electrostatic deposition (ESD) and showed the presence of  $\text{Cu}^{2+}$  without  $\text{Cu}^{1+}$  in its XPS spectrum.<sup>45</sup> In addition, the temperature programmed desorption of  $\text{O}_2$  from  $\text{Cu}_2\text{V}_2\text{O}_7$  performed by Machida *et al.* reveals that  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  does not release oxygen until above 600 °C under ultra-high vacuum.<sup>46</sup> Therefore, we believe that the reduction of  $\text{Cu}^{2+}$  is a result of oxygen vacancies created by the basic reaction conditions. Also, due to the fact that the reduction potential of  $\text{V}^{5+}$  is higher than that of  $\text{Cu}^{2+}$ , the charge imbalance was compensated by  $\text{Cu}^{1+}$ , unlike  $\text{ZnV}_2\text{O}_7$  (*vide supra*).

To further confirm the presence of oxygen-deficient sites in  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  synthesized at 500 °C, time-of-flight (TOF) neutron powder diffraction ( $\lambda = 0.8 \text{ \AA}$  and  $2.665 \text{ \AA}$ ) data were collected.

Table 1 Selected XPS data for  $\text{M}_2\text{V}_2\text{O}_7$  (M = Zn and Cu)

$\text{M}_2\text{V}_2\text{O}_7$	Annealing temperature (°C)	% concentration		
		Oxygen defects <sup>a</sup>	( $\text{V}^{4+}$ ) <sup>b</sup>	( $\text{Cu}^{1+}$ ) <sup>c</sup>
$\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$	500	18.12	42.44	—
	600	23.34	45.77	—
	700	25.35	47.09	—
$\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$	400	39.45	—	22.67
	500	49.00	—	26.22

$$^a \text{ \% concentration (O defects)} = \frac{\text{area under the peak with binding energy at 531.5 eV}}{\text{sum of area under peaks with binding energy at 531.5 and 529.7 eV}} \times 100(\%).$$

$$^b \text{ \% concentration (V}^{4+}\text{)} = \frac{\text{area under the peak with binding energy at 515.9 eV}}{\text{sum of area under peaks with binding energy at 515.9 and 517.1 eV}} \times 100(\%).$$

$$^c \text{ \% concentration (Cu}^{1+}\text{)} = \frac{\text{area under the peak with binding energy at 931.6 eV}}{\text{sum of area under peaks with binding energy at 931.6 and 934.0 eV}} \times 100(\%).$$





