

Arc Synthesis, Crystal Structure, and Photoelectrochemistry of Copper(I) Tungstate

Miguel Tayar Galante,[○] Aleksandar Živković,[○] Jéssica Costa Alvim, Cinthia Cristina Calchi Kleiner, Márcio Sangali, S. F. Rebecca Taylor, Adam J. Greer, Christopher Hardacre, Krishnan Rajeshwar, Rubens Caram, Rodnei Bertazzoli, Robin T. Macaluso, Nora H. de Leeuw,^{*} and Claudia Longo^{*}



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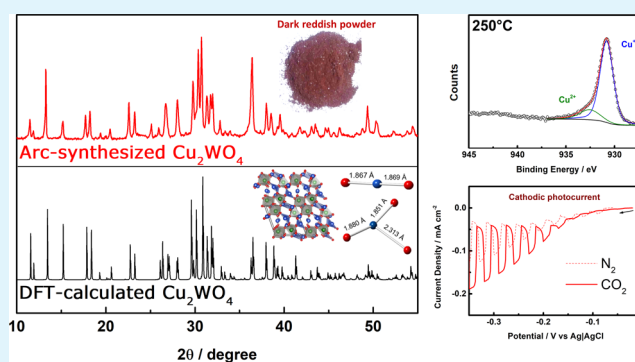
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ABSTRACT: A little-studied p-type ternary oxide semiconductor, copper(I) tungstate (Cu_2WO_4), was assessed by a combined theoretical/experimental approach. A detailed computational study was performed to solve the long-standing debate on the space group of Cu_2WO_4 , which was determined to be triclinic $P\bar{1}$. Cu_2WO_4 was synthesized by a time-efficient, arc-melting method, and the crystalline reddish particulate product showed broad-band absorption in the UV–visible spectral region, thermal stability up to $\sim 260^\circ\text{C}$, and cathodic photoelectrochemical activity. Controlled thermal oxidation of copper from the Cu(I) to Cu(II) oxidation state showed that the crystal lattice could accommodate Cu^{2+} cations up to $\sim 260^\circ\text{C}$, beyond which the compound was converted to CuO and CuWO_4 . This process was monitored by powder X-ray diffraction and X-ray photoelectron spectroscopy. The electronic band structure of Cu_2WO_4 was contrasted with that of the Cu(II) counterpart, CuWO_4 using spin-polarized density functional theory (DFT). Finally, the compound Cu_2WO_4 was determined to have a high-lying (negative potential) conduction band edge underlining its promise for driving energetic photoredox reactions.

KEYWORDS: ternary copper oxide, copper tungstate, arc synthesis, p-type semiconductor, photoelectrochemistry, solar fuels, crystal structure, density functional theory



1. INTRODUCTION

This paper presents computational aspects, solid-state chemistry, and photoelectrochemical (PEC) properties of copper(I) tungstate, Cu_2WO_4 , a little-studied inorganic compound of both fundamental and technological import. It belongs to the family of multinary copper oxides that have interesting electronic, magnetic, and optical properties for a range of applications, including in sensors/biosensors,¹ electronic devices,² and photoelectrochemical (PEC) solar energy conversion.^{3,4} Ternary copper-based inorganic p-type semiconductors have been studied extensively for their applicability in solar hydrogen generation.^{5–7} In these compounds, copper commonly occurs in the +1 or +2 oxidation states, in turn dictating their electronic properties, i.e., copper(II)-based oxides can be found both as n-type⁸ or p-type^{9,10} semiconductors, while copper(I)-based compounds are p-type semiconductors, as exemplified by the parent oxide, Cu_2O .¹¹

Much of the literature on ternary oxides based on copper oxide and tungsten trioxide as the binary components is related to copper(II) tungstate, CuWO_4 . This compound is an n-type semiconductor¹² with an optical band gap of ~ 2.2 eV. It has

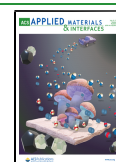
been considered as a photoanode for water oxidation^{12–14} and as a photocatalyst for hydrogen evolution in aqueous suspension.¹⁵ In contrast, little research has been reported on copper(I)-based tungstate, Cu_2WO_4 , which is a p-type semiconductor.

In contrast to the copper(II) tungstate counterpart, literature reports are sparse on the copper(I) compound (Cu_2WO_4), of focus in this study. Two studies^{16,17} on its crystal structure appeared in the 1980s followed by a rather long hiatus. Further, these two structural reports disagreed on the space group, $P\bar{1}$ or $P\bar{1}$. Marinder¹⁶ was the first to investigate the structure of Cu_2WO_4 and used X-ray diffraction to propose the $P\bar{1}$ space group, but failed to rigorously refine the structural model via Rietveld refinement. On the other

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hand, Mumm¹⁷ used single-crystal X-ray diffraction to assign the space group $P\bar{1}$, ruling out $P\bar{1}$ on the basis that the $d(W-O) = 1.42 \text{ \AA}$ and $d(\text{Cu}-\text{Cu}) = 2.42/2.51 \text{ \AA}$ distances were too short. Motivated by this historical ambiguity in the space group determination of the Cu_2WO_4 structure, a major objective of the present study was to computationally predict the thermodynamic and spectroscopic properties of each structure candidate, and thereby unequivocally assign the space group to which Cu_2WO_4 belonged.

Time-efficient synthesis of the little-studied Cu_2WO_4 and its PEC characterization constituted the other primary objectives of this study. Considering that computer-aided materials genome for a targeted application is still an evolving “blind art”,¹⁸ iterative time-efficient synthesis approaches are needed in the interim timeframe to allow for fast and efficient screening of possible candidates. Therefore, in this paper, a single-step, arc synthesis of Cu_2WO_4 , using Cu_2O and WO_3 as precursors, is introduced. The commonly employed solution (e.g., sol-gel synthesis) or even, ceramic (solid-state) synthesis routes require hours to days to derive the targeted product, most often in an amorphous state. Also, a postsynthesis thermal annealing step is then needed in these cases to improve the sample morphology. In contrast, in the arc synthesis, the product can be obtained in the crystalline state within minutes.

2. EXPERIMENTAL AND THEORETICAL METHODS

2.1. Computational Aspects. Spin-polarized density functional theory (DFT) calculations were performed using the Vienna Ab initio Simulation Package (VASP)^{19–21} with the projector-augmented wave (PAW) method and a plane-wave cutoff of 500 eV. For the PAW potentials, the valence electronic configurations used were $4s^13d^{10}$ for Cu, $2s^22p^4$ for O, and $6s^15d^55p^65s^2$ for W.

