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TiO₂/Cu₂O/CuO Multi-Nanolayers as Sensors for H₂ and VOCs: An Experimental and Theoretical Investigation

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ABSTRACT

Highly sensitive TiO₂/Cu₂O/CuO multi-nanolayers have been grown in various thicknesses by a cost-effective and reproducible combined spray-sputtering-annealing approach. The ultra-thin TiO₂ films were deposited by spray pyrolysis on top of sputtered-annealed Cu₂O/CuO nanolayers to enhance their gas sensing performance and to improve their protection against corrosion at high operating temperatures. The prepared heterostructures have been investigated using scanning electron microscopy (SEM), X-ray diffraction (XRD), ultraviolet visible (UV-Vis) and microRaman spectroscopy. The gas sensing properties were measured at several operating temperatures, where the nanolayered sensors with oxide thicknesses of between 20 and 30 nm (Cu₂O/CuO nanolayers) exhibited a high response and excellent selectivity to ethanol vapour only after thermal annealing at 420°C. The results obtained at an operating temperature of 350 °C demonstrate that the CuO/Cu₂O nanolayers with a thickness between 20 and 30 nm are sensitive mainly to ethanol vapour, with a response of ~150. The response changes from ethanol vapors to hydrogen gas as CuO/Cu₂O nanolayers thickness changes from 50 nm to 20 nm.

Density functional theory-based calculations were carried out of the geometries of the CuO($\bar{1}11$)/Cu₂O(111) and TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) heterostructures and their sensing mechanism towards alcohols of different chain lengths and molecular hydrogen. The reconstructed hexagonal Cu₂O(111) surface and the reconstructed monoclinic CuO($\bar{1}11$) and TiO₂(111) facets, all terminated in an O layer, lead to the lowest surface energies for each isolated material. We studied the formation of the binary and ternary heteroepitaxial interfaces for the surface planes with the best matching lattices. Despite the impact of the Cu₂O(111) substrate in lowering the atomic charges of the CuO($\bar{1}11$) adlayer in the binary sensor, we found that it is the different surface structures of the CuO($\bar{1}11$)/Cu₂O(111) and TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) devices that are fundamental in driving the change in the sensitivity response observed experimentally.

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The experimental data presented here, supported by the computational results, promote the use of the multi-nanolayered films tested in this work as reliable, accurate and selective sensor structures for the tracking of gases at low concentrations.

KEYWORDS: Nanolayers, Nanomaterials, Multilayered Films, CuO, *p*-type, Cu₂O, TiO₂, sensor

1. Introduction

Functional nanomaterials, including semiconducting oxide heterostructures with tunable performances, are an essential part of semiconductor-powered devices. However, synthesizing such nanocomposites has to be highly specific with respect to phase control at the nanoscopic level. Heterojunctions between different semiconductor oxide nanocrystals, especially based on ultrathin films with mixed phases, may improve the characteristics of gas sensors due to their unique detecting mechanism¹⁻⁴. The specific features of the heterojunction in nanocrystalline multi-layered composites is crucial to control the gas sensing characteristics, i.e. selectivity and gas response of the sensor, as a result of the top surface and interface phenomena^{1,5,6}. A seminal work by Brattain and Bardeen⁷ reported that gas adsorption on semiconducting surfaces produces a change of its electrical conductance⁷, which has contributed to the further development of the sensor industry based on solid state materials.

From the nanotechnology point of view, metallic copper (Cu) and its oxides have received much attention due to their variety of real applications, particularly in the field of new nanotechnology components for microelectronics^{4,8,9}. Copper oxides are *p*-type semiconducting oxides and can be obtained in forms such as cuprite (Cu₂O) and cupric oxide (CuO), which depends

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on the valence state of copper. The cuprous oxide cuprite (Cu₂O) is amongst the earliest semiconducting oxides used in solid state electronics^{6,10,11}. Although cuprite has been the focus of numerous experimental and theoretical studies^{1,7,12–14}, including its vibrational and optical properties, the electronic properties of Cu₂O continue to puzzle the scientific community. As applications of Cu₂O in nanoelectronics, photovoltaics, solid-state electronics, biosensing and spintronics emerge,^{4,9,15} as well as light-driven purification of wastewater,⁸ understanding at the atomic level of the electronic structure of Cu₂O is important for the control its properties and of identification of future applications in devices or nanodevices.

CuO has a band gap in the range 1.2 – 2.1 eV, with *p*-type semiconducting properties¹² with significant potential for sensing applications, especially in mixed CuO/Cu₂O phases^{6,11}. A summary of gas sensor structures based on copper oxide nanomaterials can be found in several reviews^{16–20}.

The (111) surface has been found, both in simulations^{13,21} and experimentally²², to be the most stable cuprite (Cu₂O) plane under a range of different conditions. Furthermore, the almost complementary ($\bar{1}11$) facet was observed to be highly prominent in the crystals of the more oxidised tenorite (CuO) phase²³. Although less stable than other planes, the anatase TiO₂(111) surface has been reported as one of the most reactive for photocatalytic applications²⁴ and in H₂ evolution²⁵.

Inorganic UV absorbers such as TiO₂, ZnO and CeO₂ are generally employed in shielding applications for the ultraviolet (UV) protection of different surfaces^{26,27}, where effective physical nano-coating barriers are needed for high temperature applications²⁸. Titania (TiO₂) nano-coatings are used extensively to increase surface hardness and adhesive strength; to provide long term and high temperature protection against corrosion; to enhance tribological properties; and to improve the design of the transparent coatings of self-cleaning surfaces^{27–30}.

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Detection and discrimination of volatile organic compounds (VOCs), which are classified as hazard vapors with adverse short- and long-term consequences on the environment and human health³¹, is important in the continuous monitoring of indoor air quality, which therefore requires reliable sensors. Moreover, exhaled VOCs can serve as biomarkers to assist in the non-invasive identification of various diseases. For example, acetone indicates diabetes^{14,32–35}, whereas isoprene, toluene, and acetic acid are signals of lung cancer,^{31,36–38} and breath testing is therefore a highly promising approach for non-invasive cancer screening³⁹. VOC analysis in patient breath offers insight into the metabolic processes in the anatomy/physiology that are altered by underlying diseases^{40,41}, although a detailed impression of the metabolic route leading to these molecules is still under investigation³⁹.

