

Cite this: *J. Mater. Chem. A*, 2020, **8**, 12538

# Elucidating the optical, electronic, and photoelectrochemical properties of p-type copper vanadate (p-Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub>) photocathodes†

Angang Song,<sup>ab</sup> Sean P. Berglund,<sup>a</sup> Abdelkrim Chemseddine,<sup>a</sup> Dennis Friedrich,<sup>a</sup> Fatwa F. Abdi<sup>a</sup> and Roel van de Krol<sup>ab\*</sup>

P-type copper vanadate (Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub>) photoelectrodes made by spray pyrolysis were developed and evaluated as a potential photocathode material for photoelectrochemical (PEC) water splitting. Using fluorine-doped tin oxide as a substrate, highly phase-pure p-Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub> thin films were obtained after annealing at 550 °C for 4 hours in air. Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub> has a small bandgap energy in the range of 1.8–2.0 eV, allowing it to absorb visible light and making it potentially interesting for solar water splitting applications. The p-Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub> films were characterized by photoelectrochemical techniques in order to provide insight into the critical PEC properties such as the flat-band potential, chemical stability, and incident photon-to-current efficiency (IPCE). The best-performing films showed a photocurrent density of up to 0.5 mA cm<sup>-2</sup> under AM1.5 simulated sunlight, and an IPCE value up to 14% for 450 nm light at 0.8 V<sub>RHE</sub> with H<sub>2</sub>O<sub>2</sub> as an electron scavenger. Despite the narrow band gap and suitable conduction band edge position for PEC H<sub>2</sub> production, these p-type films were unstable under constant illumination in aqueous electrolyte (pH 6.8) due to the reduction and dissolution of Cu. Based on our findings, the suitability of Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub> as a photocathode material for photoelectrochemical water splitting is critically discussed.

Received 21st April 2020  
Accepted 11th June 2020

DOI: 10.1039/d0ta04250a

rsc.li/materials-a

## Introduction

The direct storage of solar energy in the form of chemical bonds by photoelectrochemical (PEC) water splitting systems has been regarded as a promising way to replace fossil fuels.<sup>1–4</sup> In particular, the reduction of water to produce hydrogen, a clean and sustainable fuel, can be driven by p-type photoelectrodes with suitable band positions. Among all classes of materials investigated as photoelectrodes for solar water splitting, metal oxide semiconductors are particularly attractive mainly due to their relative abundance and low cost as well as their resistance to further oxidation in aqueous solutions. Currently, the highest performing p-type metal oxide photoelectrode is cuprous oxide (Cu<sub>2</sub>O); a photocurrent as high as ~10 mA cm<sup>-2</sup> and an onset potential as high as 1 V vs. RHE have been reported.<sup>5</sup> However, stability remains a major issue. Although it has been demonstrated that the stability can be enhanced by depositing conformal protection layers on top of the Cu<sub>2</sub>O photocathodes,<sup>6</sup> the coating needs to be sufficiently thin (to minimize ohmic

losses) and pinhole-free (to prevent degradation). This complicates the deposition process and potentially increases the cost of the photoelectrodes, which further motivates the search for an intrinsically stable p-type photoelectrode. One particular approach is to look for alternative Cu-based ternary metal oxides with a second cation (besides Cu<sup>2+</sup> or Cu<sup>+</sup>) forming the conduction band. This is expected to inhibit the reduction of Cu<sup>2+</sup> or Cu<sup>+</sup> to Cu at the semiconductor/electrolyte interface, which is the main degradation mechanism in Cu<sub>2</sub>O.<sup>6</sup> Indeed, copper niobates (CuNbO<sub>3</sub>) and copper ferrites (CuFeO<sub>2</sub>), in which Nb and Fe d orbitals dominate the conduction band character, show better stability compared to Cu<sub>2</sub>O.<sup>7–11</sup>

Another possible candidate is copper vanadate (Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub>), a CuO-based material that is expected to be chemically robust. It has suitable indirect and direct optical band gaps in the range of 1.8–2.0 eV and a moderate absorption coefficient ( $\alpha \approx 10^4$  cm<sup>-1</sup>),<sup>12</sup> allowing for the utilization of a significant portion of the solar spectrum to generate photocurrent. However, at present, there are only ~20 published reports on Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub>, only 4 of which are relevant to photoelectrochemistry or photocatalysis.<sup>12–14</sup> Therefore, many key physical and photoelectrochemical parameters of this material are not yet known. The dominant conductivity type is also still not clear, since both p-type and n-type conductivity have been reported in the literature.<sup>15–19</sup> Even less is known about the charge carrier transport properties (*e.g.*, mobility and lifetime), despite Cu<sub>5</sub>V<sub>2</sub>O<sub>10</sub> being

<sup>a</sup>Institute for Solar Fuels, Helmholtz-Zentrum Berlin für Materialien und Energie GmbH, Hahn-Meitner-Platz 1, 14109 Berlin, Germany. E-mail: roel.vandekrol@helmholtz-berlin.de

<sup>b</sup>Institut für Chemie, Technische Universität Berlin, Straße des 17. Juni 124, 10623 Berlin, Germany

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ta04250a



a potentially interesting candidate as a photoelectrode material for solar water splitting.

In this work, we have developed an attractive, low-cost spray pyrolysis recipe for the deposition of thin film  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photoelectrodes on FTO-coated glass (F-doped  $\text{SnO}_2$ ). Our spray pyrolysis method has been demonstrated as a means for homogeneously depositing large area photoelectrodes<sup>20</sup> that can be integrated with other components in solar fuels devices. The sprayed  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films show p-type conductivity and have a bandgap of 1.86 eV. We determine the lifetime and mobility of the photogenerated carriers by performing time-resolved microwave conductivity (TRMC) measurements. On the basis of these findings, a first assessment of the suitability of p-type  $\text{Cu}_5\text{V}_2\text{O}_{10}$  as a photoelectrode material for solar water splitting is provided.

## Results and discussion

### Structural and chemical characterization

To investigate the structure and chemical composition of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin film photoelectrodes, X-ray diffraction (XRD), Raman spectroscopy, and XPS measurements were carried out. Details of the experimental methods are included in the ESI.† The X-ray diffractograms of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films are shown in

Fig. 1a. The as-deposited  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film, which was deposited onto the substrate at 300 °C, is structurally amorphous as evident from the XRD pattern (blue curve). The deposition temperature of 300 °C is obviously not enough to crystallize the film. The red curve shows the diffractogram of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin film with post-deposition annealing at 550 °C for 4 hours in air. It matches very well with the reference pattern (JCPDS no. 33-0504), indicating that the film consists of single phase of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  with no discernable crystalline impurities. This is also the case for films with various thicknesses (from 50 to 500 nm), as shown in Fig. S1.† The  $\text{Cu}_5\text{V}_2\text{O}_{10}$  phase has been previously found to crystallize in the monoclinic crystal system (space group  $P2_1/c$ ) with the cell parameters of  $a = 8.3938 \text{ \AA}$ ,  $b = 6.0659 \text{ \AA}$ ,  $c = 16.1601 \text{ \AA}$  and  $\beta = 108.0857^\circ$ .<sup>15,21–23</sup>

Fig. 1b shows the Raman spectra of our sprayed  $\text{Cu}_5\text{V}_2\text{O}_{10}$  samples with different thicknesses. The main Raman bands for  $\text{Cu}_5\text{V}_2\text{O}_{10}$  centered at  $\sim 794$ , 899, and 944  $\text{cm}^{-1}$  were observed for all films, which is fully in agreement with reports in the literature.<sup>13,14</sup> These scattering peaks are associated with tetrahedral  $\text{VO}_4$  stretching modes,<sup>22</sup> whereas peaks at 280, 342 and 512  $\text{cm}^{-1}$  are assigned to the bending of the same structural motif. Tetrahedral  $\text{VO}_4$  is the predominant coordination polyhedron in many metal oxides containing  $\text{V}^{5+}$ .<sup>24–26</sup> No Raman peaks for CuO and  $\text{V}_2\text{O}_5$  are observed, again demonstrating that