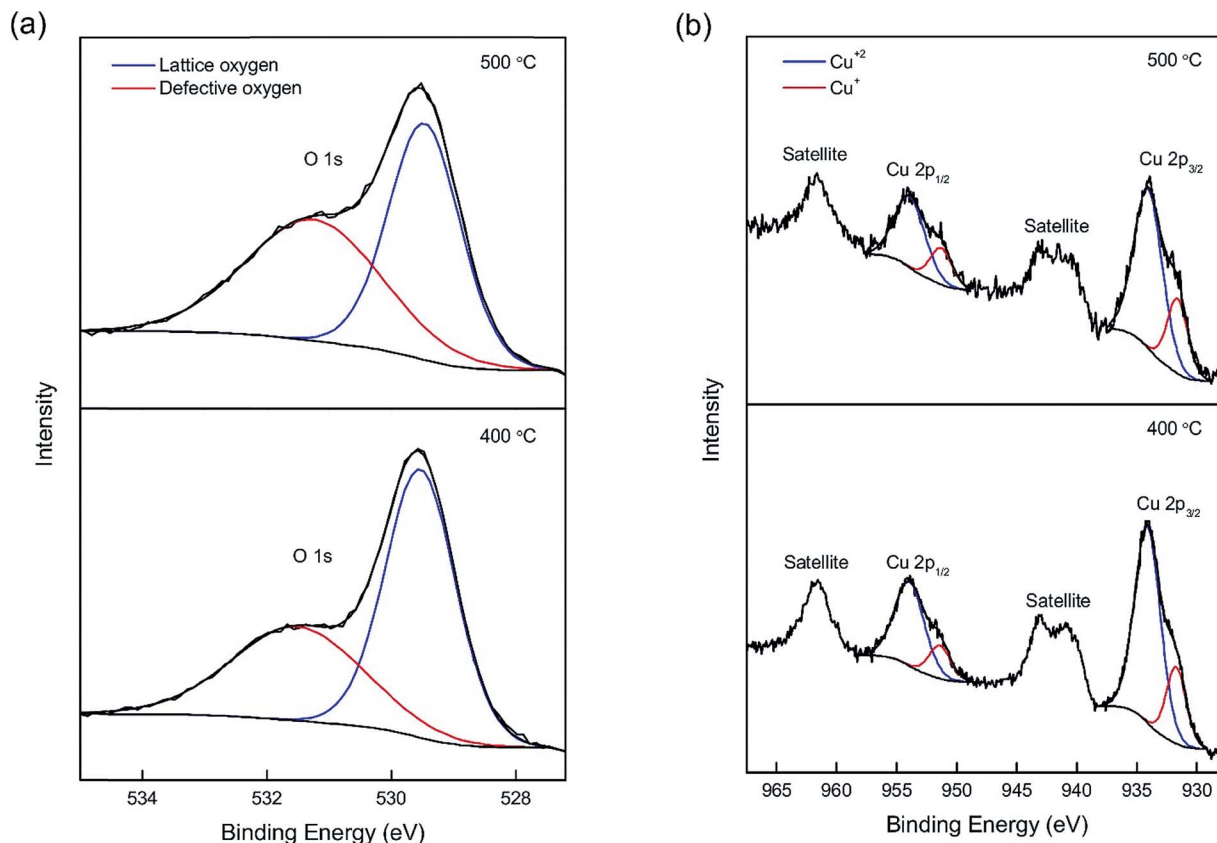


Fig. 7 XPS data of the Cu 2p region (a) and O 1s region (b) of  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ .

Because of the negative neutron scattering length of vanadium, synchrotron X-ray powder diffraction ( $\lambda = 0.412749 \text{ \AA}$ ) data were also collected and refined jointly with the neutron diffraction data (Fig. 8). The Rietveld refinement results indicate a phase pure sample of monoclinic  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ . The refinement of atomic displacement parameters (ADPs) for all atoms was carried out isotropically. In addition, site occupancy factors (SOFs) and atomic displacement parameters (ADPs) were refined in separate refinement cycles to prevent correlations for all atomic sites. The refinement of SOFs for metal sites did not result in any deviation from unity. The refinement of SOFs for four crystallographically independent oxygen sites indicates that they are partially occupied with the O4 sites bearing the largest amount of vacancies (Table 2). The relatively high ADP value for the O1 site (Table 2) is not from significant thermal motion of oxygen but likely from disordered oxygen ion displacements.<sup>47,48</sup> All oxygen positions are corner-shared sites of either vanadium or copper centered polyhedra, such as two  $\text{VO}_4$  tetrahedra sharing O1 sites. Both O2 and O3 are shared between a  $\text{VO}_4$  tetrahedron and a  $\text{CuO}_5$  distorted square pyramid. One  $\text{VO}_4$  tetrahedron and two  $\text{CuO}_5$  distorted square pyramids share the O4 site. This O4 site has a higher concentration of oxygen vacancies, in line with its coordination with two mixed valent Cu centers. Unsurprisingly, the O1 site, which is coordinated to V metal centers, has a site fraction close to unity. It should be noted that when the SOF for all O sites is set to 1, such a refinement resulted in a poorer fit with a high  $R_w$