In the DFT calculations, the general gradient approximation (GGA) for the exchange–correlation (XC) functional was employed within the Perdew–Burke–Ernzerhof (PBE) parametrization.²² For the hybrid DFT calculations, the Heyd–Scuseria–Ernzerhof (HSE)^{23–25} XC functional was used, with an inverse screening length of $\mu = 0.2 \text{ \AA}^{-1}$ and 15% ($\alpha = 0.15$) of the nonlocal exact Hartree–Fock (HF) exchange. The method of Gajdoš was used to obtain the real and imaginary parts of the dielectric function,²⁶ from which the absorption coefficient was derived. Long-distance dispersion corrections were included using the D3 approach of Grimme.²⁷ The conjugate gradient method was used for structural optimization, with the total energy and force convergence criteria set to 10^{-5} eV and 0.01 eV \AA^{-1} , respectively. The Brillouin zone was sampled by $2 \times 2 \times 4$ and $5 \times 5 \times 5$ Γ -centered Monkhorst–Pack meshes.²⁸ The extent of charge distribution was studied using the Bader scheme as implemented in the Henkelman code.^{29–31} Graphical drawings were produced using VESTA,³² the electron and hole effective masses were calculated using Sumo.³³ The transition dipole moment (TDM) was obtained using the VASPKIT program.³⁴

Vibrational frequency calculations were performed using the all-electron code CRYSTAL (2017 release)^{35,36} with the hybrid B3LYP exchange–correlation functional,^{37,38} in conjunction with triple- ζ -valence + polarization Gaussian-type basis sets employed to treat the Cu and O atoms, while a scalar relativistic pseudopotential was adopted to treat the core electrons of W. The optimal basis sets of Cu and O were taken without further modification from Linnera,^{39,40} while for W, the effective core pseudopotential derived by Hay and Wadt was chosen.^{41,42} This protocol has been shown to reproduce the structure as well as the vibrational properties of ionic and semi-ionic compounds in good agreement with experimental data.^{43,44}

In CRYSTAL, the convergence of the real-space summation of the Coulomb and exchange contributions to the Hamiltonian matrix is controlled by five overlap criteria. The values used in this study were 10^{-6} , 10^{-6} , 10^{-6} , 10^{-6} , and 10^{-12} . The threshold on the self-consistent

(SCF) energy was set to 10^{-7} Ha. For the compounds of interest, the convergence with respect to k -points was checked. Monkhorst–Pack meshes of $3 \times 3 \times 7$ for Cu_2WO_4 and $7 \times 7 \times 7$ for CuWO_4 were used to sample the first Brillouin zone.³⁵ Relative infrared and Raman intensities were computed analytically, based on coupled-perturbed Hartree–Fock/Kohn–Sham (CPHF/KS) treatments implemented in the code.^{45–48} To confirm the space group of the crystal, the FINDSYM suite was used on top of the structure obtained with fully relaxed DFT geometries.⁴⁹

2.2. Arc Synthesis of Cu_2WO_4 . Cu_2O (425 mg) and WO_3 (600 mg) (2.3 Cu:1 W atom molar ratio) were mixed and ground in an agate mortar until a homogeneous, finely dispersed powder was obtained. The mixture was made into a pellet using a hydraulic press and then placed inside a water-cooled copper crucible housed in an arc-melting furnace. The furnace was sealed, evacuated, and then flushed with argon to create an inert, deoxygenated atmosphere. The pellet was then melted by applying an electric arc for 30 s. The power source was set to provide an arc current of 90 A. After cooling, the molten material was ground in an agate mortar. Less than ~2% mass loss was observed.

Powder X-ray diffraction (XRD) data were collected at room temperature on a PANalytical Empyrean diffractometer equipped with Cu $K\alpha$ radiation, a Bragg–Brentano HD optical module, and an X'Celerator detector. Data were collected at $10 \leq 2\theta \leq 70^\circ$ with a step size of $2\theta = 0.008^\circ$ and a step rate of 0.2 s per step. Ultraviolet–visible diffuse reflectance measurements were performed on a Shimadzu (UV-2460) spectrometer, equipped with a diffuse reflection accessory for solids analysis. Differential scanning calorimetry (DSC) analysis was performed in DSC 2910 device (TA Instruments). Thermogravimetric analysis (TGA) was performed on a TA Instruments Model 2950 instrument. Both sets of thermal analyses were performed in air, at a heating ratio of $10 \text{ }^\circ\text{C min}^{-1}$.

X-ray photoelectron spectroscopy (XPS) was performed on a Kratos Axis Ultra DLD instrument using an Al- $K\alpha$ monochromatic source and hemispherical electron energy analyzer. Peak profile fitting was done using XPSPeak 4.1 software considering a Shirley-type baseline. Binding energy values were corrected using the peak from adventitious carbon (C 1s binding energy = 284.8 eV) as reference.

2.3. Photoelectrochemical Measurements. Thin-film electrodes were prepared by suspending the arc-synthesized Cu_2WO_4 powder sample in isopropanol, followed by sonication for 30 min. The suspension was cast dropwise over the conductive surface of previously washed fluorine-doped tin oxide (FTO)-glass slides and dried at room temperature in flowing air before thermal annealing.

Electrochemical measurements were performed in a single-compartment 3-electrode cell using 0.5 mol L^{-1} NaHCO_3 aqueous solution as supporting electrolyte. Platinum wire was used as the counterelectrode and Ag/AgCl/KCl 3 M as the reference electrode. An Autolab PGSTAT128N potentiostat was used for the electrochemical and photoelectrochemical (PEC) measurements. An Asahi Spectra HAL-100 solar simulator calibrated at 1 Sun, AM 1.5G was used as the light source.

3. RESULTS AND DISCUSSION

3.1. $P\bar{1}$ or $P1$? The two calculated crystal structures are depicted in Figure 1 for Cu_2WO_4 corresponding to the two space groups in question. Starting from the respective experimental atomic positions and lattice parameters, a full geometrical relaxation was undertaken, with the final structures in the $P\bar{1}$ and $P1$ candidates exhibiting essentially similar features. The W atom was always coordinated by six O atoms in an octahedral environment regardless of the space group assignment; however, there were differences in the Cu coordination.