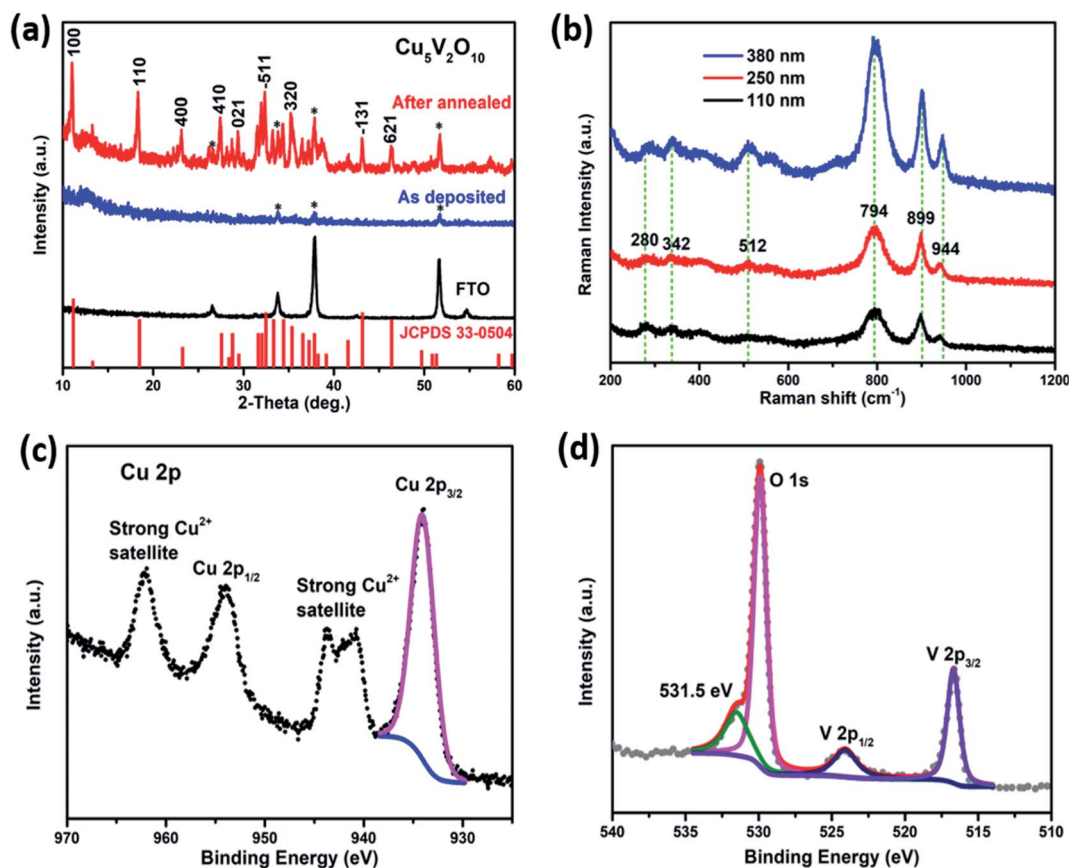


Fig. 1 (a) XRD diffractogram for a  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode synthesized by spray pyrolysis on an FTO substrate. Red vertical lines represent the reference pattern for  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photoelectrode (PDF no. 33-0504) (b) Raman spectra of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films (annealed at 550 °C for 4 hours in air) with different thicknesses. XPS spectra of (c) Cu 2p, (d) V 2p and O 1s for  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film.



our synthesis method yields highly phase-pure thin films of  $\text{Cu}_5\text{V}_2\text{O}_{10}$ .

The core level XPS spectra for Cu 2p, V 2p, and O 1s are compared in Fig. 1c and d. The Cu 2p<sub>3/2</sub> spectrum can be deconvoluted into a single peak with a binding energy of 934.1 eV, and strong satellite peaks in the range of 940–945 eV are observed (Fig. 1c). This suggests that the oxidation state of Cu is +2, which is in agreement with the expected formal oxidation states of copper in  $\text{Cu}_5\text{V}_2\text{O}_{10}$ . The V 2p<sub>3/2</sub> and V 2p<sub>1/2</sub> signal (Fig. 1d) can also be deconvoluted into individual peaks with a binding energies of 516.7 eV and 524.1 eV, respectively, indicating an oxidation state of V<sup>5+</sup> based on reports in the literature.<sup>27</sup> The O 1s peak is located at 529.9 eV, which is expected for a surface lattice oxygen peak in the film. A minor peak at a higher binding energy of 531.5 eV is also present, which corresponds to surface adsorbed water or hydrogen-bearing groups.<sup>28</sup>

### Morphology characterization

Fig. 2a shows the SEM image of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photoelectrode prepared on FTO glass upon post-deposition annealing. The film consists of nanoparticles with diameters of 100–500 nm. The film shows good overall uniformity, and the average thickness of the film is estimated to be ~250 nm using 200 mL precursor solution from the cross-section SEM image shown in Fig. 2b. In comparison with previously reported films prepared by inkjet printing,<sup>13</sup> these films deposited by spray pyrolysis exhibit a significantly more compact and homogeneous morphology. As a comparison, the SEM images of the bare FTO substrate and as-deposited films are shown in Fig. S2;† the as deposited film consists of larger and round grains with some

degree of surface roughness. We also obtained thickness values of ~110 nm and ~500 nm (Fig. S3†) from the samples sprayed with 100 mL and 400 mL of the precursor, which indicates a linear relationship between the precursor volume and film thickness. Fig. 2c shows a photograph of a thin film  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photoelectrode deposited on an FTO-coated glass substrate. The film has a brown color, which appears to be due to the reflected light in the 500–700 nm range, indicating good visible light absorption. Energy dispersive spectroscopy (EDS) was performed on the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin films (see Fig. S4†). The average Cu/(Cu + V) ratio is about  $(0.70 \pm 0.02)$  measured from several films of different thicknesses, which is consistent with the expected value of 0.71 for  $\text{Cu}_5\text{V}_2\text{O}_{10}$ . EDS mapping was also performed on a representative film area as shown in Fig. S5.† The EDS signals from Cu, V, and O appear to be homogeneously distributed over the entire film area.

The surface roughness of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film was measured using atomic force microscopy (AFM); a representative image on a  $5 \times 5 \mu\text{m}^2$  area is shown in Fig. 2d. AFM reveals that the root mean squared (RMS) roughness and real surface area of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films are 44.9 nm and  $27.3 \mu\text{m}^2$ , respectively. A roughness factor (real surface area divided by geometric surface area) of 1.09 is obtained, which confirms that the film is indeed relatively compact. As a comparison, the AFM image of a bare FTO substrate is shown in Fig. S6,† which shows a roughness factor of 1.06.