value, thus further confirming the presence of partially occupied oxygen sites. Similar partial occupancies ( $\sim 0.95(2)$ ) for all oxygen sites were obtained, when the ADPs for all four oxygen sites were constrained to be the same and refined to be  $0.0120(2) \text{ \AA}^2$ . It is noteworthy that the expanded unit cell volume and elongated unit cell parameters ( $a$ ,  $b$ , and  $c$ ) as compared to the reported value ( $V = 581.9 \text{ \AA}^3$ ;  $a = 7.687(5) \text{ \AA}$ ,  $b = 8.007(3) \text{ \AA}$ , and  $c = 10.09(2) \text{ \AA}$  from ICSD no. 23479 as compared to unit cell parameters in Table 2) might also be due to the oxygen defects. Similar enlargement of the unit cell volume was reported for vacancy-containing  $\text{ZnO}$ <sup>49,50</sup> and indium-tin-oxide (ITO) thin films.<sup>51</sup> Moreover, the absence of cation vacancies as confirmed by neutron and synchrotron X-ray diffraction data refinement supports unit cell expansion due to oxygen vacancies.

The composition calculated based on the total oxygen content obtained from the Rietveld refinement is  $\text{Cu}_2\text{V}_2\text{O}_{6.69(2)}$  (Table 2), which is consistent with  $\text{Cu}_2\text{V}_2\text{O}_{6.74}$ , calculated using the  $\text{Cu}^+$  and  $\text{Cu}^{2+}$  ratios from the XPS result and assuming charge balance.

To study the effect of oxygen vacancies and mixed oxidation states of metal centers on the electronic structure, the optical properties of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  and  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  samples were evaluated from diffuse reflectance UV-vis absorption spectroscopy (Fig. 9). A red shift of the main absorption edge in  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  was observed as the annealing temperature increases: 443.5 nm for 500 °C, 471.6 nm for 600 °C, and 500.6 nm for 700 °C. To determine optical indirect band gaps, the data were plotted as