Cu atoms could be found linearly coordinated by two O atoms, with distances of 1.82–1.88 and 1.84–1.98 \AA for $P\bar{1}$ and $P1$, respectively. However, the Cu2 atom in the $P\bar{1}$ model was coordinated to four O atoms in a distorted tetrahedron,

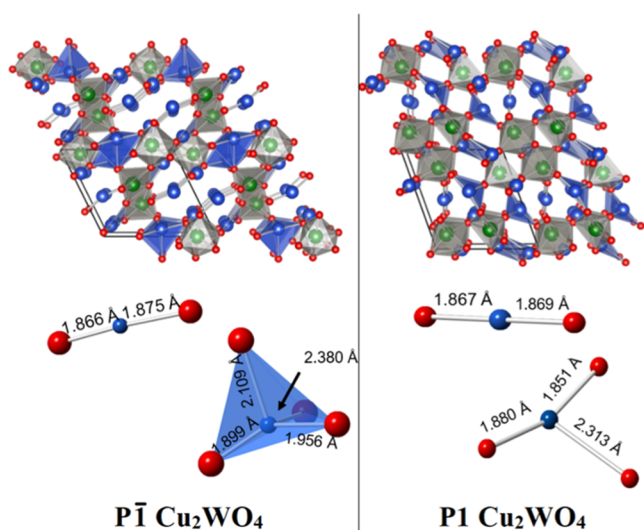


Figure 1. Crystal structures of fully relaxed copper(I) tungstate geometries: a detailed view of $P\bar{1}$ Cu_2WO_4 (left) and $P1$ Cu_2WO_4 (right). The Cu atoms are in distorted tetrahedral and linear geometries for $P\bar{1}$ and distorted T-shaped and linear geometries for $P1$. Cu, W, and O atoms are shown as blue, green, and red spheres, respectively.

with Cu–O distances ranging between 1.89 and 1.95 Å. Two additional O atoms lie 2.11 and 2.38 Å away. In the $P1$ model, half of the Cu atoms were coordinated to three oxygen atoms in a trigonal geometry. The 3-fold coordination (distorted T-shaped geometry) is expected for Cu^+ species rather than the unusual distorted tetrahedron in the $P\bar{1}$ structure, which is expected for Cu^{2+} species. The Cu–O distances in the $P1$ model were 1.86–1.89 Å with an additional O atom separation of 2.32 Å.

Furthermore, the bond lengths from the experimentally available data of the $P\bar{1}$ structure, namely, $d(\text{W–O}) = 1.42$ Å and $d(\text{Cu–Cu}) = 2.42/2.51$ Å¹⁶ were not found in the computed ground state structure. Our optimized bond lengths: $d(\text{W–O}) = 1.74$ Å and $d(\text{Cu–Cu}) = 2.71/3.06$ Å for Cu_2WO_4 ($P\bar{1}$), were comparable to the values of $d(\text{W–O}) = 1.78$ Å and $d(\text{Cu–Cu}) = 2.62/2.78$ Å in Cu_2WO_4 ($P1$). Based on the structural data outlined above, we suggest that Cu_2WO_4 is most likely to crystallize in $P1$ as proposed by Mumm,¹⁷ while the ($P\bar{1}$) structure of Marinder¹⁶ undergoes strong relaxation into an arrangement very close to the $P1$ structure.

The computed cell parameters for copper(I) tungstate in $P\bar{1}$ and $P1$ are listed in Tables 1 and 2, respectively. Three different functionals—PBE, HSE, and B3LYP, yielded unit cell parameters equivalent to one another for both $P\bar{1}$ and $P1$. However, the XC functionals yielded cell parameters that deviated by 8–9% for $P\bar{1}$, substantially higher than the 1–2% deviation for $P1$.

Figure 2 maps the computed variation of the total energy with respect to the unit cell volume, which also indicated that

the $P1$ space group was thermodynamically favored over the $P\bar{1}$ space group. This energy computation corroborates the fact that arc-melted samples (see below) typically yield thermodynamically stable phases.

To further clarify the differences between the two possible space groups, vibrational calculations were performed on both the unrelaxed CIF structures available from the American Mineralogist Database, as well as the fully DFT optimized structures. The calculated spectra of Cu_2WO_4 in the two considered space groups are shown in Figure S1. The spectra obtained from the CIF structure without further relaxation contain several negative frequencies (21 in the case of $P\bar{1}$ and 3 in the case of $P1$), indicating that the structures belonged to transition states or were unstable. Upon full geometry optimization, two frequencies in the $P\bar{1}$ structure remained negative, while all frequencies in the $P1$ structure turned positive. This further illustrates that the $P\bar{1}$ structure was probably not fully refined and did not correspond to the most likely structure of Cu_2WO_4 . However, it is also worth noting that a calculation of all positive frequencies does not ensure that the structure is at the “global” minimum, but only at a “minimum”, which could still be a local one.^{50,51} The fully relaxed geometries of both discussed structures are available in the Supporting Information.

Raman spectra for both the $P1$ and $P\bar{1}$ structures were also calculated and compared with the experimental Raman spectrum of Cu_2WO_4 (obtained from an arc-melted Cu_2WO_4 sample, to be discussed in the following section) (Figure 3). The experimental Raman spectrum (albeit, significantly broadened) clearly shows better conformity with the calculated spectrum for the $P1$ structure.

3.2. Arc Synthesis, Structure, and Optical Behavior of Cu_2WO_4 . A representative XRD pattern for the powder sample is shown in Figure 4a. All peaks matched with the calculated $P1$ structure. Importantly, no peaks associated with Cu_2O and WO_3 precursors were observed, indicating total consumption of the starting materials. To ensure that the obtained structure was not a kinetically trapped metastable phase rather than the thermodynamically stable one determined by DFT, XRD analyses were performed before and after annealing at 250 °C. Figure S2 shows these results in lower angles, where $P1$ and P structures most differ from each other. No changes are observed, confirming that the obtained material was indeed a thermodynamically stable $P1$ structure. It should also be noted that a small excess of Cu_2O was used in the precursor mixture (see Section 2). Arc-melting a stoichiometric 2 Cu:1 W atom ratio resulted in a complex mixture (Figure S3). This may be due to the volatility of metal oxides.

Strong light absorption was observed over the entire UV–vis region, as seen in the diffuse reflectance spectrum in Figure 4b. A Tauc plot⁵² assuming a direct optical transition is presented, where a main optical transition was observed with an absorption onset at approximately 1.97 eV, together with an absorption “tail”⁵³ extending to the near-infrared region (see

Table 1. Experimental and Theoretical Unit Cells of Cu_2WO_4 in $P\bar{1}$

	<i>a</i> (Å)	<i>b</i> (Å)	<i>c</i> (Å)	α	β	γ	<i>V</i> (Å ³)
Marinder ¹⁶	5.8578	7.9140	8.2050	110.360	96.450	93.970	351.948
PBE	5.7868	7.7199	8.8347	64.830	88.554	88.470	357.043
HSE	5.7130	7.6947	8.8396	65.220	89.363	88.840	352.741
B3LYP	5.7322	7.3662	8.8970	66.6446	89.9870	88.6059	344.780