This study reports the fabrication of stable ethanol sensors using multi-nanolayered films consisting of titania/cuprite/cupric oxide (TiO₂/CuO/Cu₂O/glass). The cuprite/cupric oxide structures are produced by a method based on sputtering combined with thermal annealing, before spraying titania on the surface of the CuO/Cu₂O/glass substrate. Interdigitated Au-electrodes are deposited on the top surface of the specimens to connect heterostructures and perform gas sensing tests, and the morphology composition, structure, electrical transport and gas sensing properties of the materials are studied. Moreover, a theoretical discussion developed from density functional theory (DFT) calculations of the binary heterojunction CuO($\bar{1}11$)/Cu₂O(111) and the ternary heterostructure TiO₂(111)/CuO ($\bar{1}11$)/Cu₂O(111) have been performed. First, we have investigated the structures and relative stabilities of the hexagonal and monoclinic surfaces for each isolated material, before elucidating the arrangement of the heteroepitaxial junctions and reporting their work function values and scanning tunnelling microscopy (STM) images. The calculated values of the work function rationalise the reactivity trends of the binary and ternary heterojunctions, whereas their computed STM images compare well with the experimental SEM

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images. The adsorption energies have been computed for molecular hydrogen (H₂), ethanol (C₂H₅OH) and *n*-butanol (*n*-C₄H₉OH) and we have plotted the electron charge density flow after the interaction with the heterojunctions in order to interrogate their selectivity changes.

2. Experimental Section

Glass slides from ThermoScientific (2.5 × 7.5 cm) were employed as the substrates for the device development. The glass substrates were cleaned by dipping in HCl (11%) and then rinsed with distilled H₂O and acetone for 11 min, which was followed by ultrasonic bath in ethanol for 11 min and then rinsed in deionized H₂O^{1,42}. Afterward, the CuO-Cu₂O ultrathin layers with thicknesses of 20, 30, 40, 50 and 60 nm were prepared on top of the clean glass by sputtering metallic copper under vacuum conditions. The Cu sputtering was carried out using a custom-built RF-magnetron system at 25°C, a pressure of 3.6 × 10⁻³ mbar, an Argon gas flux of 22 sccm and a power of 51 W. Evochem GmbH, Germany supplied the copper metal (with high purity: 5n = 99,999%, radius of 2.5 cm). The deposition rate of 6 nm/min was determined experimentally by using a profilometer. Then, the metallic copper layers deposited on the glass substrates were thermally treated at 420°C under normal atmospheric conditions for 30 or 60 min. The temperature for the thermal treatment was in accordance with our previous work of the ultrathin mixed CuO-Cu₂O oxide film^{6,43}. Five different sample sets were produced with thicknesses of 20, 30, 40, 50 and 60 nm for the CuO-Cu₂O layer on the microscopic glass substrate. Next, titania (TiO₂) films were spray-pyrolysis-deposited on top of the mixed copper oxide phase layers to prepare another five sample sets. Afterwards, all materials were mounted on a thermal heating plate, where we kept the temperature at 420°C for 25 min prior to starting the spray pyrolysis process, as published previously by Pauporté *et al.*^{27,42,44}. For the spray pyrolysis, the precursor was delivered as a mixture of 7.1 ml

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of isopropanol, 0.62 ml of titanium (IV) tetra-isopropoxide (TTIP) and 0.41 ml of acetylacetone.

The carrier gas used was an oxygen flow, which was selected as previously reported^{42,44}, in order to blow the mixed aerosol through a valve with a diameter of 10 mm directly on the surface of CuO/Cu₂O/glass composite, which remained on the heated hot plate at 420°C for the entire spray process. The CuO/Cu₂O samples were grown with different thicknesses of 20, 30, 40, 50 and 60 nm, labeled as Cu20, Cu30, Cu40, Cu50 and Cu60, respectively. Afterwards, the TiO₂/CuO/Cu₂O heterostructures were treated under air at 420°C for another 30 min (Cu20, Cu30, Cu40) or 60 min (Cu50, Cu60), depending on their thickness, prior to allowing them to cool down spontaneously. The thickness of the TiO₂ layer was monitored throughout the duration of the spray, as reported before^{42,44}.

After the preparation of materials, Au electrodes were grown on top of the nanolayered TiO₂/CuO/Cu₂O samples through an Al meander-shaped mask^{6,43}. The Au top-contacts have a thickness of is ~ 180 nm and a separation of 1 mm between gold contacts. The Au target (purity: 4n = 99,99%, radius: 2.5 cm) which was produced by Evochem GmbH, Germany, was mounted on the magnetron (DC). The chamber pressure during Au sputtering was 3.55×10^{-3} mbar, whereas the flowrate of Ar was about 16 sccm and the power of sputtering was set at 51 W, enabling a deposition rate of about 47 nm/min⁴². The morphological, structural, chemical, micro-Raman (MR) measurements and X-ray photoelectron spectroscopy (XPS) investigations were carried out as before^{1,45}. MR experiments were done using a Raman WITec Alpha300 RA spectrometer at 22°C, as reported before⁴⁶. A graphite monochromatized CuK₁ radiation (1.5405 Å) at 40 kV and 40 mA were used for the X-ray diffraction (XRD), which was carried out using a Seifert 3000 TT instrument,³³ and X-ray photoelectron spectroscopy (XPS) was used to measure the thickness of the TiO₂/CuO thin films by using an Omicron Nano-Technology GmbH, Al-anode, P=240 W, as reported previously⁴². We charge calibrated the documented spectra using the signal at 284.5

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eV corresponding to the aliphatic carbon C-1s and the “CasaXPS”, software version 2.3.16. We employed a Varian Cary 5000 spectrophotometer to carry out the characterization of the optical properties of the samples, where we used the integrating sphere supplied in the wavelength values between 300 and 2500 nm, as before⁴⁷. The gas detection characteristics were obtained using the set-up and protocol described previously^{6,43,48,49} at 30% relative humidity (RH). A computer-controlled Keithley2400 sourcemeter at 0.25 V applied bias voltage was used to record continuously the electrical measurements, which were processed through the LabView software (from National Instruments). The responses to gas and VOCs were defined as the ratio $\left(\frac{R_g - R_a}{R_a} \cdot 100\%\right)$, where R_g and R_a - the electrical resistances of the specimens exposed to gas/VOC and air under normal environmental conditions, respectively^{6,43}.

The surface properties of the binary and ternary heteroepitaxial interfaces were simulated using unrestricted density functional theory (DFT) simulations, which are described in **Text S1** (Supporting Information).