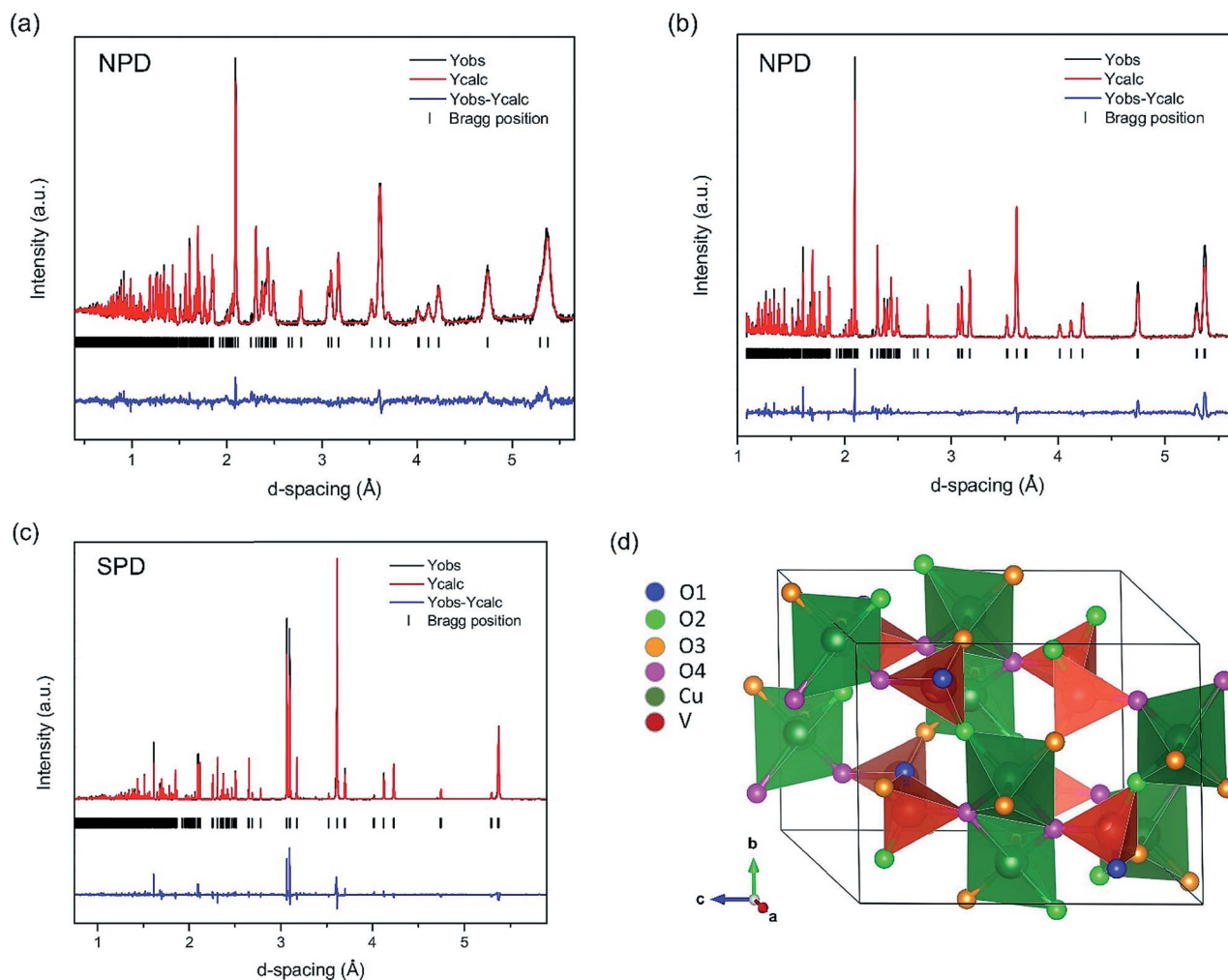


Fig. 8 TOF neutron powder diffraction (NPD) data using (a)  $\lambda = 0.8$  Å and (b)  $\lambda = 2.665$  Å for  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  synthesized at 500 °C. (c) Synchrotron powder diffraction (SPD) data ( $\lambda = 0.412749$  Å) and (d) crystal structure of  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ .

( $\alpha \times h\nu$ )<sup>1/2</sup> vs.  $h\nu$  (Tauc plot), where  $\alpha$  is absorbance and  $h\nu$  is the excitation energy in eV. The estimated bandgaps are 2.63(3) eV, 2.25(3) eV, and 1.97(3) eV for 500 °C, 600 °C, and 700 °C, respectively. The bandgap of  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  annealed at 500 °C is close to the reported value (2.75 eV),<sup>25</sup> but a smaller bandgap

was observed for the samples synthesized at 600 °C and 700 °C. This is due to oxygen vacancy related disorder being responsible for band tailing.<sup>50</sup> Interestingly, an additional absorption edge at 682.3 nm for the 600 °C annealed sample and 674.7 nm for the 700 °C annealed sample is observed, corresponding to mid-

Table 2 Structural results for  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  from combined Rietveld refinement of neutron powder diffraction and synchrotron powder diffraction data<sup>a</sup>

Atom	Wyckoff site	Coordinates			Site occupancy factor (SOF)	$U_{\text{eq.}}, \text{\AA}^2$
		$x/a$	$y/b$	$z/c$		
Cu	8f	0.30950(7)	0.92687(6)	0.51372(6)	1	0.0096(1)
V	8f	0.2753(1)	0.72239(9)	0.21290(8)	1	0.0047(2)
O1	4e	1/2	0.6369(3)	1/4	0.999(4)	0.0254(7)
O2	8f	0.2666(2)	0.9027(2)	0.1336(2)	0.951(3)	0.0171(5)
O3	8f	0.1221(2)	0.5893(2)	0.1041(2)	0.950(3)	0.0102(4)
O4	8f	0.2364(2)	0.7522(2)	0.3708(2)	0.934(3)	0.0067(3)

<sup>a</sup> Space group:  $C2/c$ ,  $a = 7.7014(1)$  Å,  $b = 8.03305(4)$  Å,  $c = 10.1166(2)$  Å,  $\beta = 110.2794(3)^\circ$ ,  $V = 587.079(8)$  Å<sup>3</sup>,  $R_w = 9.36\%$ ,  $R_p = 7.58\%$ , and G.O.F = 3.52.