### Optical properties

Fig. 3a shows the absorbance spectrum of a 250 nm  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin film deposited on a quartz substrate as a function of wavelength (absorbance spectra for films with other

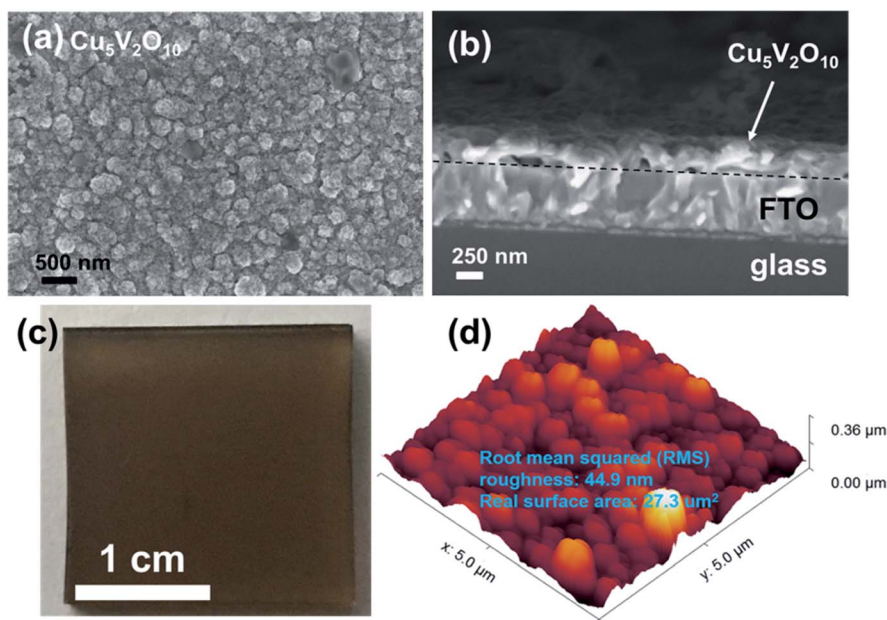


Fig. 2 (a) Top-view and (b) cross-section SEM images of a  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin film deposited on FTO annealed at 550 °C in air for 4 hours. (c) Photograph of a  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin film deposited on an FTO-coated glass substrate. (d) Atomic force microscopy (AFM) image of a ~250 nm  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film deposited on an FTO substrate.



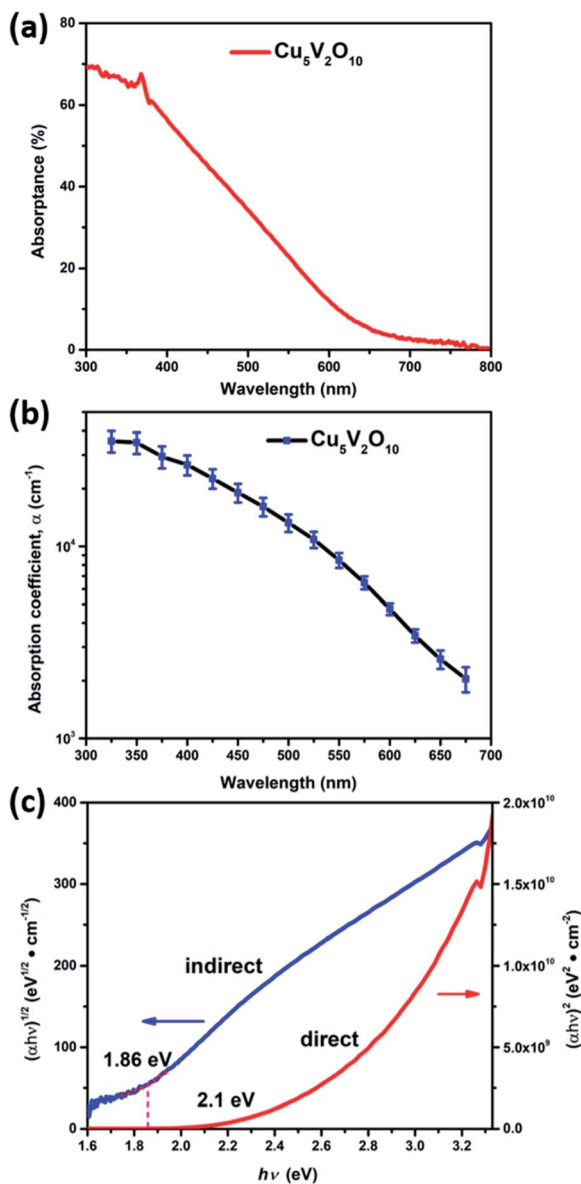


Fig. 3 (a) Absorbance spectra and (b) absorption coefficient for 250 nm thick  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film deposited on quartz substrate. (c) Tauc plots for bandgap analysis of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film. The indirect and direct bandgaps are estimated from inflection point and the intercept of the  $(\alpha h\nu)^{0.5}$  and  $(\alpha h\nu)^2$  curves with the  $x$ -axis, respectively.

thicknesses are shown in Fig. S7a†). The absorption increases sharply at approximately 650 nm, which is in agreement with earlier reports.<sup>12</sup> The weak absorption tail between 700–800 nm arises from localized ligand field excitation at the  $\text{Cu}^{2+}$  ( $3d^9$ ) cations, and therefore does not generate mobile charge carriers.<sup>12</sup> Fig. 3b shows the absorption coefficient ( $\alpha$ ) as a function of wavelength, which was calculated from a fit (for each wavelength) of the absorption data of 5 different film thicknesses by using the following equation:  $\alpha = -\ln(\text{TR})/d$ . TR is the transmittance at a single wavelength and  $d$  is the film thickness determined from cross-section SEM. At 450 nm the absorption coefficient for the film is  $(1.92 \pm 0.21) \times 10^4 \text{ cm}^{-1}$

(Fig. S7b†), which corresponds to a light penetration depth of  $\alpha^{-1} = 521 \text{ nm}$ , *i.e.*, a film thickness of  $1.2 \mu\text{m}$  ( $2.3 \alpha^{-1}$ ) is required to absorb 90% of the incident light having a wavelength of 450 nm. Tauc analysis was performed on the absorption coefficient spectra to determine the optical bandgaps of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  as shown in Fig. 3c. Both direct and indirect optical transitions can be discerned. The indirect and direct bandgaps for the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin film are estimated to be 1.9 and 2.1 eV, respectively, which is promising for visible light absorption for a metal oxide photoelectrode material.<sup>29,30</sup>

### Charge carrier transport

The charge carrier transport properties of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin films were investigated using time-resolved microwave conductivity (TRMC) measurements. For this purpose, the films were deposited on quartz substrates, since a non-conducting substrate is necessary for TRMC. However, only moderate TRMC signals can be detected from the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin film, indicating relatively low photoconductivity. Fig. 4a shows the TRMC signal for a 250 nm  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film under an incident laser pulse intensity of  $3 \times 10^{14} \text{ cm}^{-2}$  at four different excitation wavelengths (410, 450, 500, and 550 nm). Mobility values of  $1.1$ – $2.3 \times 10^{-5} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$  were obtained from the peaks of the

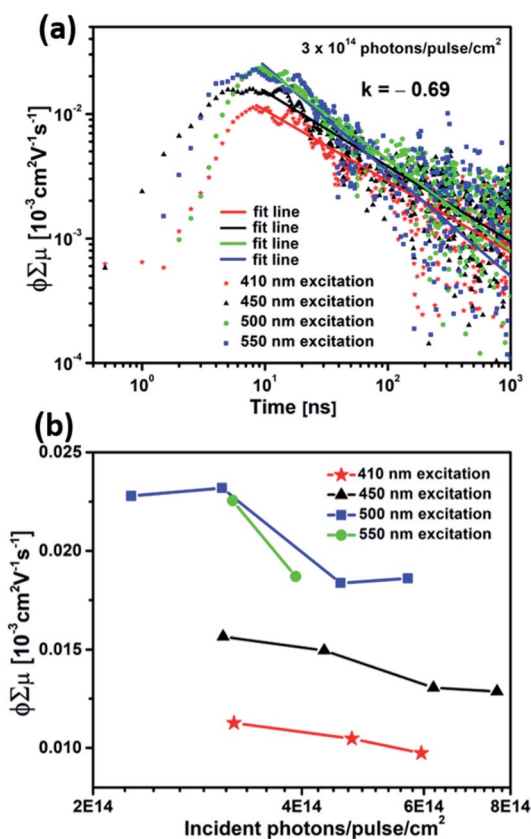


Fig. 4 (a) TRMC signal under various incident laser pulse intensities for a  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film synthesized by spray pyrolysis on a quartz substrate at different wavelengths. (b) Maximum TRMC signal  $(\phi \sum \mu)_{\text{max}}$  vs. incident laser pulse intensity for  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films deposited on quartz substrates under different laser wavelengths.