Table 2. Experimental and Theoretical Unit Cells of Cu_2WO_4 in $P1$

	a (Å)	b (Å)	c (Å)	α	β	γ	V (Å ³)
Mumtaz ¹⁷	7.9018	8.2008	5.8718	96.506	94.033	110.270	352.176
PBE	7.8804	8.2054	5.9174	95.941	93.421	109.503	356.928
HSE	7.9182	8.1943	5.8830	96.071	93.436	109.618	355.678
B3LYP	7.8398	8.2148	5.7540	97.725	95.585	109.929	341.125

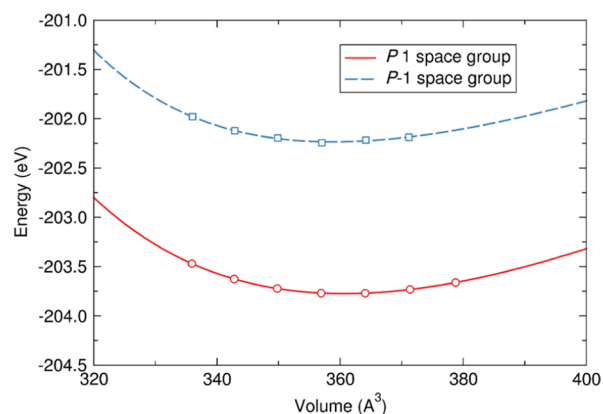


Figure 2. Calculated energy versus unit cell volume for copper(I) tungstate crystallizing in two distinct space groups. The values were fitted with a Birch–Murnaghan equation of state.

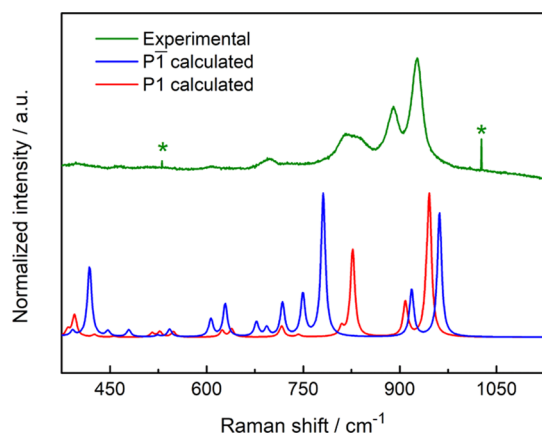


Figure 3. Calculated Raman spectra of Cu_2WO_4 for $P1$ and $P\bar{1}$ space groups, and experimental data obtained from an arc-melted Cu_2WO_4 sample. Asterisks on the experimental spectrum indicate equipment artifacts.

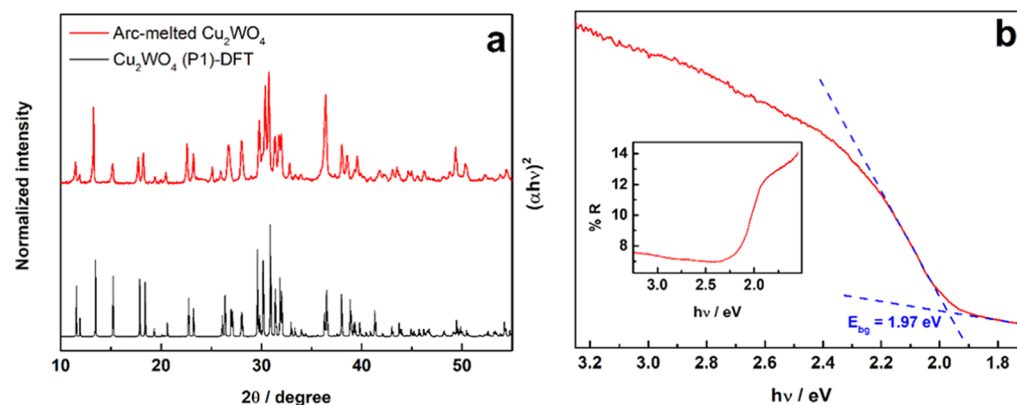


Figure 4. Powder X-ray diffractogram of arc-synthesized Cu_2WO_4 , with a calculated pattern of $P1$ Cu_2WO_4 (a) and a Tauc plot from a diffuse reflectance spectrum assuming direct transition (b) for arc-melted Cu_2WO_4 . The inset in (b) shows the raw reflectance data.

inset showing the raw reflectance data). The optical behavior may be compared against the electronic band structure of the compound which is presented next.

3.3. Electronic Band Structure of Cu_2WO_4 and Comparison with CuWO_4 .

The DFT Kohn–Sham electronic band structure of Cu_2WO_4 , together with the accompanying densities of states, are shown in Figure 5a. For comparison, the same quantities were calculated for CuWO_4 using an equivalent methodology (Figure 5b). The nonmagnetic ground state of Cu_2WO_4 featured a direct band gap between the valence band maximum (VBM) and the conduction band minimum (CBM) of 1.98 eV located at the high-symmetry X-point (Table 3). The top of the valence band consisted of mostly Cu-3d states weakly mixed with O-2p states, while the conduction band was composed predominantly of W-5d states strongly hybridizing with empty O-2p states. Such behavior stood in complete contrast to CuWO_4 , where O-2p states dictate the VBM and mixed strongly with Cu-3d states. The conduction band in this case, which can be viewed as an intermediate band, was predominantly made of empty Cu-3d e_g states and further separated by around 1 eV from the subsequent conduction band states. These results indicate the fundamentally distinct nature of the two different oxide compounds.

Upon electron excitation (for instance, via incoming electromagnetic radiation), the foremost transition in Cu_2WO_4 is a d-d transfer between the Cu and W states, while in CuWO_4 , electrons hop between the p levels of O and the d levels of Cu (visual representation through partial charges shown in Figure S4). As such, CuWO_4 can be identified as a charge-transfer (p-d)-type insulator.⁵² Furthermore, contrary to earlier suppositions,⁵³ all of the aforementioned transitions were found to be dipole-allowed in both compounds. Bader charges calculated for each structure are shown in Table S1.