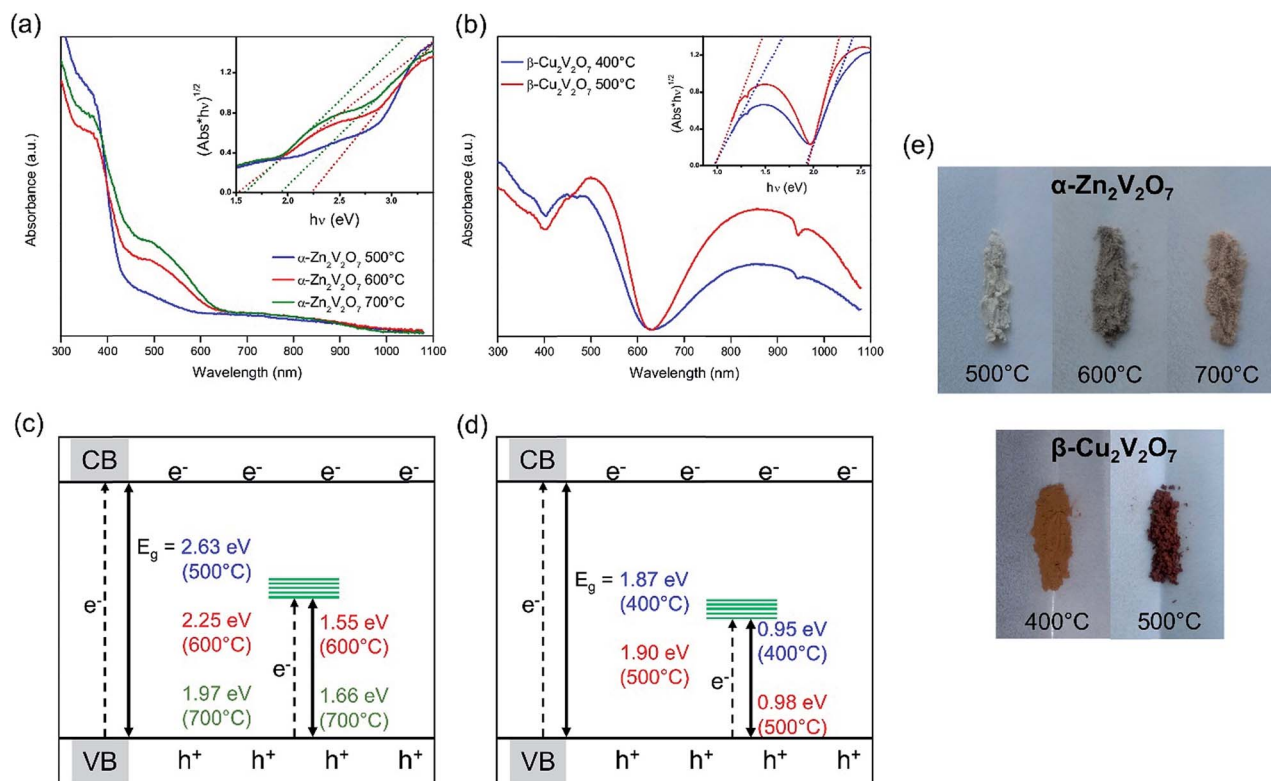


Fig. 9 UV-vis absorption spectra of  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> (a) and  $\beta$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub> (b). The inset shows Tauc plots and their linear fits (dotted lines) to determine band gaps and mid gaps, corresponding to the additional absorption edge. A schematic of band structures (not to scale) of  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> (c) and  $\beta$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub> (d), showing band gaps and mid-gap states. Optical images of  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> and  $\beta$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub> (e).

gap states at 1.55 eV and 1.66 eV, respectively. This additional absorption in the visible-light range is attributed to the d-states from the reduced metal centers ( $V^{4+}$ ) generated to compensate oxygen vacancies. Similar to the band tailing originating from the oxygen defects,<sup>30</sup> the mid-gap state position changes with different reaction conditions. This could be due to different mid-gap defect densities of the reduced metal centers affecting the bandwidth of the mid-gap states.