curves. The decay curves of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  sample can be fitted with a power law relationship ( $\varphi \sum \mu = \gamma \times t^k$ ,  $k = -0.69 \pm 0.1$ ,  $\gamma = 2.3 \times 10^{-10}$ ,  $t < 1000$  ns). The power-law behavior is assigned to the occurrence of trap-limited recombination.<sup>31–33</sup> TRMC measurements were also performed under various incident laser pulse intensities as shown in Fig. S8.† Fig. 4b shows the maximum TRMC signal ( $\varphi \sum \mu$ )<sub>max</sub> versus the incident laser pulse intensity at four different laser wavelengths. With increasing light intensities, the mobility ( $\varphi \sum \mu$ )<sub>max</sub> decreases gradually. This can be explained by the occurrence of non-geminate higher order electron–hole recombination. In addition, Fig. 4b shows that there is a clear dependency between the carrier mobility in  $\text{Cu}_5\text{V}_2\text{O}_{10}$  and the excitation wavelength. We tentatively attribute this to a wavelength-dependent photo-generation yield in  $\text{Cu}_5\text{V}_2\text{O}_{10}$ , similar to what was shown before for hematite.<sup>34,35</sup> Indeed, a d–d electronic transition is expected in  $\text{Cu}_5\text{V}_2\text{O}_{10}$ , which in the case of hematite has been suggested to not contribute to the generation of mobile carriers.

The effective lifetime of the charge carriers in  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films can be defined as the time that it takes for the signal to decrease to 50% of its starting value, and an average value of 30 ns for different excitation wavelengths is found for the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film. Based on the effective lifetime and the mobility, one might calculate an “effective” diffusion length of the photoexcited carriers. When doing this, we find a value of 1.3 nm. This value is clearly unrealistic. Even if we account for some surface roughness in our films (Fig. S2b and S3b†), the amount of light absorbed in the first 1.3 nm of the films would lead to negligible photocurrents. As mentioned above, trapping plays an important role in our films, and trapped carriers will have a much lower mobility than (quasi-) free carriers. With this in mind, we attribute the TRMC decay to a decay in mobility, as opposed to a decay in the number of carriers.

While slow carriers might still reach the interface and contribute to the photocurrent, it is reasonable to expect that the chance of recombination increases with the time that the carriers need to reach the interface. Nanostructuring strategies will, therefore, help to reduce the diffusion path lengths and may be needed to improve the charge carrier separation efficiency.

### Band positions

The conductivity type and flat band potential of our  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films were estimated by performing Mott–Schottky analysis. Fig. 5a shows a Mott–Schottky plot for the 250 nm  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode measured at different frequencies. The negative slopes in the Mott–Schottky curves indicate that the obtained  $\text{Cu}_5\text{V}_2\text{O}_{10}$  is a p-type semiconductor. In addition, an open-circuit potential (OCP) measurement under chopped AM1.5 illumination confirms the p-type conductivity, *i.e.*, the OCP shifts to a more positive value upon illumination as shown in Fig. S9.† It is worth noting that p-type conductivity for  $\text{Cu}_5\text{V}_2\text{O}_{10}$  is rarely reported in the literature; n-type  $\text{Cu}_5\text{V}_2\text{O}_{10}$  is more common.<sup>12,13</sup> The p-type conductivity may be caused by the presence of Cu vacancies in the photoelectrode. Cu vacancies have been shown to define the p-type nature of other Cu-based

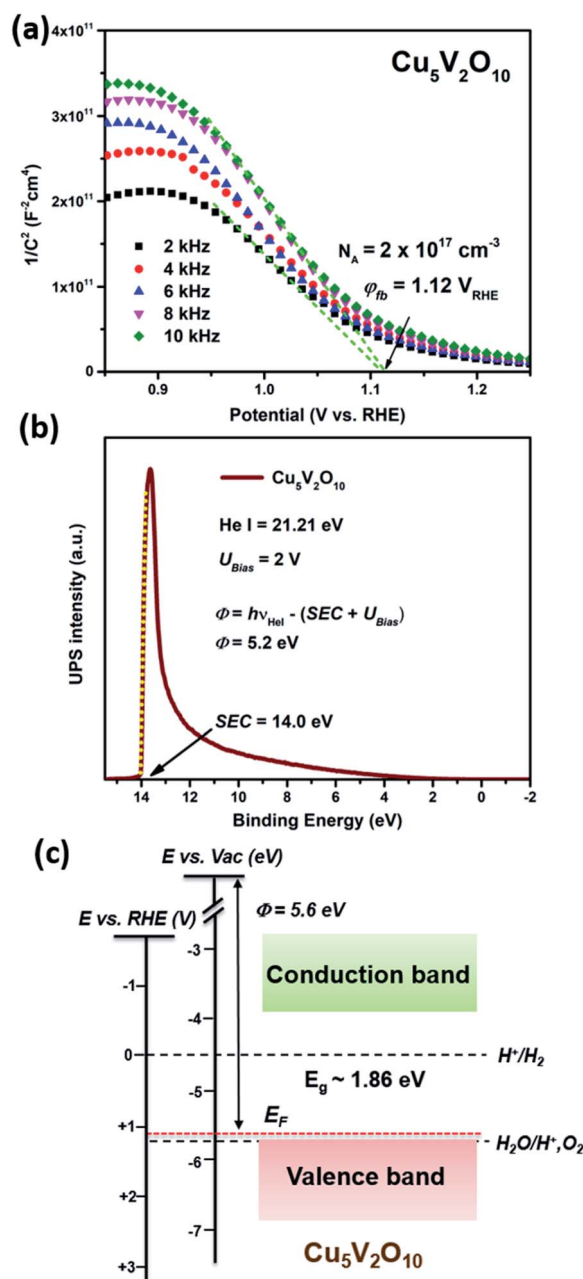


Fig. 5 (a) Mott–Schottky plot for a  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode in 0.3 M  $\text{K}_2\text{SO}_4$  and 0.2 M phosphate buffer (pH6.8) at 2 kHz, 4 kHz, 6 kHz, 8 kHz and 10 kHz. (b) UPS cutoff spectra measured with a 2 V bias for  $\sim 250$  nm  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film on FTO. (c) Schematic band diagram of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film constructed from UPS and Mott–Schottky measurements. The positions of the valence band, Fermi level  $E_F$  and the conduction band with respect to the vacuum level, the hydrogen electrode (RHE) potential, as well as water oxidation and reduction potentials are depicted.

photocathodes, such as  $\text{CuBi}_2\text{O}_4$  and  $\text{CuInS}_2$ .<sup>36,37</sup> Using eqn (6) in the ESI,† the flat band potential ( $\phi_{fb}$ ) and acceptor density ( $N_A$ ) can be obtained from the x-axis intercept and the slope of the Mott–Schottky plot, respectively. All Mott–Schottky curves at different frequencies in Fig. 5a intercept the same x-axis value, and the obtained  $\phi_{fb}$  value is  $1.12 \pm 0.02$  V vs. RHE, which is



consistent with the photocurrent onset potential ( $\sim 1.1$  V *vs.* RHE) of the film (*vide infra*). The frequency dispersion in the Mott–Schottky plot may be due to surface defects or a frequency-dependent dielectric constant; a detailed analysis of this is beyond the scope of this study and the reader is referred to the classic papers of Gomes and Cardon for an in-depth treatment on such phenomena.<sup>38,39</sup>

It should be noted that the Mott–Schottky curve flattens at the most negative potentials in Fig. 5a. We attribute this to the space charge region becoming large enough for the entire film to be depleted.<sup>40</sup> A fully depleted film will act as an insulator with a constant capacitance given by  $C = \epsilon_0 \epsilon_r A/d$ , where  $d$  is the thickness of the film and the other variables are described in the ESI.† Using the plateau values in Fig. 5a and the known value for the film thickness (250 nm), this allows us to calculate the relative permittivity (dielectric constant) of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  as  $\epsilon_r = 520 \pm 40$  in a frequency range from 4–10 kHz. Note that the  $\beta\text{-Cu}_2\text{V}_2\text{O}_7$  phase has a similarly high dielectric constant.<sup>26</sup> Using this value, the  $N_A$  of the film is estimated to be about  $(2.0 \pm 0.4) \times 10^{17} \text{ cm}^{-3}$ , which is relatively low but well within the range of values commonly observed for metal oxide semiconductors.<sup>30,41</sup> Since this low value results in a relatively wide space charge region even at moderate band bending, the space charge region will be insensitive to surface roughness effects and the films can be considered as ‘flat’ for the purpose of the Mott–Schottky analysis. Assuming a typical value of  $5 \times 10^{19} \text{ cm}^{-3}$  as the effective density of states at the valence band ( $N_V$ ), the Fermi level is expected to be located  $\sim 110$  mV above the valence band maximum (see ESI†).