The orientation-dependent hole and electron effective masses were estimated from the calculated HSE band

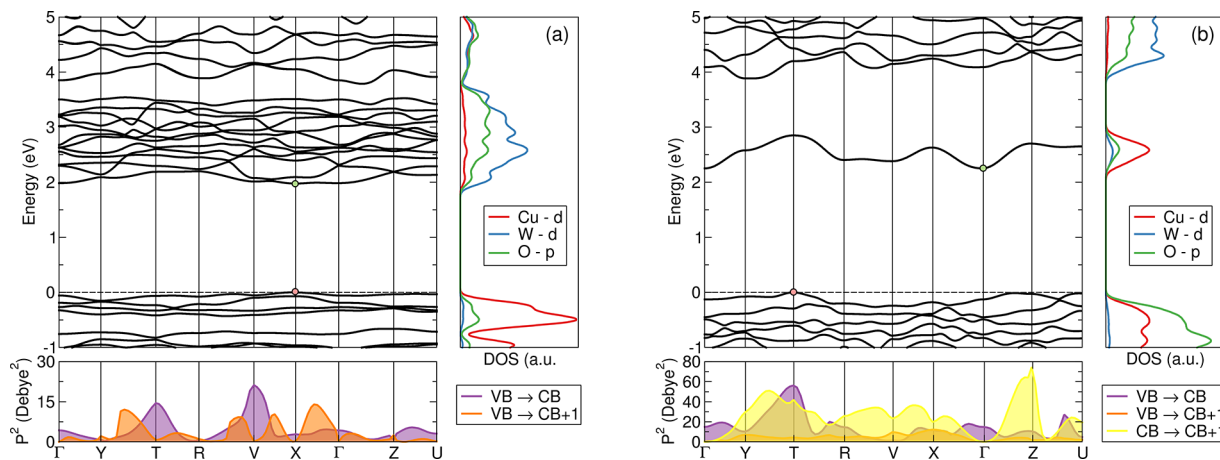


Figure 5. Electronic band structures, density of states, and transition dipole moments for Cu₂WO₄ (a) and CuWO₄ (b).

Table 3. Computed Kohn–Sham DFT Electronic Band Gap Values

	Cu ₂ WO ₄		CuWO ₄	
	indirect gap	direct gap	indirect gap	direct gap
PBE	1.19 eV (X–V)	1.24 eV (X)	0.26 eV (T–X)	0.56 eV (Γ)
HSE		1.98 eV (X)	2.25 eV (T–X)	2.40 eV (Γ)
exp.	~1.97 eV (this work)		~2.2 eV (ref 12)	

structure, using a parabolic fit to the band edges, and these are listed in Table 4. The obtained effective mass values were

Table 4. Calculated Hole and Electron Effective Masses for the Two Copper Tungstate Compounds under Scrutiny, from the HSE Band Structure

	effective mass (<i>m</i> ₀)			
	valence band		conduction band	
Cu ₂ WO ₄	X → V	−5.524	X → V	2.942
	X → Γ	−5.884	X → Γ	2.942
CuWO ₄	T → Y	−1.830	Γ → X	2.734
	T → R	−1.049	Γ → Y	1.440
			Γ → Z	1.247

relatively high, reflecting the flatness of the bands as well as the significant contributions of the d states of Cu and W to the top and bottom of the valence and conduction bands, respectively. Both effective masses further showed strong isotropy, indicating that there was no preferred direction within the crystal for the conduction of either electrons or holes.

3.4. Thermal Stability of Cu₂WO₄ and Decomposition to CuWO₄. Heating the obtained material up to ~260 °C did not change the XRD profile in Figure 4a, indicating that the crystal structure was stable up to ~260 °C. The thermal stability was further evaluated by differential scanning calorimetry/thermogravimetric analyses (DSC/TGA, Figure S5) with samples heated in an oxidative atmosphere. A set of observed exothermic events were associated with significant mass gain, which was attributed to oxygen incorporation into the compound structure.

To investigate this further, the Cu₂WO₄ films were preannealed at different temperatures in the range, 200–350 °C and analyzed by XRD and XPS. XRD analyses at different annealing temperatures (Figure 6) showed structural stability up to 260 °C. Further heating promoted thermal conversion to

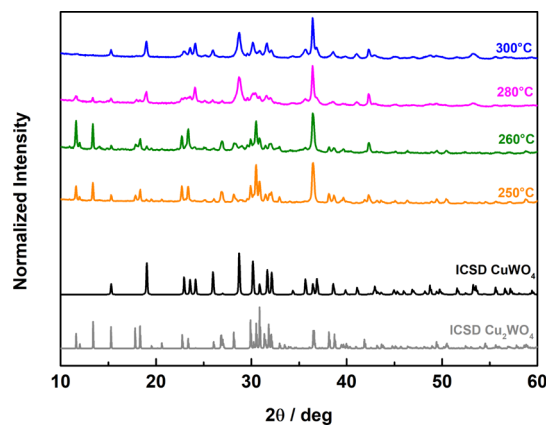


Figure 6. XRD patterns of Cu₂WO₄ films on glass, preannealed at different temperatures in air for 30 min.

CuWO₄, which belongs to the same crystal structure as Cu₂WO₄ (triclinic). ICSD references of both phases are shown together with the diffractograms of the annealed samples in Figure 6. A closer inspection of XRD data indicated that after thermal oxidation, a peak at 35.68 °C was seen (Figure S6), which is characteristic of the Cu(II) binary oxide, CuO. In this case, based also on mass gain as observed in TGA data (Figure S5), we conclude that thermal oxidation of Cu₂WO₄ occurred via oxidation of Cu⁺ to Cu²⁺, followed by expulsion of CuO (eq 1). Similar oxidation routes were also reported for Cu⁺ niobates,⁵⁴ tantalates,⁵⁵ and vanadates.⁵⁶

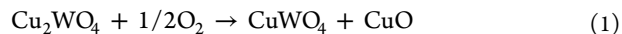


Figure S7 maps the changes in the optical absorption profiles (as established by diffuse reflectance spectroscopy, DRS) of the arc-synthesized Cu₂WO₄ samples as a function of the thermal annealing temperature. No optical changes were observed up to 260 °C, and the absorption cutoff gradually broadened thereafter with significant absorption below the band gap of the original compound, followed by darkening of the film. The DRS data of the fully oxidized sample at 300 °C do not match those of pure CuWO₄, which is expected to show an optical band gap of ~2.2 eV. This finding further supports thermal oxidation of Cu₂WO₄ occurring via oxidation to CuWO₄ and expulsion of CuO from the compound framework (eq 1), the latter being known to absorb light at wavelengths longer than 800 nm.⁵⁷

Scanning electron microscopy (SEM) images were obtained for the electrodes annealed at 250 and 300 °C (Figures S8 and S9, respectively). For the electrode annealed at 250 °C, SEM image showed a porous film with effective substrate coverage, mainly constituted of particles with ill-defined morphology and particle size in the range of submicrometer to few microns (Figure S8a). Different points of the film were analyzed by energy-dispersive X-ray (EDS) spectroscopy (Figure S8b) to obtain elemental composition of the electrode surface. As shown in Table S2, the Cu/W mass ratio was 0.43, which strongly agreed with the expected ratio of 0.41, and within the uncertainty limits inherent in the EDS data. Cross-sectional images of the photoelectrode are shown in Figure S8c,d, which indicated a porous film of Cu_2WO_4 with homogeneous thickness in the range of 7–8 μm .