UV-vis diffuse reflectance spectra of  $\beta$ -Cu<sub>2</sub>V<sub>2</sub>O<sub>7</sub> show the main absorption edge at 615.0 nm and 615.5 nm for the samples synthesized at 400 °C and 500 °C, respectively (Fig. 9). The estimated band gaps of 1.90(3) eV (400 °C) and 1.87(3) eV (500 °C) are smaller than the reported values of 2.20 eV (ref. 52) and 2.22 eV.<sup>38</sup> Similar to  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub>, there is an additional absorption edge at a higher wavelength: 1166.2 nm for the 400 °C sample and 1150.5 nm for the 500 °C sample, corresponding to mid-gap states at 0.95 eV and 0.98 eV, respectively. The additional absorption and the distinct color difference between the 400 °C and 500 °C samples are due to the mid-gap states located in between the valence band and the conduction band contributed by d-states from the reduced metal centers ( $Cu^{+}$ ). Also, it is observed that the 500 °C sample exhibits higher absorption in the wavelength range from 700 nm to 1100 nm compared to the 400 °C sample, suggesting that a greater amount of oxygen vacancies is present in the 500 °C sample, thus a greater fraction of  $Cu^{+}$ , consistent with XPS results.

Surface photovoltage spectroscopy (SPS) measurements were conducted to probe the ability of the vanadates to generate and separate photochemical charge carriers. In SPS, the contact potential difference (CPD) of a sample film on top of a conductive substrate is recorded with a contactless Kelvin probe as a function of the incident photon energy (Fig. 10).<sup>53–55</sup> The change of the CPD under illumination corresponds to a photovoltage. The sign, onset, and size of this voltage provide information about the majority carrier type, the effective band gap, defects, and the photochemistry of the sample, as detailed in previous studies.<sup>54–61</sup> SPS data for the vanadate powders on FTO substrates are shown in Fig. 10. All samples generate a negative photovoltage caused by majority charge carrier injection into the substrate. This confirms that all vanadates are n-type semiconductors. Using the tangent method, the effective bandgaps are estimated from the major photovoltage feature to be 2.85 eV, 2.81 eV, and 2.95 eV for the  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> samples annealed at 500 °C, 600 °C and 700 °C, respectively. These onset values exceed the optical bandgaps of these samples, especially at higher annealing temperatures. The 600 °C and 700 °C annealed samples also produce much smaller photovoltages than the  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> film annealed at 500 °C. This suggests that the photovoltaic properties of  $\alpha$ -Zn<sub>2</sub>V<sub>2</sub>O<sub>7</sub> are degraded by the higher temperature annealing step. Tentatively, we attribute these observations to electron trapping and/or charge recombination at 1.55 eV and 1.66 eV mid gap states of  $V^{4+}$  that are seen in the optical spectra of the 600 °C and 700 °C annealed