Figure 1 shows the technological flow for the manufacture of the mixed-phase: (a) CuO/Cu₂O (Final device set #1), and (b) TiO₂/CuO/Cu₂O (Final device set #2) nanofilm devices. The general process can be described as follows: Step 1; Pre-cleaned glass substrate is sputtered with copper nanoparticles with a radius of about 2-5 nm to obtain ultrathin films of copper with thicknesses between 20 and 60 nm; Step 2; thermal annealing in a furnace at the temperature of 420°C for 30 min (for sample sets Cu20, Cu30 and Cu40) or 60 min (for sample sets Cu50 and Cu60) in air to develop the CuO/Cu₂O nanolayer-heterojunctions. Steps 3 and 4 in **Figure 1a** are the deposition of the Au contacts by shading a metallic meander mask with a 1 mm gap. In step 5, we obtain the device 1 based on the CuO/Cu₂O nanolayers. Following steps 3 and 4 in **Figure 1b**, where the thin films of TiO₂ are deposited with a thickness of 20 nm and then thermally treated in a furnace at the temperature of 420°C for 30 min (for sample sets Cu20, Cu30 and Cu40) or 60

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min (for sample sets Cu50 and Cu60) in air, we obtain the sensitive multi-nanolayered TiO₂/CuO/Cu₂O heterojunction film. Steps 5 and 6 in **Figure 1b** are the deposition of the Au contacts as in **Figure 1a**. Finally, step 7 represents the final device set #2, i.e. the TiO₂/CuO/Cu₂O heterostructure. It is important to mention that the layers are drawn as straight lines in **Figure S1** but, in reality, they are rough polycrystalline layers (**Figure 2**). **Figure S1** displays the cross-section view of the devices from the set #1 made of CuO/Cu₂O nanolayers (Device 1), and set #2, which are based on heterolayers of TiO₂/CuO/Cu₂O heterolayer films (Device 2).

3. Results and Discussion

3.1. Morphology and composition analyses

Figure 2a-f presents the low and high (in the insets) magnification SEM images of the nanocrystallite CuO/Cu₂O samples, which were grown using the reproducible spraying/sputtering/annealing approach, owing to the possibility of controlling the speed of sputtering growth. We thus obtained five samples simultaneously from a single substrate before thermal annealing at 420°C for 30 min (for sample sets Cu20, Cu30 and Cu40) or 60 min (for sample sets Cu50 and Cu60) in air. It can be clearly observed that the volume of nanoparticles essentially changes with the film thickness, which also affects the sensing properties. For the CuO nanoparticles, their larger electrical conductivity leads to some bright dots, which are seen in **Figure 2a-c**,⁴³ in the areas exposed to the electron beam of the SEM. The nanoparticles, which appear much smaller in size in the SEM images due to thinner films, oxidize completely during the 30 min of annealing.

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The ultra-thin films possess very good adhesion to the microscopic glass substrates, as we have not seen any signs of delamination during the three years that we have been investigating these types of samples ^{1,12}.

3.2. X-ray photoelectron spectroscopy and Micro-Raman characterization

For the TiO₂/CuO/Cu₂O layered thin films sensor, the X-ray photoelectron spectra (XPS) is presented in **Figure 3**, where the overview spectrum, shown in **Figure 3a**, attests to the presence of the Cu, O, Ti, Na and C elements. Cu, O and Ti originate from the TiO₂/CuO layers whereas the signal from carbon is as reported before ^{1,48}. The Na peak corresponds to the glass substrates ⁴² used for the deposition of the sensor structures, since samples were cut for XPS.

High-resolution Cu-2p and Ti-2p lines are depicted in **Figure 3b**. A closer look at the Cu-2p line clearly reveals the Cu-2p_{1/2} and Cu-2p_{3/2} satellite lines, which are shifted to higher binding energies. The observed satellite peaks in the spectrum are commonly noted a signature for the presence of CuO, ^{43,50} which is found in the base layer of our samples.

The signals between 459.6 - 458.0 eV are due to Ti-2p_{3/2}, which is usually assigned to Ti in TiO₂. The evaluation of the high resolution Ti-2p XPS spectra reveals that the line at 458.3eV corresponds to Ti-2p_{3/2}. The separation of 5.6 eV between the Ti-2p_{3/2} and Ti-2p_{1/2} peaks and the position of the Ti-2p_{3/2} signal, illustrate the presence of Ti as TiO₂ in the base layer, according to the literature ^{51,52}.

The Raman scattering method is a useful spectroscopic technique to measure the vibrational modes and phase of ultra-thin layers and nanomaterials, owing to its ability to determine the vibrational modes of the heterostructure ^{11,45}. Micro-Raman spectroscopy was employed to investigate the characteristics at the nano-scale, namely the lattice dynamics

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(electron-phonon interaction) of the CuO/Cu₂O and TiO₂/CuO/Cu₂O nano-materials. The micro-Raman spectra in the range 100-1000 cm⁻¹ were obtained at room temperature for the CuO/Cu₂O and TiO₂/CuO/Cu₂O nano-materials, as shown in **Figure S2** and **Figure S3**.

The Raman studies clearly show the formation of the mixed copper oxide phases, namely CuO/Cu₂O, following thermal annealing at 420°C under ambient conditions, as well as the TiO₂/CuO/Cu₂O heterostructure after spraying a TiO₂ nano-layer on top of the binary films, which show the existence of heterostructured mixed phase films. A detailed description of the aforementioned results is given in **Text S2** (Supporting Information).

3.3. Ultraviolet, Visible and Near Infrared spectroscopy

Ultraviolet, Visible and Near InfraRed (UV-Vis-NIR) absorption spectroscopy is a characterization method used to study energy level as well as optical properties of transparent semiconducting oxide materials. The room temperature spectra of the CuO/Cu₂O and TiO₂/CuO/Cu₂O heterostructures allowed the detection of the optical absorption and excitonic transitions characteristic of the nanolayers. The transmission and absorption spectra are presented in **Figure S4**, and the plots of the $(\alpha h\nu)^2$ versus the photon energy ($h\nu$) for the CuO/Cu₂O and TiO₂/CuO/Cu₂O heterostructures are shown in **Figure S5** and **Figure S6**. More details of the UV-Vis-NIR characterization are provided in **Text S3** (Supporting Information).

3.4. Gas sensing properties

First, we will focus on the CuO/Cu₂O samples and their gas sensing performances, followed by the gas sensing performances of the TiO₂/CuO/Cu₂O nanolayered materials, which we will compare to the base layer.

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Gas sensing results of CuO/Cu₂O. The CuO/Cu₂O based Cu₂₀, Cu₃₀, Cu₄₀, Cu₅₀ and Cu₆₀ nanolayer samples were connected in the sensor structures as indicated schematically in **Figure S1**.