Fig. 5b shows the secondary electron cut-off UPS spectrum of a 250 nm  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film measured with a 2 V bias. The work function (difference between the vacuum energy level and Fermi level) can be determined from the high binding energy onset of the secondary electron cut-off.<sup>42</sup> The work function is found to be  $21.21 - 2 - 14 = 5.21$  eV, given that the photon energy of the UV source (He I discharge) is 21.21 eV. Using 4.5 eV *vs.* vacuum as the reference value for the electrochemical reduction of water (0.0 V *vs.* RHE),<sup>1</sup> this places the work function obtained by UPS at 0.71 V *vs.* RHE. This is different by  $\sim 0.4$  V as compared to the  $\phi_{\text{fb}}$  value of 1.12 V *vs.* RHE obtained by the Mott–Schottky analysis. We attribute this to the different sample environment in the two measurements: Mott–Schottky measurement was performed in aqueous electrolyte, whereas UPS measurement was conducted on the surface of the samples in vacuum. As is well known, the work function of a material is extremely sensitive to adsorbate coverage. The drop in work function measured by UPS can also be attributed to the adsorption of carbonaceous species and other atmospheric contaminants on  $\text{Cu}_5\text{V}_2\text{O}_{10}$  surface, since it has been fully exposed to air before the measurement. The valence band maximum (VBM) position of 0.08 eV below  $E_F$  is determined by the linear extrapolation of the UPS spectrum at the low binding energy side to the binding energy axis (see Fig. S10†), which is in a good agreement from our previous estimation from Mott–Schottky analysis.

By combining the various parameters that were experimentally determined by UV-vis, Mott–Schottky, and UPS measurements we can now construct a detailed band diagram for

$\text{Cu}_5\text{V}_2\text{O}_{10}$  photoelectrode. Fig. 5c shows the band diagram of our spray deposited  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film with respect to the electrochemical redox potentials of the water oxidation ( $\text{H}^+$ ,  $\text{O}_2/\text{H}_2\text{O}$ ) and reduction ( $\text{H}^+/\text{H}_2$ ) reactions. It shows that the valence band (VB) of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  is approximately equal to the water oxidation potential. Considering that the  $E_g$  is 1.9 eV, we also estimate that the position of conduction band minimum ( $E_{\text{CBM}}$ ) is at about  $-0.62$  V *vs.* RHE or  $-3.88$  eV *vs.*  $E_{\text{vac}}$ , which is more negative than water reduction potential for hydrogen production. In other words, photo-excited electrons in the conduction band of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  thin films are thermodynamically able to reduce protons to hydrogen.

### Photoelectrochemical properties

To investigate the photoelectrochemical performance of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films, chopped linear sweep voltammetry (LSV) measurements were conducted under simulated AM1.5 illumination with the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode as the working electrode, Pt wire as the counter electrode, and Ag/AgCl as the reference electrode. The measurements were performed in 0.3 M  $\text{K}_2\text{SO}_4$  and 0.2 M phosphate buffer (pH 6.8) either with  $\text{H}_2\text{O}_2$  added as an electron scavenger to remove the reaction kinetics and surface recombination limitations, or with Ar bubbling (without  $\text{H}_2\text{O}_2$ ) to remove dissolved  $\text{O}_2$  and test for proton reduction.

Fig. 6a shows linear sweep voltammetry (LSV) scans for 250 nm  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode films measured with Ar bubbling. Photocurrent densities of  $0.02 \text{ mA cm}^{-2}$  and  $0.01 \text{ mA cm}^{-2}$  were obtained at 0.8 V *versus* RHE under back and front illumination, respectively. The current spikes are attributed to electron trapping on the photoelectrode's surface, followed by charge recombination.<sup>43–45</sup> Overall, moderate photocurrents are achieved. Interestingly, the photocurrent onset potential is relatively positive at  $\sim 1.1$  V *vs.* RHE. This value, *i.e.*, the difference between the thermodynamic reduction potential of water, 0 V *vs.* RHE, and the photocurrent onset potential, also serves as an indication of the photovoltage generated in the material. For comparison, other promising photocathodes such as CuO, Si, and  $\text{Cu}_2\text{ZnSnS}_4$  typically show a photocurrent onset of  $\sim 0.7$  V *vs.* RHE for water reduction,<sup>46–51</sup> although similarly high photocurrent onset potentials have recently been reported for  $\text{Cu}_2\text{O}$  and  $\text{CuBi}_2\text{O}_4$ .<sup>52,53</sup> Chopped LSV scans with  $\text{H}_2\text{O}_2$  were also performed as shown in Fig. 6b. Under optimum conditions (see Fig. S11† for the individual LSV scans for different  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film thicknesses), we could achieve a photocurrent density of  $0.52 \text{ mA cm}^{-2}$  and  $0.39 \text{ mA cm}^{-2}$  at 0.8 V *versus* RHE under back and front illumination, respectively. This is a factor of  $>10$  higher than without  $\text{H}_2\text{O}_2$ , indicating that the performance of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  is severely limited by surface reaction kinetics. Chopped LSV scans were also performed for the as deposited  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode (Fig. S12†); the light current density is barely distinguishable from the dark current density, which shows that the post-deposition annealing step that crystallizes the film is essential in order to get photoactivity.

To get an estimate for the amount of recombination in the films, we compare the measured photocurrents to the