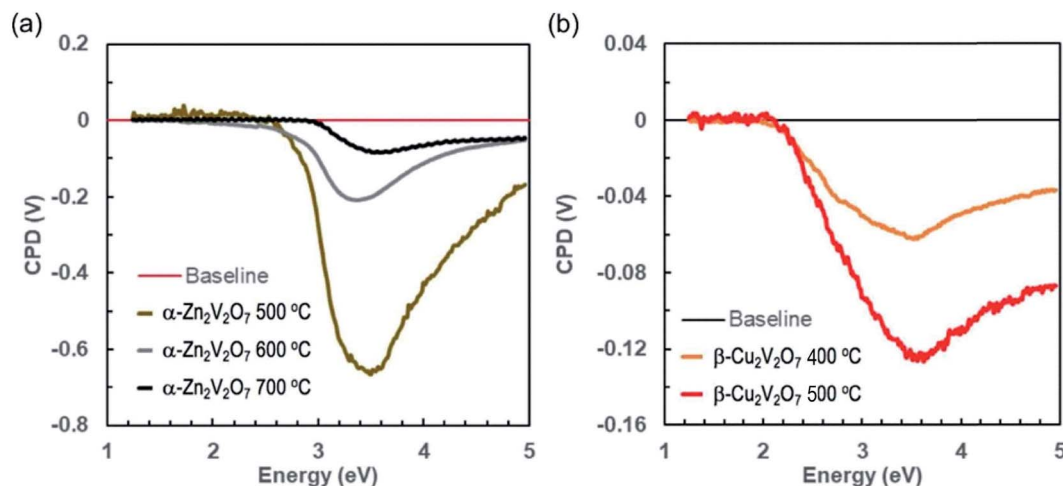


Fig. 10 Surface photovoltage spectra (SPS) of (a)  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$  and (b)  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$  on FTO substrates in an air atmosphere ( $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ ) and under vacuum ( $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$ ). Contact potential difference (CPD) values are shown relative to those in the dark.

samples. Trap states near the absorber mid gap energy are known to be particularly effective for recombination because they are equally accessible to electrons and holes.<sup>62–64</sup> The SPS data for copper vanadates on FTO (Fig. 10b) resemble those of zinc vanadates; however, the effective band gap of 2.12 eV for both samples agrees relatively well with the optical band gaps. But again, the photovoltage values achieved with the 400 °C and 500 °C annealed copper vanadates (−0.06 V and −0.12 V at 3.6 eV) are much smaller than that for the 500 °C annealed  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_7$ . The photovoltage appears to be limited by electron trapping and/or recombination at the 0.95 eV  $\text{Cu}^+$  mid gap state. This behavior is analogous to  $\text{CuWO}_4$ , whose photocatalytic properties are also diminished by electron trapping at the 1.8 eV  $\text{Cu}^{2+}$  mid gap-state.<sup>65</sup> Cathodic photocorrosion *via* electron trapping at copper d states is a general property of copper containing metal oxides.<sup>66–68</sup>

## Conclusion

Ternary metal oxides  $\text{M}_2\text{V}_2\text{O}_{7-\delta}$  ( $\text{M} = \text{Zn}$  and  $\text{Cu}$ ) were synthesized using a deep eutectic solvent (DES): a urea : choline chloride mixture. Based on thermal stability of the products, their syntheses were optimized. We have shown for the first time the presence of oxygen vacancies and reduced oxidation states of metal ions in ternary metal oxides synthesized by this route. The mixed-valence oxidation states of vanadium ( $\text{V}^{4+}$  vs.  $\text{V}^{5+}$ ) in  $\alpha$ - $\text{Zn}_2\text{V}_2\text{O}_{7-n}$  and copper ( $\text{Cu}^+$  vs.  $\text{Cu}^{2+}$ ) in  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_{7-m}$  corroborate partial reduction of the metal ions needed for charge compensation due to the oxygen vacancies. Further investigation of oxygen deficiencies in  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_{7-m}$  by neutron powder diffraction reveals that the oxygen site coordinated to two copper metal centers possesses a significant number of oxygen vacancies. By simply changing the annealing temperature of metal precursors in the DES, various concentrations of oxygen vacancies are achieved, thereby tuning the electronic structure of metal oxides. A high concentration of oxygen vacancies in  $\text{M}_2\text{V}_2\text{O}_{7-\delta}$  ( $\text{M} = \text{Zn}$  and  $\text{Cu}$ ) leads to narrowed band

gaps and it generates mid-gap states of reduced metal centers in between the valence band and conduction band. Surface photovoltage spectroscopy shows that the mid-gap states reduce the photovoltage of the materials, indicating that  $\text{Cu}^+$  and  $\text{V}^{4+}$  ions can serve as electron/hole trap and recombination sites. This limits solar energy conversion applications of the vanadates. Most importantly, the results demonstrate the potential of deep eutectic solvents for the synthesis of semiconducting metal oxides containing oxygen vacancies.

## Conflicts of interest

There are no conflicts to declare.

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