Figure 4a represents the responses at the standard deviations between multiple measurements of the CuO/Cu₂O samples to hydrogen and ethanol, at the operating temperature of 350°C, from which it is clear that Cu₂₀ and Cu₃₀ samples show the highest response to ethanol. From **Figure 4a** we observe that, for all thicknesses, the CuO/Cu₂O specimens are more selective to ethanol compared to the other tested gases (hydrogen, ethanol, 2-propanol, *n*-butanol, acetone and ammonia). However, the response and selectivity of the sensors decrease as the thickness of the layers increases (more than 30–40 nm), because with increasing thickness of the layers, the electrical resistance of the structures decreases, which is in accordance with previously reported data ⁶. The optimum thickness is therefore in the range 20–30 nm (sample sets Cu₂₀ and Cu₃₀). The response values of CuO/Cu₂O nanolayers to different analytes are depend on their thickness, specifically when the thickness is in the order of the Debye length ^{43,53}. The sensing mechanism proposed for these results from the binary heterostructure is explained in **Text S4** (Supporting Information).

Figure 4b represents that the current-voltage (*I-V*) curves of the prepared structures, and from **Figure S7a**, it can be seen that the characteristics are linear at room temperature for all samples. According to the data in **Figure 4b** and **Figure S7a**, the *I-V* characteristics for all the CuO/Cu₂O samples of different thicknesses show Ohmic contact behavior between the CuO/Cu₂O nanolayers and the Au top contact at room temperature. However, at the operating temperature of 350 °C (**Figure S8a**) the *I-V* current-voltage characteristics is non-linear, which can be attributed to a conductivity effect driven by the energy barrier. The variation of the electrical current with temperature, see **Figure 4c**, indicates that the heterostructure is also convenient for temperature

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measurement. We have used equation (1) to express the electrical resistivity of our *p*-type semiconducting oxides⁵⁴:

$$\rho \equiv \frac{1}{\sigma} = \frac{1}{q\mu_p p} = \frac{1}{q\mu_p \left(N_V e^{\left[-\left(\frac{E_F - E_V}{kT} \right) \right]} \right)} = \frac{e^{\left(\frac{E_F - E_V}{kT} \right)}}{\sigma \mu_p N_V} \quad (1)$$

where ρ is the electrical resistivity, σ is the electrical conductivity, q is the charge of the electron, μ_p is the holes mobility, p is the holes concentration, N_V is the acceptors concentration, E_F is the Fermi energy, E_V is the maximum energy of the valence band, T is the absolute temperature and k is the Boltzmann constant.

The response variation of the structures to 100 ppm of ethanol as a behavior of the operating temperature was studied to find the optimal working conditions. **Figure 4c** shows the response of a Cu₂₀ (CuO/Cu₂O sample) to different gaseous compounds (hydrogen, *n*-butanol, 2-propanol, ethanol, acetone and ammonia) at the operating temperatures between 250–350 °C. The standard deviation between multiple measurements of the same sample set is indicated with error bars. For ethanol vapors, the response of the sensor at 350 °C is larger compared to the other gases. Sample sets tested at 250, 300 and 350 °C have the highest response to ethanol with values of about 24%, ~121% and ~140%, respectively, whereas **Figure 4c** also shows that the hydrogen response improved with the operating temperature. The sensing response of the CuO/Cu₂O heterostructure with a thickness of between 20 and 30 nm demonstrates the *p*-type behavior of this material. The response variation of the structure to 100 ppm of ethanol depending on its operating temperature was investigated to find the optimal working conditions, **Figure 4d** indicates the dynamic response of the Cu₂₀ sample to 100 ppm of ethanol. We observe that the response time (t_r) and recovery time (t_d) are relatively small under the operating temperatures of 250 °C, 300 °C and 350 °C (t_r = 18.8, 13 and 18.5 s, t_d = 43.3, 46.4 and 49.6 s, respectively). Then, we have prepared more complex

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structures based on TiO₂/CuO/Cu₂O and studied their sensing properties towards H₂ and ethanol.

After this, the effect of each layer can be explained in detail.

Gas sensing performances of TiO₂/CuO/Cu₂O. The hydrogen and ethanol responses of the samples consisting of the TiO₂/Cu₂O/CuO multi-nanolayered films with different thicknesses are studied to determine the impact of adding TiO₂ to the top layer. The ternary heterojunction can also act as a self-cleaning surface or to enhance the gas detection and to improve the protection against corrosion at high temperatures^{27–30,55,56}.

Figure 5a shows the response to hydrogen and ethanol of the TiO₂/CuO/Cu₂O Cu20, Cu30, Cu40, Cu40 and Cu60 samples, measured at the operating temperature of 350 °C. From **Figure 5a** we can see that the TiO₂/CuO/Cu₂O Cu20 and Cu30 samples are more sensitive towards ethanol compared to the hydrogen gas. However, increasing the thickness of CuO/Cu₂O to Cu50 and Cu60 changes the sensing performance of TiO₂/CuO/Cu₂O structures and the most sensitive gas becomes to hydrogen, although the response values are smaller. However, the response values of sensors with the thicker layers of CuO/Cu₂O are lower towards hydrogen and ethanol compared to that of Cu20 and Cu30. The best sensing performances to ethanol were obtained for the samples with the thickness of 30 nm (sample Cu30). **Figure 5b** illustrates the electrical current-voltage plot of the TiO₂/CuO/Cu₂O samples, and the linear Ohmic behavior at room temperature can be seen in **Figure S7b**. However, as already stated, at the operating temperature of 350 °C (**Figure S8b**) the *I-V* current-voltage characteristics is non-linear due to a conductivity effect driven by the energy barrier. **Figure 5c** presents the responses to different chemical compounds (hydrogen, ethanol, 2-propanol, *n*-butanol, acetone and ammonia) versus the operating temperature of the TiO₂/CuO/Cu₂O Cu20 sample set. At all operating temperatures, i.e. 250 °C, 300 °C and 350 °C, we see that the samples have the highest response to ethanol with responses of ~38%, ~115% and ~121%, respectively. Comparatively (the data in **Figure 4c** and **Figure 5c**), the response value

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was enhanced with increasing operating temperatures for the TiO₂/CuO/Cu₂O nanomaterial. Thus, there was no noticeable response for these sensor-structures below OPTs of 250 °C due to surface reactions with different oxygen species. The maximum response is achieved at an operating temperature of 350 °C, which is the highest temperature allowed by our gas test equipment. **Figure 5d** indicates the dynamic response to 100 ppm of ethanol of the TiO₂/CuO/Cu₂O samples (prepared based on the Cu20). For the working temperatures 250, 300 and 350 °C, we observed that the response times (τ_r) are 18.9, 16.3 and 13.8 s) and recovery times (τ_d) are 35.5, 28.3 and 37.3 s, respectively) are relatively low.