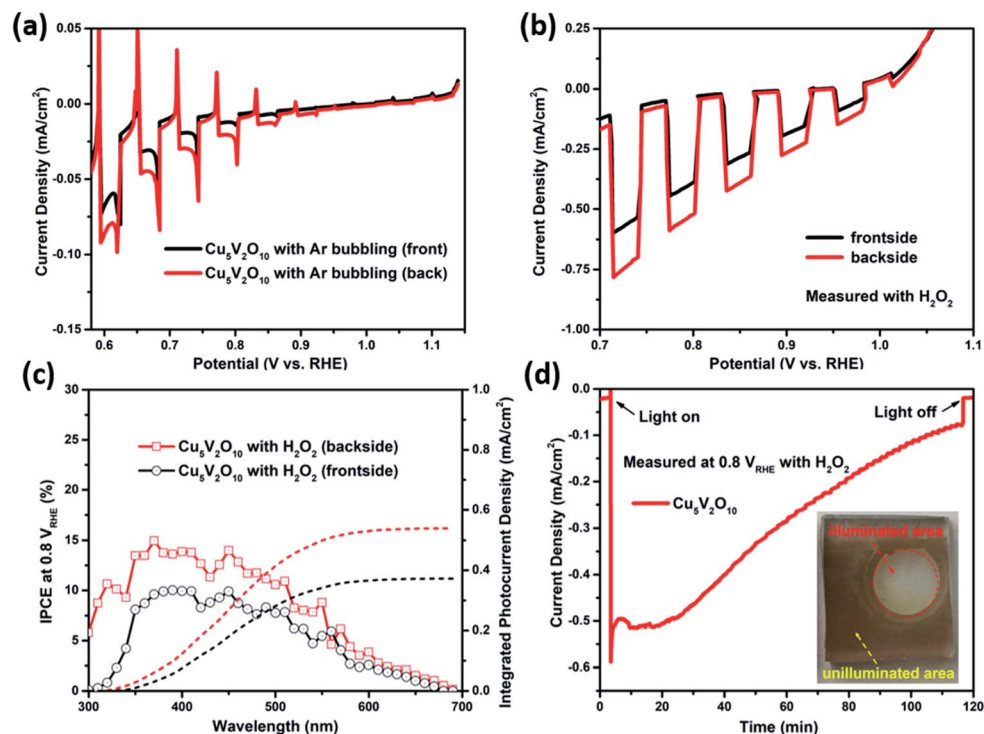


Fig. 6 Chopped LSV scans for  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathodes synthesized to a thickness of  $\sim 250$  nm on FTO measured with Ar bubbling (a) and with  $\text{H}_2\text{O}_2$  as an electron scavenger (b) under backside (red line) and frontside illumination (black line). (c) IPCE spectra of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  measured with  $\text{H}_2\text{O}_2$  at a potential of 0.8 V vs. RHE, corresponding integrated photocurrents are represented by the dashed lines. (d) Constant potential measurement at 0.8 V vs. RHE for 250 nm  $\text{Cu}_5\text{V}_2\text{O}_{10}$  films in the dark and light (AM1.5 irradiation). All measurements were done in 0.3 M  $\text{K}_2\text{SO}_4$  and 0.2 M phosphate buffer (pH 6.8).

photocurrent that can theoretically be generated in a 250 nm thick film (assuming that every absorbed photon yields a photoexcited electron-hole pair and that there is no recombination). The latter can be calculated from the absorption coefficient as a function of wavelength (Fig. 3b) and the AM1.5 solar spectrum. The results, shown in Fig. S13,† reveal that the maximum photocurrent for a 250 nm film is  $4.91 \text{ mA cm}^{-2}$ . The highest photocurrent observed for our films is  $0.52 \text{ mA cm}^{-2}$ , which means that  $\sim 89\%$  of the carriers recombine.

To gain additional information about the wavelength-dependence of the quantum efficiency of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathodes, we measured the incident photon-to-current conversion efficiency (IPCE) and calculated the absorbed photon-to-current efficiency (APCE) either with  $\text{H}_2\text{O}_2$  or with Ar bubbling. Fig. 6c shows the IPCE spectra of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film measured at 0.8 V versus RHE with  $\text{H}_2\text{O}_2$  under back and front illumination. The onset of the IPCE was 680 nm and it increases gradually toward shorter wavelengths. The  $\text{Cu}_5\text{V}_2\text{O}_{10}$  film produces IPCE values of  $<15\%$  for all wavelengths. The highest IPCE values of 14% and 10% were measured at around 400 nm under back and front illumination, respectively. The predicted AM1.5 photocurrent density ( $J_{\text{AM1.5}}$ ) of the photocathodes can be estimated by multiplying the IPCE values by the AM1.5 solar photon flux and subsequently integrating this for wavelengths below 700 nm according to eqn (5).† Integrating these spectra with the standard AM 1.5G solar spectrum yielded predicted photocurrent densities of  $0.54 \text{ mA cm}^{-2}$  and  $0.37 \text{ mA cm}^{-2}$  for  $\text{Cu}_5\text{V}_2\text{O}_{10}$

photocathode measured under back and front illumination, respectively. These values are consistent with the photocurrent densities of  $0.52$  and  $0.39 \text{ mA cm}^{-2}$  observed for the chopped LSV measurements at the same potential (Fig. 6b). IPCE spectra of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  were also measured without  $\text{H}_2\text{O}_2$  (Fig. S14 in the ESI†); in this case, the IPCE values remain lower than 1% for all wavelengths. Fig. S15† shows the APCE spectra for  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode measured with  $\text{H}_2\text{O}_2$  at 0.8 V vs. RHE under front and back illumination. The APCE shows the highest values in the region between 500–600 nm, which is consistent with the TRMC data in Fig. 4b. Overall, the low quantum efficiency is consistent with the very short effective carrier diffusion length, indicating that majority of the photoexcited carriers recombine before they even reach the interfacemodest AM1.5 photocurrents and the extensive trapping we observed from the TRMC measurements.

We note that the photoelectrochemical performance, both the AM1.5 photocurrent and the IPCE, of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  under back illumination is always larger than that under front illumination. Considering that our films are compact and that the absorption coefficient is high enough to have a non-constant light intensity profile across the film thickness, we can attribute this difference to the transport distance of electrons and holes in  $\text{Cu}_5\text{V}_2\text{O}_{10}$ . Under front illumination, holes have to traverse the entire thickness of the film, while electrons only need to travel a short distance. The situation is reversed under back illumination. Therefore, we can conclude that electron transport is more



efficient, and hole transport is the limiting factor in  $\text{Cu}_5\text{V}_2\text{O}_{10}$ . The poor hole transport properties may seem surprising in view of the fact that  $\text{Cu}_5\text{V}_2\text{O}_{10}$  is a p-type semiconductor, in which holes are the majority carriers. We attribute this to the fact that the film has a rather low acceptor density ( $\sim 2 \times 10^{17} \text{ cm}^{-3}$ ) and shows near-insulating behavior when fully depleted (*vide supra*). This also suggests that p-type doping might be a promising strategy to improve the performance of these film; this strategy has been successfully applied to a variety of n-type photoelectrodes, most notably  $\text{Fe}_2\text{O}_3$  and  $\text{BiVO}_4$ .<sup>54,55</sup>

In order to investigate the stability of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathodes, the photocurrent density was measured as a function of time at a fixed constant potential of 0.8 V vs. RHE under back illumination. The photocurrents show a fast decaying feature at the initial stage after illumination, which is often observed for unstable semiconductors (see Fig. S16†). The photocurrent density decreases by 85% within 10 min. This indicates extensive photo-corrosion, which is a common problem for Cu-based photocathodes.<sup>41,45,56</sup> On the other hand,  $\text{Cu}_5\text{V}_2\text{O}_{10}$  can transfer photo-generated electrons into the electrolyte much more efficiently when  $\text{H}_2\text{O}_2$  is present. The results are shown in Fig. 6d. The photocurrent density shows relatively good stability in the first 20 min and decreases by only 14%. After 20 min, however, it rapidly degrades to about 13% of its original value within 120 min and the illuminated area of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  electrode has completely disappeared (see the inset of Fig. 6d). We integrated the photocurrent density of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode over time to obtain the total charge that passed for the red curve of Fig. 6d. The obtained value,  $0.021 \text{ mmol cm}^{-2}$ , is over 7.6 times larger than the total charge needed to reduce all cations including  $\text{Cu}^{2+}$ ,  $\text{V}^{5+}$ , in the composite photocathode ( $0.0028 \text{ mmol cm}^{-2}$ ). Thus, about  $(7.6 - 1)/7.6 = 87\%$  of the photocurrent density went toward  $\text{H}_2\text{O}_2$  reduction, while 13% of the charge was presumably used to dissolve the film. The occurrence of photocorrosion even in the presence of an efficient electron scavenger indicates that p- $\text{Cu}_5\text{V}_2\text{O}_{10}$  is intrinsically unstable in aqueous solutions and that suitable protection layers are needed for this material. In contrast, the stability of n-type copper vanadate photoanodes, such as  $\alpha$ - $\text{CuV}_2\text{O}_6$ ,  $\beta$ - $\text{Cu}_2\text{V}_2\text{O}_7$ ,  $\gamma$ - $\text{Cu}_3\text{V}_2\text{O}_8$ , and  $\text{Cu}_{11}\text{V}_6\text{O}_{26}$  appears less problematic. Although protection layers may still be required for practical applications, several reports have demonstrated stabilities of several hours for n-type copper vanadates.<sup>12,26,57,58</sup> To compare the oxidation states changes in  $\text{Cu}_5\text{V}_2\text{O}_{10}$  electrode, XPS spectra of the sample after 1 hour of PEC measurements (*i.e.*, before the films were completely dissolved) were recorded (Fig. S17†). The Cu 2p peaks clearly shift toward lower binding energies after the PEC measurement, and fitting of the peaks show that the surface now contains a larger amount of  $\text{Cu}^+$  species ( $\text{Cu}^{2+} : \text{Cu}^+$  ratio of 1) (Fig. S17a†). This confirms that the  $\text{Cu}^{2+}$  at the surface of the  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode suffers cathodic photo-corrosion during PEC measurement in aqueous electrolyte, which is consistent with the fact that the electrochemical reduction potentials for  $\text{Cu}^{2+}$  are more positive than the reduction potential for  $\text{H}^+$ . This photoelectrochemical reduction of  $\text{Cu}^{2+}$  is clearly a relatively fast reaction, since it cannot be prevented even with the presence of an electron scavenger (*i.e.*,  $\text{H}_2\text{O}_2$ ) in our electrolyte. The XPS spectra for the O 1s and V 2p regions of PEC tested area are