Figure S8 shows the Cu_2WO_4 photoelectrode surface after annealing at 300 °C. The oxidation process clearly induced a change in morphology, with increased particle size and formation of aggregates. The separate frames on the right-hand side of Figure S9 are zoomed-in sections of the oxidized film, where small, spherical domains decorate larger particles of oxides. This feature is similar to that observed in other oxidized Cu(I) multimetallic oxides, such as Cu/Nb and Cu/Nb/Ta oxides.^{54,55}

X-ray photoelectron spectroscopy (XPS) data are presented in Figure 7. Typically, copper exhibits a doublet associated with Cu 2p electrons at ~ 933.0 eV (Cu $3p_{3/2}$). Copper(II) species presented additional strong satellite peaks, which can be used to distinguish them from the monovalent copper species (which show only weak satellite peaks). The two oxidation states also differ in binding energy and full width at half-maximum (i.e., broader peak for the Cu^{2+} signal).⁵⁸

For the sample annealed at 250 °C (Figure 7), Cu sites were seen mainly in the +1 oxidation state, as expected for Cu_2WO_4 . Trace amounts of Cu^{2+} could be attributed to charge compensation, owing to oxygen vacancies in the crystal structure. As the sample was preannealed at different temperatures, the contribution of the +2 oxidation state gradually increased. Concomitantly, the intensity of satellite peaks in the range of 940–945 eV, attributed to Cu(II), significantly increased as well. A major change in the $\text{Cu}^+/\text{Cu}^{2+}$ ratio was observed after annealing at 270 °C.

Different binding energies were also observed for the oxygen atoms in the O 1s binding energy regime (Figure S10). Such a profile is also expected for a mixed-valent copper oxide, as previously reported in the literature for the controlled reduction of CuO to Cu_2O .⁵⁸ Interestingly, all of the charge compensation stemming from oxygen deficiency in the tungstate compound was provided by copper in different oxidation states, while tungsten atoms remained solely in the +6 oxidation state, as shown in Figure S11.

The observation of Cu(I) in Cu_2WO_4 in the XPS data is supported by the crystal structure, where the low coordination numbers (2 and 3) around Cu sites¹⁷ are reminiscent of the coordination environments observed in CuAl_2O_2 ⁵⁹ and SrCu_2O_2 .⁶⁰ These XRD and XPS results also demonstrate that controlled $\text{Cu}^+/\text{Cu}^{2+}$ mixed-valence behavior in copper tungstate can be obtained by simple thermal annealing in air.

The combined results obtained from XRD and XPS show that pure, crystalline Cu_2WO_4 can be obtained by arc synthesis and that it can be thermally oxidized to systematically change the $\text{Cu}^+/\text{Cu}^{2+}$ ratio. The crystal structure of Cu_2WO_4 can accommodate Cu^{2+} and O vacancies to a certain extent in the

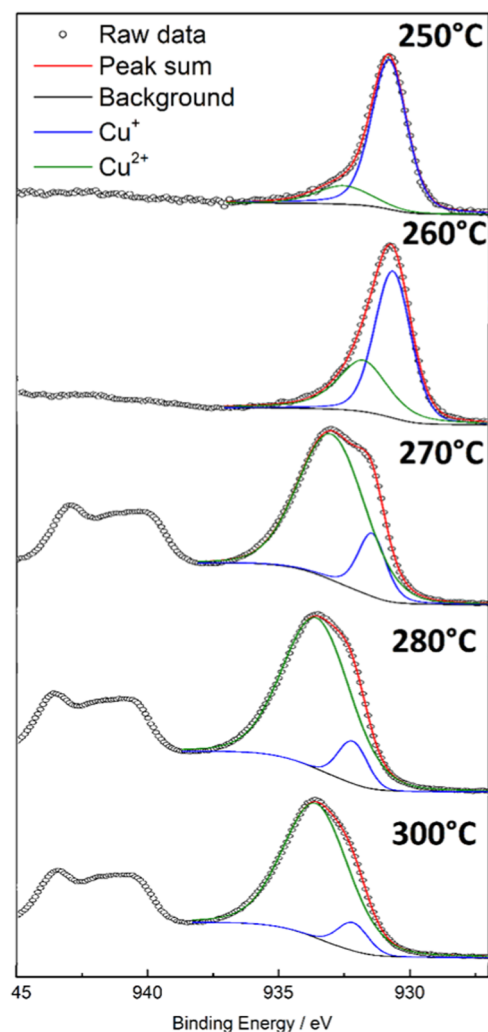


Figure 7. High-resolution XPS Cu 2p analyses of the Cu_2WO_4 films after annealing at different temperatures: 250, 260, 270, 280, and 300 °C.

mixed-valence case. However, ultimately, a switch in the array of atoms in space takes place, culminating in an XRD pattern with characteristic diffraction peaks of CuWO_4 above 260 °C.

3.5. Photoelectrochemical Behavior of Cu_2WO_4 and Effects of Thermal Preannealing. Figure 8 shows the photoelectrochemical (PEC) properties of electrodes constituted from arc-synthesized copper tungstate particles, immobilized on an FTO-glass substrate. Aqueous NaHCO_3 solution was used as the supporting electrolyte and saturated with dissolved CO_2 . The linear sweep voltammograms (at 3 mV s^{-1}) shown in Figure 8 were acquired by scanning the applied potential from the rest value (E_{eq}) to -0.5 V under intermittent illumination provided by a solar simulator (100 mW cm^{-2}). Prior thermal annealing at 150 °C did not cause any visible change in the XRD, XPS, and DRS analyses compared to the as-synthesized sample, while adhesion to the FTO substrate was promoted. Therefore, this sample could be represented as Cu_2WO_4 . Since a cathodic photocurrent was observed which increased at more negative (reverse bias) potentials, arc-synthesized Cu_2WO_4 was diagnosed as a p-type semiconductor.