Comparison of the gas sensing features of Cu₂O/CuO and TiO₂/Cu₂O/CuO. Here we compare the gas response of the multi-nanolayered Cu₂O/CuO and TiO₂/Cu₂O/CuO heterostructured films.

Comparing the data in **Figure 4c** and **Figure 5c** indicates that the response to hydrogen has increased for the ternary system with respect to the heterojunction at all operating temperatures. **Figure 6a** represents the relationship between the response to 100 ppm of ethanol vapors and the power consumption versus the applied voltage for the CuO/Cu₂O Cu20 samples. **Figures 6a** and **6b** show that when the applied bias voltage decreases, the response to the ethanol vapors increases and the power consumption decreases, which effect can be attributed to the nonlinear I-V current-voltage characteristics at OPT 350 °C (**Figure S8a** and **S8b**). For an applied bias voltage of 0.5, 0.01 and 0.005 V, the power consumption is about 88, ~0.02 and ~0.004 μ W (approximately 4 nW), and response is about 140%, ~160% and ~300%, respectively. This power consumption is part of the energy consumption of the entire sensor system, because the energy consumption to heat the sample to OPT 350 °C is not calculated here.

Figure 6b represents the response to 100 ppm of ethanol vapors and the dependence of the power consumption versus the applied bias voltage for the TiO₂/CuO/Cu₂O Cu20 sample set. This

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figure shows that when the applied voltage decreases, the power consumption also decreases. For example, the power consumption is about 1.5, ~0.33, ~0.05 and ~0.001 μ W (approximately 1 nW) at the applied voltage of 0.25, 0.1, 0.05 and 0.01 V, respectively. However, the gas response is ~120% for the applied bias voltage of 0.25 V, which decreases to ~97% under 0.1 V of an applied bias voltage, which increased to ~160% and ~222% at the applied bias voltage to 0.05 V and 0.01 V, respectively.

Figure 6c represents the variation of the response of the CuO/Cu₂O Cu₂₀ (curve 1) and TiO₂/CuO/Cu₂O Cu₂₀ (curve 2) samples to 100 ppm of ethanol vapors for 105 days. **Figure 6c** shows that the response to ethanol vapors has bisected for the (curve 1) samples, whereas it remained unchanged in the case of the TiO₂/CuO/Cu₂O (curve 2) samples significant changes were not observed due to the self-cleaning effect of the ternary heterostructure^{55,56}.

According to the results presented in this study, changes in the conductance of semiconductor oxide-based chemical sensors are caused by interactions with environmental chemical compounds with the sensor surface, which are strongly influenced by the operating temperature^{2,57}. An efficient way to improve the characteristics of the sensor is through control of the catalytic properties of the oxide surface^{2,42}.

Figure 7 shows the gas response to 1, 5, 10, 50, 100, 500 and 1000 ppm ethanol () versus the type of CuO/Cu₂O and TiO₂/CuO/Cu₂O samples Cu₂₀, Cu₃₀, Cu₄₀, Cu₅₀ and Cu₆₀ measured at the working temperature of 350 °C. **Figure 7** suggests that all samples responded to all concentrations of ethanol.

Figure 8 shows the dynamic response of the CuO/Cu₂O (**Figure 8a**) and TiO₂/CuO/Cu₂O (**Figure 8b**) samples with the thickness of 20 nm (Cu₂₀) and 30 nm (Cu₃₀) to 1, 5, and 10 ppm of ethanol at 350 °C of working temperatures. From the plot it can be seen that the response of the sensors is quite high at very low concentrations of ethanol. The sensing parameters of CuO/Cu₂O

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and TiO₂/CuO/Cu₂O samples with the thickness of 20 nm (Cu20) and 30 nm (Cu30) at operating temperature of 350 °C to 1, 5, and 10 ppm of ethanol are shown in **Table S1**. For the CuO/Cu₂O and TiO₂/CuO/Cu₂O Cu20 and Cu30 samples, the dynamic response to 1 ppm of ethanol is shown in **Figure S9**.

The dynamic response to various concentrations of ethanol (1, 5, 10, 50, 100, 500 and 1000 ppm) by the CuO/Cu₂O and TiO₂/CuO/Cu₂O Cu20 and Cu30 samples is shown in **Figure S10**. The gas response to high concentrations of hydrogen is presented in **Figure S11**, which shows the effect of different concentrations of hydrogen (100, 500 and 1000 ppm) versus the type of CuO/Cu₂O and TiO₂/CuO/Cu₂O Cu20, Cu30, Cu40, Cu50 and Cu60 sample sets measured at 350 °C of working temperature. **Figure S11** displays that the response of the samples increases strongly at high concentrations of hydrogen.

Overall, the chemical detection mechanism proposed for CuO/Cu₂O relies on surface physico-chemical reactions, which strongly depend on the operating temperature. More details of the gas detection mechanism are described in **Text S4** (Supporting Information) and the energy band diagrams of heterostructures TiO₂/CuO/Cu₂O in air and ethanol vapors are represented in **Figure S12**.

3.5. Simulation of bulk phases

We first investigate the optimised bulk structures of the materials used to build the heteroepitaxial structures, which are later used to simulate the molecular adsorption to compare with our experiments. Our starting point for the substrate is Cu₂O, characterised by the space group $Pn\bar{3}m$ (no. 224) and the cuprite structure, with Cu in the lowest oxidation state of 1+⁵⁸. Erreur ! Source du renvoi introuvable. **Figure 9a** presents the conventional cubic unit cell of Cu₂O with two formula units (f.u.). The O atoms are distributed in a body-centred cubic (*bcc*) sublattice, whereas