shown in Fig. S17b.† A higher proportion of the adsorbed hydroxyl species after PEC measurement is evidenced from the O 1s peaks, as a result of the exposure to the aqueous electrolyte solutions. In the V 2p spectra, the peaks shifted to higher binding energy, suggesting that PEC measurement resulted in an increase in oxidation state of V to perhaps (partially) compensate the charge imbalance (*i.e.*, due to the reduction of copper). The simultaneous oxidation of the co-cation of Cu has also been observed for  $\text{CuBi}_2\text{O}_4$  photocathodes, in which the Bi  $4f_{5/2}$  and  $4f_{7/2}$  peaks shifted to higher binding energy after PEC experiments.<sup>30,59</sup>

## Conclusions

In this work we have identified the optoelectronic and PEC properties of a p-type  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathode synthesized by a scalable spray pyrolysis method, including bandgap, absorption coefficient, band positions, carrier mobility, flat band potential. Our study shows that the valence and conduction bands of  $\text{Cu}_5\text{V}_2\text{O}_{10}$  are nearly ideally positioned for PEC water splitting as they straddle the water oxidation and reduction potentials and allow for visible light absorption, which makes  $\text{Cu}_5\text{V}_2\text{O}_{10}$  a promising candidate to act as a top absorber layer in a dual absorber device for high-efficiency water reduction. However, the moderate absorption coefficient prevents it from absorbing a significant portion of the AM1.5 spectrum. Moreover, extensive trapping will likely prevent the photogenerated carriers from reaching the interface in these spray-deposited p-type films. Bare p-type  $\text{Cu}_5\text{V}_2\text{O}_{10}$  is also limited by its poor chemical stability, since we observed extensive photo-corrosion, even in the presence of  $\text{H}_2\text{O}_2$  as an electron scavenger. Further strategies such as the application of HER catalysts, protection layers, and interfacial layers can be used to improve the photoconversion efficiency of p-type  $\text{Cu}_5\text{V}_2\text{O}_{10}$  photocathodes. Nevertheless, given the severity of the carrier transport and stability limitations, it will be a major challenge to address both of these limitations at the same time. Efforts might be better spent in developing other complex metal oxide photocathodes, such as  $\text{CuBi}_2\text{O}_4$ ,  $\text{CaFe}_2\text{O}_4$ , and  $\text{CuFeO}_2$ .<sup>60,61</sup>

## Author contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

Angang Song thanks the China Scholarship Council for financial support (File no. 201607040078). We thank Christian Höhn for assistance with XPS and UPS measurements.