On band-gap photoexcitation, the photogenerated holes migrate to the back contact, while the electrons at the

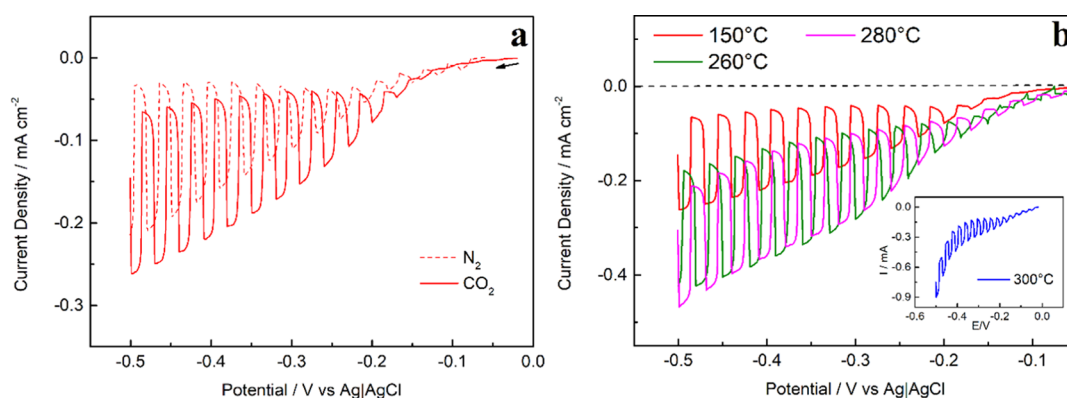


Figure 8. (a) Photovoltammograms obtained under intermittent light incidence in aqueous 0.5 M NaHCO₃ solution from Cu₂WO₄ electrodes preannealed at 150 °C, in electrolyte saturated with N₂ and CO₂. (b) Photovoltammograms for FTO-Cu₂WO₄ electrodes preannealed at 150, 260, and 280 °C (red, green, and magenta lines, respectively) in solutions saturated with CO₂. The inset shows data for the sample preannealed at 300 °C.

electrode surface react with dissolved CO₂ in the aqueous solution interface. It is worth noting here that the photocurrent was significantly higher in the presence of CO₂, compared to the N₂-bubbled electrolyte case (Figure 8a). We assume that the photogenerated species reduce dissolved CO₂ species in solution via redox processes that are beyond the scope of discussion in the present study.

As the sample was further oxidized (e.g., after 260 °C thermal preannealing), the photocurrent was slightly enhanced, while the dark current also increased to a significant extent (Figure 8b). The behavior now reflects that of the mixed-valence tungstate. At some point, the amount of Cu²⁺ sites became detrimental to the photoactivity, and electroactivity in the dark became dominant (300 °C annealing, Figure 8b, inset), although some cathodic photocurrent is still observed, which agrees with CuWO₄ being formed along with p-type CuO via the oxidation pathway described in eq 1.

As copper(II) tungstate is known to be an n-type semiconductor, metal-like behavior is expected at forward bias and this is indeed reflected in the data in Figure 8 (inset). This switch in both structures (cf. Figure 6) and electronic properties accounts for the trends with annealing temperature, seen in Figure 8. Figure S12 shows the correlation between Cu²⁺ content extracted from XPS data and the dark current at the selected potential of -0.45 V vs Ag/AgCl. Interestingly, different ratios of mixed valence in a broad range did not alter the dark current values noticeably.

Finally, as shown in Figure S13, flat band potentials (V_{fb}) of Cu₂WO₄ were experimentally obtained by the Butler–Gärtner methodology⁶¹ (see the Supporting Information). Interestingly, the values for V_{fb} of Cu₂WO₄ and the mixed-valence copper tungstates were the same. The V_{fb} and E_{bg} values can be combined to obtain band edge positions (VB and CB) for a given oxide. Figure 9 shows the band positions of Cu₂WO₄ thus obtained in this work, compared to literature values for Cu₂O and other p-type, Cu⁺ complex oxides.^{54,56,62,63} In constructing this diagram using the data from Figure S13 and literature sources, a Nernstian correction (of -0.059 V/pH) was also made when needed, to obtain values on the reversible hydrogen electrode (RHE) scale.

The literature data¹² for CuWO₄ band edge positions indicate a much more positive conduction band minimum. However, Cu₂WO₄ has a 3d¹⁰ electronic configuration; therefore, the CB is mainly composed of W 5d⁰ orbitals. On

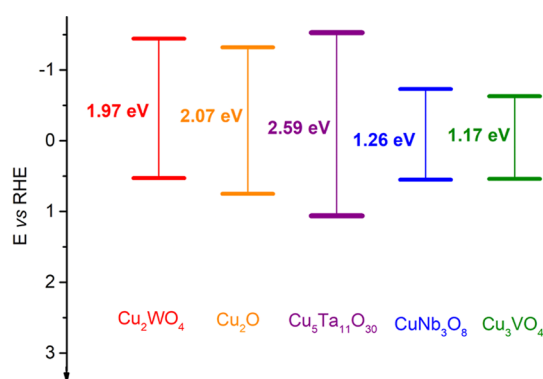


Figure 9. Band positions of arc-melted Cu₂WO₄ determined by combining optical band gap measurements and flat band potentials (V_{fb}) extracted from Butler–Gärtner plots.

the other hand, CuWO₄ has a 3d⁹ configuration leading to a different orbital composition for the corresponding CB. As can be seen in Figure 5, our calculations show that CuWO₄ has a mid-gap state mainly composed of Cu-d orbitals. Experimental determination of CuWO₄ CB energy¹² did not initially consider this feature. In fact, the same authors later discussed this mid-gap state in a more detailed EIS study.⁶⁴ Insights on the influence of mid-gap states when determining CBM values of CuWO₄ are provided in more recent work from the same authors.⁶⁵

In terms of structural aspects, the copper coordination environment has an important influence on the energy levels of metal oxide semiconductors. For instance, a higher coordination of copper atoms tends to provide narrow band gaps for some family of copper complex oxides,⁵ although more commonly by shifting the valence band.

The very negative CB edge positions for the p-type copper oxides (Figure 9) underline why they are of much current interest. A high driving force (i.e., overpotential) for photoredox reactions involving negative redox potentials, is intrinsic to the electronic structure of these compounds. This feature could be exploited for many practical device applications including energy conversion and sensors.

Stability issues hamper the widespread use of copper oxides, especially for photoelectrochemical applications. For example, the parent oxide Cu₂O is known to undergo photoreduction (to metallic copper) in aqueous solutions, as the potential for

this redox process lies inside the energy band gap.^{66–68} Therefore, the photoelectrochemical stability of Cu_2WO_4 was compared with that of Cu_2O ; the data are contained in Figure 10.