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the Cu ions are arranged in a face-centred cubic (*fcc*) sublattice. The 2-fold Cu atoms occupy the 4*b* Wyckoff linear positions with coordinates at the origin of the unit cell, while the 4-fold O ions occupy the 2*a* tetrahedral crystallographic sites at (1/4, 1/4, 1/4). CuO, the material in the middle of the ternary heterostructure with Cu in the higher oxidation state of 2+, crystallises in the tenorite structure with space group *C2/c* (no. 15)⁵⁹. **Figure 9b** shows the conventional monoclinic unit cell of CuO with 4 f.u. In this structure, the 4-fold Cu atoms are located at the 4*c* Wyckoff square planar positions with coordinates (1/4, 1/4, 0), whereas the 4-fold O counter-ions fill the 4*e* distorted tetrahedral holes at (0, *y*, 1/4). CuO forms two sets of $\infty^1[\text{CuO}_{4/2}]$ chains perfectly aligned along the [110] and [1 $\bar{1}$ 0] directions⁶⁰. The deviation of the *y* value from 1/2, in fractional coordinates, represents the staggering of the $\infty^1[\text{CuO}_{4/2}]$ chains along the [001] direction. The crystal structure of anatase TiO₂, the topmost material in the ternary heterojunctions, is tetragonal with space group *I4₁/amd* (no. 141)⁶¹. The conventional unit cell contains 4 f.u. of TiO₂, as depicted in **Figure 9c**. The distorted octahedral Ti²⁺ cations are in the 4*b* crystallographic positions with coordinates (0, 1/4, 3/8), and the O anions are in the 8*a* distorted trigonal planar sites at (0, 1/4, 1/6). The anatase TiO₂ structure displays channels in the [100] and [010] directions and dual chains $\infty^1[\text{TiO}_{6/2}]$ along the [2,2,13] direction.

Table S2 shows the optimised and experimental lattice parameters for the cubic unit cell of Cu₂O, the monoclinic unit cell of CuO and the tetragonal unit cell of TiO₂. Our calculations indicate that the lattice parameters were overestimated by 0.41% for CuO⁵⁹ and by 1.59% for TiO₂⁶¹, while it was underestimated by 0.28% for Cu₂O⁵⁸. The internal coordinates of the three materials were allowed to relax, with all ions showing a large preference for staying in their perfect Wyckoff crystallographic positions. The value calculated for the parameter *y* is 0.004 larger than in experiments, indicating that we predict a smaller staggering for the $\infty^1[\text{CuO}_{4/2}]$ chains along the

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[001] direction in the simulated cell of CuO. The shapes of the conventional unit cells were fully optimised, but Cu₂O, CuO and TiO₂ remained in the ideal cubic, monoclinic and tetragonal structures, respectively. The perfect match between the simulated and experimental angle β further supports the undistorted monoclinic shape of CuO. More details and a description of the atomic Bader charges, atomic magnetic moments and band gaps are provided in **Text S5** (Supporting Information).

3.6. Simulation of isolated surfaces

We have also simulated the pristine Cu₂O(111) surfaces with hexagonal symmetry as well as the CuO($\bar{1}11$) and TiO₂(111) facets with monoclinic symmetry, that were used to construct the binary and ternary heterojunctions. The surface slabs were constructed using METADISE⁶² to cut the geometry optimized bulks. A vacuum gap of 20 Å was added above the surfaces to avoid spurious interactions between the periodic supercells. The two bottommost layers were kept at their relaxed atomic bulk positions, to simulate the bulk phase, while the rest of the slab was allowed relax explicitly. We have applied dipole corrections in the direction perpendicular to the surface^{63,64} to improve the description of the total energy of our single surface models^{65–69}. According to this formalism, a planar dipole sheet was introduced in the centre of the vacuum region, and its strength was calculated self-consistently to compensate the artificial adsorbate-induced dipole.

The Cu₂O(111) and CuO($\bar{1}11$) surface slabs have the smallest surface areas of 31.378 and 32.438 Å², respectively. The surface cell of Cu₂O contains 24 atoms, whereas CuO comprises 32 atoms distributed in four stacking sequences consisting of 2 and 4 f.u., respectively. The TiO₂(111) surface was modelled using a slab with an area of 54.580 Å² and 36 atoms occupying 12 stacking sequences of a single stoichiometric unit each. With this setup, we ensured that all surfaces were symmetric along the z axis and their widths were between 7.4 and 10.1 Å. The vacuum thickness

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as well as the total and relaxed number of surface layers were carefully tested until convergence to within 1 meV per atom was reached.

The stacking of the atomic layers is (O)–(Cu₄)–(O) for the Cu₂O surfaces in the (111) direction, with the atoms within parenthesis lying approximately in the same plane as shown in **Figure S13**. Termination *A* is a type 2 Tasker surface⁷⁰, with a top surface layer that has a bulk-like structure terminated in 3-fold under-coordinated O atoms and an 0.25 monolayer (ML) of monocoordinated Cu atoms with a single dangling bond. Termination *B* is a reconstructed type 3 Tasker surface⁷⁰, where the dipole moment was quenched by shifting half of the mono-coordinated Cu atoms from the topmost stoichiometric stacking sequence at the relaxed to the unrelaxed side of the slab, which created 2-fold O anions. Following relaxation of termination *A*, both the exposed 3-fold and subsurface 4-fold O atoms moved outwards by an average of 0.052 Å, while the cations migrated towards the bulk by only 42% of the displacement of the anions. We found that for termination *B*, the displacement of the atomic layers is larger than for termination *A*. The 0.5 ML mono-coordinated Cu atoms shifted their position horizontally by 0.689 Å, with half of them even coordinating the 2-fold O atoms, that only moved 0.141 Å towards the vacuum. This surface reconstruction and atomic displacements effectively increased the coordination number of both the exposed O and Cu atoms for the relaxed termination *B*.

Despite their different crystal structures, the stacking of the atomic layers for CuO in the ($\bar{1}11$) direction is similar to Cu₂O in the (111) direction, except for their different stoichiometric ratios, as illustrated in **Figure S14**. Unsurprisingly, both terminations *A* and *B* of the CuO($\bar{1}11$) surface display many of the same characteristics as its Cu₂O(111) counterpart. For example, termination *A* is a type 2 Tasker⁷⁰ surface presenting bulk-like structured 3-fold O and Cu atoms, of which 0.5 ML are 3-fold. Similar to the reduced copper oxide phase, the exposed atoms exhibit the lowest coordination numbers in termination *B* of CuO($\bar{1}11$), which is also a reconstructed type

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3 Tasker surface⁷⁰. Due to the surface construction, the 0.5 ML of Cu atoms left exposed are 2-fold, whereas 0.25 ML of the counteranions are 2-fold, 0.50 ML are 3-fold and the remaining have kept the distorted tetrahedral configuration of the bulk. After relaxation, the exposed O atoms of both terminations moved outwards by an average distance of ~0.2 Å, whereas the cations shifted horizontally their locations by 0.164 Å towards the vacuum in termination *A* and by 0.518 Å inwards in termination *B*, where they also became 3-fold.