## References

- 1 R. van de Krol and M. Grätzel, *Photoelectrochemical hydrogen production*, Springer, 2012.
- 2 M. G. Walter, E. L. Warren, J. R. McKone, S. W. Boettcher, Q. Mi, E. A. Santori and N. S. Lewis, *Chem. Rev.*, 2010, **110**, 6446–6473.
- 3 T. Bak, J. Nowotny, M. Rekas and C. C. Sorrell, *Int. J. Hydrogen Energy*, 2002, **27**, 991–1022.
- 4 B. Alfaifi, H. Ullah, S. Alfaifi, A. Tahir and T. Mallick, Photoelectrochemical solar water splitting: from basic principles to advanced devices, *Veruscript Functional Nanomaterials*, 2018, **2**, BDJOC3.
- 5 J. Luo, L. Steier, M.-K. Son, M. Schreier, M. T. Mayer and M. Grätzel, *Nano Lett.*, 2016, **16**, 1848–1857.
- 6 A. Paracchino, V. Laporte, K. Sivula, M. Grätzel and E. Thimsen, *Nat. Mater.*, 2011, **10**, 456–461.
- 7 Y. J. Jang and J. S. Lee, *ChemSusChem*, 2019, **12**, 1835–1845.
- 8 U. A. Joshi, A. M. Palasyuk and P. A. Maggard, *J. Phys. Chem. C*, 2011, **115**, 13534–13539.
- 9 M. S. Prévot, N. Guijarro and K. Sivula, *ChemSusChem*, 2015, **8**, 1359–1367.
- 10 I. Sullivan, B. Zoellner and P. A. Maggard, *Chem. Mater.*, 2016, **28**, 5999–6016.
- 11 K. Rajeshwar, M. K. Hossain, R. T. Macaluso, C. Janáky, A. Varga and P. J. Kulesza, *J. Electrochem. Soc.*, 2018, **165**, H3192–H3206.
- 12 C.-M. Jiang, G. Segev, L. H. Hess, G. Liu, G. Zaborski, F. M. Toma, J. K. Cooper and I. D. Sharp, *ACS Appl. Mater. Interfaces*, 2018, **10**, 10627–10633.
- 13 P. Newhouse, D. Boyd, A. Shinde, D. Guevarra, L. Zhou, E. Soedarmadji, G. Li, J. B. Neaton and J. M. Gregoire, *J. Mater. Chem. A*, 2016, **4**, 7483–7494.
- 14 T. Kawada, S. Hinokuma and M. Machida, *Catal. Today*, 2015, **242**, 268–273.
- 15 T. Hillel and Y. Ein-Eli, *J. Power Sources*, 2013, **229**, 112–116.
- 16 J. Dai, M. Lai, R. M. LaFollette and D. Reisner, *ECS Trans.*, 2011, **33**, 3–9.
- 17 Y. Sakurai, H. Ohtsuka and J. i. Yamaki, *J. Electrochem. Soc.*, 1988, **135**, 32–36.
- 18 Z. He, C. Lin, W. Cheng, A. Okazawa, N. Kojima, J.-i. Yamaura and Y. Ueda, *J. Am. Chem. Soc.*, 2010, **133**, 1298–1300.
- 19 J. Ko, S. H. Kang, H.-W. Cheong, Y. S. Yoon, J. Ko, S. H. Kang, H.-W. Cheong and Y. S. Yoon, *J. Korean Ceram. Soc.*, 2019, **56**, 233–255.
- 20 I. Y. Ahmet, Y. Ma, J.-W. Jang, T. Henschel, B. Stannowski, T. Lopes, A. Vilanova, A. Mendes, F. F. Abdi and R. van de Krol, *Sustainable Energy Fuels*, 2019, **3**, 2366–2379.
- 21 M. Eguchi, A. Komamura, T. Miura and T. Kishi, *Electrochim. Acta*, 1996, **41**, 857–861.
- 22 R. Shannon and C. Calvo, *Acta Crystallogr., Sect. B: Struct. Crystallogr. Cryst. Chem.*, 1973, **29**, 1338–1345.
- 23 L. Denisova, Y. F. Kargin, N. Belousova, N. Galiakhmetova and V. Denisov, *Russ. J. Inorg. Chem.*, 2019, **64**, 725–728.
- 24 C.-M. Jiang, M. Farmand, C. H. Wu, Y.-S. Liu, J. Guo, W. S. Drisdell, J. K. Cooper and I. D. Sharp, *Chem. Mater.*, 2017, **29**, 3334–3345.
- 25 T. Kawada, T. Tajiri, H. Yamashita and M. Machida, *Catal. Sci. Technol.*, 2014, **4**, 780–785.
- 26 A. Song, A. Chemseddine, I. Y. Ahmet, P. Bogdanoff, D. Friedrich, F. F. Abdi, S. P. Berglund and R. van de Krol, *Chem. Mater.*, 2020, **32**, 2408–2419.
- 27 S. K. Pilli, T. E. Furtak, L. D. Brown, T. G. Deutsch, J. A. Turner and A. M. Herring, *Energy Environ. Sci.*, 2011, **4**, 5028–5034.
- 28 R. Sankari, M. Ehara, H. Nakatsuji, Y. Senba, K. Hosokawa, H. Yoshida, A. De Fanis, Y. Tamenori, S. Aksela and K. Ueda, *Chem. Phys. Lett.*, 2003, **380**, 647–653.
- 29 M. Zhang, Y. Ma, D. Friedrich, R. van de Krol, L. H. Wong and F. F. Abdi, *J. Mater. Chem. A*, 2018, **6**, 548–555.
- 30 S. P. Berglund, F. F. Abdi, P. Bogdanoff, A. Chemseddine, D. Friedrich and R. van de Krol, *Chem. Mater.*, 2016, **28**, 4231–4242.
- 31 J. E. Kroeze, T. J. Savenije and J. M. Warman, *J. Am. Chem. Soc.*, 2004, **126**, 7608–7618.
- 32 M. Kuno, D. P. Fromm, H. F. Hamann, A. Gallagher and D. J. Nesbitt, *J. Chem. Phys.*, 2000, **112**, 3117–3120.
- 33 A. Barzykin and M. Tachiya, *J. Phys. Chem. B*, 2002, **106**, 4356–4363.
- 34 A. Kay, M. Fiegenbaum-Raz, S. Müller, R. Eichberger, H. Dotan, R. van de Krol, F. F. Abdi, A. Rothschild, D. Friedrich and D. A. Grave, *Adv. Funct. Mater.*, 2019, 1901590.
- 35 D. Hayes, R. G. Hadt, J. D. Emery, A. A. Cordones, A. B. Martinson, M. L. Shelby, K. A. Fransted, P. D. Dahlberg, J. Hong and X. Zhang, *Energy Environ. Sci.*, 2016, **9**, 3754–3769.
- 36 F. M. Courtel, A. Hammami, R. Imbeault, G. Hersant, R. W. Paynter, B. t. Marsan and M. Morin, *Chem. Mater.*, 2010, **22**, 3752–3761.
- 37 F. Wang, W. Septina, A. Chemseddine, F. F. Abdi, D. Friedrich, P. Bogdanoff, R. van de Krol, S. D. Tilley and S. P. Berglund, *J. Am. Chem. Soc.*, 2017, **139**, 15094–15103.
- 38 F. Cardon and W. P. Gomes, *J. Phys. D: Appl. Phys.*, 1978, **11**, L63–L67.
- 39 W. P. Gomes and F. Cardon, *Prog. Surf. Sci.*, 1982, **12**, 155–215.
- 40 R. van de Krol, A. Goossens and J. Schoonman, *J. Electrochem. Soc.*, 1997, **144**, 1723.
- 41 A. Song, P. Plate, A. Chemseddine, F. Wang, F. F. Abdi, M. Wollgarten, R. van de Krol and S. P. Berglund, *J. Mater. Chem. A*, 2019, **7**, 9183–9194.
- 42 M. Helander, M. Greiner, Z. Wang and Z. Lu, *Appl. Surf. Sci.*, 2010, **256**, 2602–2605.
- 43 Y. Nakabayashi, M. Nishikawa and Y. Nosaka, *Electrochim. Acta*, 2014, **125**, 191–198.
- 44 Y.-H. Choi, K. D. Yang, D.-H. Kim, K. T. Nam and S.-H. Hong, *Mater. Lett.*, 2017, **188**, 192–196.
- 45 W. Septina, R. R. Prabhakar, R. Wick, T. Moehl and S. D. Tilley, *Chem. Mater.*, 2017, **29**, 1735–1743.



- 46 M. G. Kast, L. J. Enman, N. J. Gurnon, A. Nadarajah and S. W. Boettcher, *ACS Appl. Mater. Interfaces*, 2014, **6**, 22830–22837.
- 47 D. Kang, T. W. Kim, S. R. Kubota, A. C. Cardiel, H. G. Cha and K.-S. Choi, *Chem. Rev.*, 2015, **115**, 12839–12887.
- 48 A. A. Dubale, W.-N. Su, A. G. Tamirat, C.-J. Pan, B. A. Aragaw, H.-M. Chen, C.-H. Chen and B.-J. Hwang, *J. Mater. Chem. A*, 2014, **2**, 18383–18397.
- 49 C. G. Morales-Guio, S. D. Tilley, H. Vrubel, M. Grätzel and X. Hu, *Nat. Commun.*, 2014, **5**, 3059.
- 50 H. Kumagai, T. Minegishi, Y. Moriya, J. Kubota and K. Domen, *J. Phys. Chem. C*, 2014, **118**, 16386–16392.
- 51 G. Ma, T. Minegishi, D. Yokoyama, J. Kubota and K. Domen, *Chem. Phys. Lett.*, 2011, **501**, 619–622.
- 52 L. Pan, Y. Liu, L. Yao, R. Dan, K. Sivula, M. Grätzel and A. Hagfeldt, *Nat. Commun.*, 2020, **11**, 318.
- 53 R. Gottesman, A. Song, I. Levine, M. Krause, A. T. M. N. Islam, D. Abou-Ras, T. Dittrich, R. van de Krol and A. Chemseddine, *Adv. Funct. Mater.*, 2020, 1910832.
- 54 I. Cesar, A. Kay, J. A. Gonzalez Martinez and M. Grätzel, *J. Am. Chem. Soc.*, 2006, **128**, 4582–4583.
- 55 Y. Liang, T. Tsubota, L. P. A. Mooij and R. van de Krol, *J. Phys. Chem. C*, 2011, **115**, 17594–17598.
- 56 A. Song, P. Bogdanoff, A. Esau, I. Y. Ahmet, I. Levine, T. Dittrich, T. Unold, R. van de Krol and S. P. Berglund, *ACS Appl. Mater. Interfaces*, 2020, **12**, 13959–13970.
- 57 L. Zhou, Q. Yan, J. Yu, R. J. R. Jones, N. Becerra-Stasiewicz, S. K. Suram, A. Shinde, D. Guevarra, J. B. Neaton, K. A. Persson and J. M. Gregoire, *Phys. Chem. Chem. Phys.*, 2016, **18**, 9349–9352.
- 58 M. A. Lumley and K.-S. Choi, *Chem. Mater.*, 2017, **29**, 9472–9479.
- 59 D. Kang, J. C. Hill, Y. Park and K.-S. Choi, *Chem. Mater.*, 2016, **28**, 4331–4340.
- 60 M. S. Prévot, X. A. Jeanbourquin, W. S. Bourée, F. Abdi, D. Friedrich, R. van de Krol, N. Guijarro, F. Le Formal and K. Sivula, *Chem. Mater.*, 2017, **29**, 4952–4962.
- 61 D. K. Lee, D. Lee, M. A. Lumley and K.-S. Choi, *Chem. Soc. Rev.*, 2019, **48**, 2126–2157.