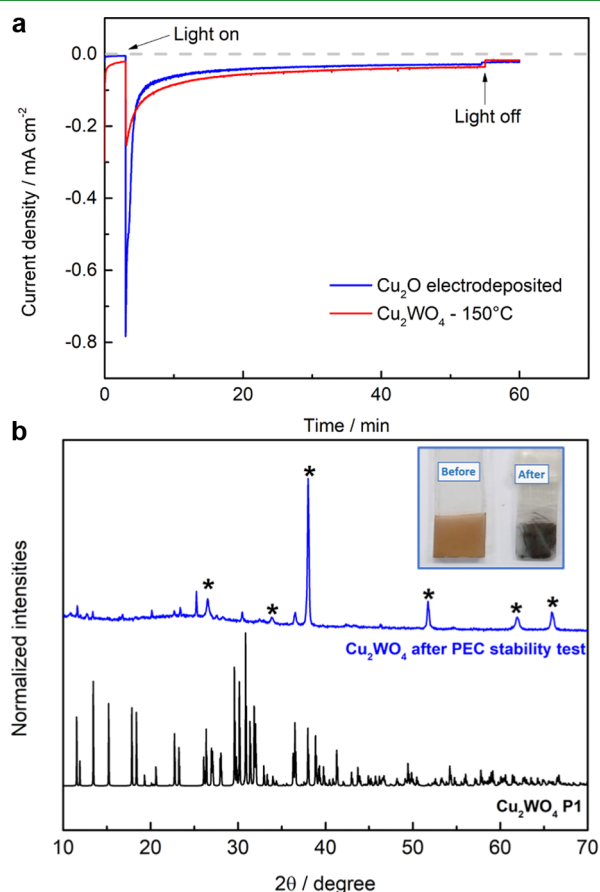


Figure 10. (a) Chronoamperometric PEC stability test performed at -0.3 V vs Ag/AgCl under 1 sun illumination for 60 min. (b) XRD analysis of Cu_2WO_4 cathode after stability test.

Electrodes of Cu_2WO_4 were preannealed at 150 °C and electrodeposited Cu_2O were polarized at -0.3 V and kept under illumination for 50 min. As expected, both Cu_2O and Cu_2WO_4 underwent photocorrosion as demonstrated by the gradual photocurrent decay with time (Figure 10a). Nonetheless, the decay was more pronounced for Cu_2O , indicating enhanced stability for the Cu_2WO_4 photocathode, although severe photocorrosion did occur. Figure 10b shows the XRD analysis of the Cu_2WO_4 photoelectrode after the stability test. The main diffraction peaks were identified as the FTO substrate, indicating detachment of copper tungstate from the conductive support. Minor diffraction peaks could be related to CuWO_4 , although not unambiguously. Formation of CuO or Cu was ruled out, which suggests photocorrosion leading to CuWO_4 accompanied by copper leaching into the solution. The mechanistic aspects of photocorrosion of Cu_2WO_4 , as well as different strategies for electrode protection against photocorrosion, constitute topics beyond the scope of this study, and will be the subject of follow-up work.

4. CONCLUSIONS

In conclusion, a hitherto little-studied Cu(I) tungstate (Cu_2WO_4) was studied by a combined theoretical/exper-

imental approach. The long-standing debate on the crystal structure was resolved, and Cu_2WO_4 was theoretically shown to belong to the $P1$ space group. Cu_2WO_4 was obtained by a simple, time-efficient synthesis using an arc furnace. This material possessed broad-band light absorption in the entire UV–vis spectrum. Photoelectrodes of Cu_2WO_4 showed cathodic photocurrent in the presence of CO_2 in aqueous solution, when polarized at more negative potentials than the rest potential; this was diagnostic of p-type semiconductor behavior. Prior thermal annealing of Cu_2WO_4 was demonstrated to induce mixed-valent behavior up to 260 °C, with higher temperatures inducing oxidation to CuWO_4 along with expulsion of CuO from the compound structure. Finally, the present study has served to address a significant literature void that existed for an interesting ternary copper oxide based on earth-abundant elements and obtained by simple processing methods, two important aspects for reaching economically feasible solar fuels technologies.⁶⁹

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acsami.1c03928>.

Calculated infrared spectra and atomic positions of relaxed geometries for structures based on the two space groups; partial charge densities at the VBM and CBM; thermal analysis data; effect of preannealing on XRD and DRS data; high-resolution XPS data in the O 1s and W 4f domains as a function of preannealing temperature; SEM image of Cu_2WO_4 photoelectrode; Butler–Gartner analyses; Bader charge analyses for copper(I) tungstate and copper(II) tungstate; and images of arc synthesis setup (PDF)

■ AUTHOR INFORMATION

Corresponding Authors

Nora H. de Leeuw – Department of Earth Sciences, Utrecht University, 3548CB Utrecht, The Netherlands; School of Chemistry, University of Leeds, Leeds LS2 9JT, United Kingdom; Email: n.h.deleeuw@uu.nl, n.h.deleeuw@leeds.ac.uk

Claudia Longo – Institute of Chemistry, University of Campinas—UNICAMP, 13083-970 Campinas, Brazil; Center for Innovation on New Energies, University of Campinas, 13083-841 Campinas, Brazil; orcid.org/0000-0001-8955-148X; Email: clalongo@unicamp.br

Authors

Miguel Tayar Galante – Institute of Chemistry, University of Campinas—UNICAMP, 13083-970 Campinas, Brazil; Center for Innovation on New Energies, University of Campinas, 13083-841 Campinas, Brazil; orcid.org/0000-0003-1697-0039

Aleksandar Živković – Department of Earth Sciences, Utrecht University, 3548CB Utrecht, The Netherlands; orcid.org/0000-0003-1347-6203

Jéssica Costa Alvim – Institute of Chemistry, University of Campinas—UNICAMP, 13083-970 Campinas, Brazil; Center for Innovation on New Energies, University of Campinas, 13083-841 Campinas, Brazil

Cinthia Cristina Calchi Kleiner – Faculty of Mechanical Engineering, University of Campinas—UNICAMP, 13083-970 Campinas, Brazil

Márcio Sangali – Faculty of Mechanical Engineering, University of Campinas—UNICAMP, 13083-970 Campinas, Brazil

S. F. Rebecca Taylor – Department of Chemical Engineering and Analytical Science, University of Manchester, Manchester M13 9PL, United Kingdom; orcid.org/0000-0002-6175-2631

Adam J. Greer – Department of Chemical Engineering and Analytical Science, University of Manchester, Manchester M13 9PL, United Kingdom; orcid.org/0000-0003-1639-5433

Christopher Hardacre – Department of Chemical Engineering and Analytical Science, University of Manchester, Manchester M13 9PL, United Kingdom; orcid.org/0000-0001-7256-6765

Krishnan Rajeshwar – Department of Chemistry and Biochemistry, The University of Texas at Arlington, Arlington, Texas 76019, United States; orcid.org/0000-0003-4917-7790

Rubens Caram – Faculty of Mechanical Engineering, University of Campinas—UNICAMP, 13083-970 Campinas, Brazil

Rodnei Bertazzoli – Faculty of Mechanical Engineering, University of Campinas—UNICAMP, 13083-970 Campinas, Brazil

Robin T. Macaluso – Department of Chemistry and Biochemistry, The University of Texas at Arlington, Arlington, Texas 76019, United States; orcid.org/0000-0002-0021-0775

Complete contact information is available at: <https://pubs.acs.org/10.1021/acsami.1c03928>

Author Contributions

[○]M.T.G. and A.Ž. contributed equally to this work.

Notes

The authors declare no competing financial interest.

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