TiO₂(111) is a type 2 Tasker surface⁷⁰ composed of thin (O)-(Ti)-(O) planes with a width of 0.220 Å and separated by 0.434 Å from the neighbouring stoichiometric units, which explains why only one termination is possible for this facet, as displayed in **Figure S15**. Cutting the bulk of TiO₂ reduces the coordination number of the exposed atoms, *i.e.* 0.5 ML of Ti become 4-fold, whereas the other half is surrounded just by three O ions, compared to the octahedral coordination environment in the bulk. Moreover, 0.75 ML of the anions are 2-fold and 0.25 ML are mono-coordinated in the freshly created surface. During relaxation, the least coordinated Ti ions moved 0.235 Å inwards. The 4-fold Ti remained approximately at the same position and the Ti of the third stoichiometric unit migrated 0.511 Å towards the surface, becoming all penta-coordinated and appearing roughly at the same layer. Moreover, all the O atoms have a coordination number of 2 after surface relaxation, with the largest average inwards displacement of 0.442 Å observed for the topmost stoichiometric unit. The anions of the third stoichiometric unit also experienced notable outwards shifts of 0.708 Å.

We have calculated the surface energy before (γ_u) and after (γ_r) relaxation as,

$$\gamma_u = \frac{E_u - n_{\text{bulk}} \cdot E_{\text{bulk}}}{2A}, \quad (2)$$

$$\gamma_r = \frac{E_r - n \cdot E_{\text{bulk}}}{A} - \gamma_u, \quad (3)$$

where n_{bulk} is the number of f.u. contained in the surface cell, E_u is the energy of the slab with all atoms at their optimised bulk positions, E_{bulk} is the energy of the bulk per f.u., E_r is the energy of

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the half-relaxed slab and A is the surface area of one side of the slab. The degree of relaxation (R) was quantified as $R = 100 \cdot (\gamma_u - \gamma_r)/\gamma_u$.

From our simulation of the surface energies, which are listed in **Table S3**, we have determined that terminations A are the most stable for Cu₂O(111) and CuO($\bar{1}11$), both before and after relaxation. The calculations also reveal that the overall most stable surface is Cu₂O(111) with a relaxed surface energy of $\gamma_r = 70 \text{ meV} \cdot \text{\AA}^2$. TiO₂(111) is the least thermodynamically stable plane in this study, since its surface energy is one order of magnitude larger than for any other material. Despite their similar relaxed surface energies, termination A of Cu₂O(111) only experiences a small degree of relaxation $R = 4.08\%$, while termination A of CuO($\bar{1}11$) suffers a relaxation approximately five times larger. Termination B of both copper oxide surfaces have the largest degree of relaxation, in agreement with the displacement of their respective atomic layers; while the TiO₂(111) facet shows a modest 9.69%. The lower charges of the undercoordinated atoms suggest that all surfaces are less ionic than their respective bulks. Our calculations indicate that the increasing order of ionic character for the surfaces is Cu₂O(111) < CuO($\bar{1}11$) < TiO₂(111), in line with the trend found for the charges in their respective bulk phases. The surface atoms of both Cu₂O(111) and TiO₂(111) are non-magnetic, as evidenced by their lack of magnetic moments. The exposed Cu atom in CuO experiences a reduction of its magnetic moment by $\sim 0.06 \mu_B \text{ atom}^{-1}$ in terminations A and B of the ($\bar{1}11$) surface. However, the magnetic moment is lower at $0.298 \mu_B \text{ atom}^{-1}$ and larger at $0.382 \mu_B \text{ atom}^{-1}$ for the O ions in terminations A and B , respectively, than in the bulk.

The work function (Φ), which measures the energy required to move an electron from the Fermi level (E_F) to the vacuum, was calculated as the difference between the potential of the vacuum (E_{vac}) and E_F . Based on this descriptor, the most reactive system is termination A of the Cu₂O(111) surface, since it only requires 4.831 eV of energy to provide the electron that can

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facilitate the detection of an adsorbed chemical species. However, termination A, which is the most stable plane of the CuO($\bar{1}11$) surface, has the largest work function at $\Phi = 5.943$ eV of any of the pristine facets reported in this study. Our results suggest that the different magnetic properties and stoichiometry of the Cu₂O(111) and CuO($\bar{1}11$) surfaces play major roles in the different trends observed for the work function values of terminations A and B of these materials. Moreover, the difference of the work function values is 0.192 eV for the least stable terminations B of the Cu₂O(111) and CuO($\bar{1}11$) surfaces. Our simulated work function values suggest that TiO₂(111) has an intermediate reactivity between the terminations A of Cu₂O(111) and CuO($\bar{1}11$).

3.7. Simulation of the multi-nanolayers-based heterojunctions

Based on the results for the pristine surfaces, we have analysed the thermodynamic stability, atomic structure and electronic properties of the binary CuO($\bar{1}11$)/Cu₂O(111) and ternary TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) multi-nanolayered interfaces, with hexagonal symmetry. For the binary CuO($\bar{1}11$)/Cu₂O(111) heterojunction, we used the 1×1 surface geometry and for the ternary heterostructure TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111), we employed the 1×2 supercell. Since in section 3.9 we will investigate the adsorption properties of the 2×2 ternary heterojunction interfaces, we decided to simulate these structures using 2, 1 and 4 stoichiometric stacking sequences for Cu₂O(111), CuO($\bar{1}11$) and TiO₂(111), respectively. This configuration ensures that we model 2×2 ternary heterojunctions of approximately 10.63 Å of width and containing 104 atoms, as bigger systems have a prohibitively larger computational cost. We report these heterostructures using the most stable termination A calculated for CuO($\bar{1}11$) and Cu₂O(111) and the only termination of TiO₂(111). All surfaces comprise incomplete top layers of O atoms, which are complementary in a “jigsaw puzzle” fashion, allowing the sub-surface Cu and Ti ions with dangling bonds to increase their coordination numbers. For the interface computations, we used

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the equilibrium lattice vectors for Cu₂O(111) to mimic the heteroepitaxial growth of CuO($\bar{1}\bar{1}1$)

and then TiO₂(111) adlayers. The Cu₂O(111) is perfectly hexagonal, but CuO($\bar{1}\bar{1}1$) and TiO₂(111)

have monoclinic symmetries, which leads to a mismatch of their lattice parameters and angles.

Figure 10 displays the side views of the supercell slabs used to simulate the binary and ternary heteroepitaxial junctions. 50% of the exposed and 25% of the subsurface Cu atoms experience an outward relaxation of approximately 0.13 Å after the formation of CuO($\bar{1}\bar{1}1$)/Cu₂O(111), with respect to the pristine surfaces. Moreover, 50% of the subsurface anions shift their positions by ~0.22 Å towards the surface in the binary device compared to the isolated materials. Our calculations suggest that the remaining atoms in the binary heterostructure suffer only a relatively smaller displacement after the deposition of the CuO($\bar{1}\bar{1}1$) layer. Next, we have applied a layer of TiO₂(111) and found that all the Ti atoms relaxed towards the bulk to roughly form a single atomic layer in the ternary heterostructure, which is similar to the atomic displacements of the pristine surface.

The scanning tunnelling microscopy images (STM) were constructed using the basic formulation of the Tersoff-Hamann approach⁷¹ and the HIVE code⁷², which has been successfully employed in previous works providing images in agreement with experiments^{66,73,74}. The sign of the sample bias applied for the production of the STM image provides information of the valence of conduction bands in the vicinity of the Fermi level (E_F). For example, the positive (negative) bias of $V = 1.0$ (-0.5) eV applied for the production of the STM image of the binary (ternary) heterostructure indicates that the electrons are moving from the probe tip to the conduction band (valence band to the probe tip). The brightest spots resolved for the surface of the binary CuO($\bar{1}\bar{1}1$)/Cu₂O(111) device correspond to the protruding O atoms located at the ridges along the $[2\bar{1}\bar{1}]$ direction, as shown in **Figure 10a**. The image also resolves the O atoms situated at the grooves, whose lowest brightness can be used to evaluate their relative position with respect to

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those anions at the tip of the ridge. The 3- and 4-fold Cu atoms can be seen, and their different sizes can also be used to differentiate them. **Figure 10b** clearly displays the honeycomb structure of the ternary TiO₂(111)/CuO($\bar{1}\bar{1}\bar{1}$)/Cu₂O(111) interface, whose corners are defined by the O represented by the brightest spots. Although Ti are the least noticeable ions, they are still well-defined circles occupying the trigonal holes created by the O atoms.

Table S4 lists the geometric misfit parameter (ζ), which was quantified as $\zeta = 100 \cdot [1 - 2\Omega/(A_1 + A_2)]$, where Ω is the overlap area and A_1 as well as A_2 are the surface areas of the materials forming the interface^{75,76}. The geometric misfit parameter is just under 1.7% for the binary heterojunction and 6.97% for the ternary interface. Note that typically, the misfit parameter is below 5% for stable heteroepitaxial junctions, but much larger values such as 18.6% and 8.3% have been measured for ZnO(0001)/Al₂O₃(0001) and MgO(111)/Al₂O₃(0001), respectively.⁷⁷ The values of this parameter indicate that the thin film CuO($\bar{1}\bar{1}\bar{1}$) suffered a smaller compression than the expansion experienced by the TiO₂-based layer upon deposition onto the substrate.

The interfacial free energy (σ_{int}) was calculated as;

$$\sigma_{\text{int}} = \gamma_r + (E_{\text{int}} - n_{\text{over}}E_{\text{over}} - E_{\text{sub}})/A, \quad (4)$$

where E_{int} and E_{sub} , are the energies of the interface and substrate, respectively; E_{over} is the energy of 1 f.u. in the bulk of the overlayer; and n_{over} is the number of f.u. in the overlayer.

The interfacial free energy calculated for both heteroepitaxial systems is positive, implying that these interfaces are stable with respect to their isolated bulk components. The difference in interfacial energy for the binary and ternary heterostructures is only 21 meV Å⁻², despite the large difference of their geometric misfit parameters. Note that the interfacial free energy of the interfaces is larger and therefore less stable than the relaxed surface energy of the substrate Cu₂O(111), explaining the carefully controlled experimental conditions required to prepare them. The calculated Bader charge is 0.139 e⁻ smaller for the exposed Cu atoms in

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CuO($\bar{1}11$)/Cu₂O(111) than in the pristine CuO($\bar{1}11$) surface. However, the subsequent deposition of the thin film TiO₂(111) increases the charge of the Ti atoms by only 0.024 e⁻ with respect to its pure surface. The formation of the binary heterojunction forces a large (modest) reduction by 0.448 (0.174) μ_B atom⁻¹ of the magnetic moment of the Cu (O) ions, compared to the value in the perfect CuO($\bar{1}11$) surface. Our simulations indicate that the magnetisation of the middle CuO($\bar{1}11$) thin film induces magnetic moments one order of magnitude smaller on the atoms of the TiO₂(111) top thin film. The DFT modelling indicates that the deposition of the CuO($\bar{1}11$) layer onto the substrate raises the work function by 1.454 eV with respect to the value of this descriptor in the pristine Cu₂O(111) surface. Likewise, the formation of the ternary heteroepitaxial junction leads to a further increase by 1.169 eV of the work function of the nanodevice.

3.8. Simulation of the molecular adsorption properties

We next investigated the adsorption properties of the binary and ternary layered heterojunction devices. We have considered the interaction of the single molecules probed experimentally, *i.e.* H₂, C₂H₅OH and *n*-C₄H₉OH, with the symmetrically inequivalent adsorption sites. We believe that these early molecular adsorption processes will determine the kinetics of the oxidation reactions that are discussed within the oxygen chemisorption model in the **Text S4** (Supporting Information). The O atoms from the VOC molecules were initially placed at 1.5 Å away from the closest transition metal atom of the sensing device and were subsequently optimised to their equilibrium configurations and energies. For the pristine heterostructures, the H₂ molecule was introduced interacting instead with the surface O atoms. Previous studies of nanosensor devices based on ZnO have shown that the interaction between these atoms of opposite Coulomb charges led to the most stable adsorption configurations^{78,79}.

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Table S5 lists the adsorption energies for the binding of the VOCs and molecular hydrogen onto both the exposed Cu and O atoms, respectively, of the binary heteroepitaxial structure CuO($\bar{1}11$)/Cu₂O(111). The largest adsorption energy $E_{\text{ads}} = -1.072$ eV was calculated for C₂H₅OH, which was followed by $E_{\text{ads}} = -0.833$ eV for *n*-C₄H₉OH on the most reactive under-coordinated 3-fold Cu site of the binary layered device. From a thermodynamic point of view, C₂H₅OH and *n*-C₄H₉OH have a strength of binding towards the undercoordinated 3-fold Cu site which is ~0.3 and ~0.05 eV larger, respectively, than their fully coordinated planar 4-fold counterpart. Moreover, the difference is also 0.05 eV for the energy released by the H₂ molecule on the two types of O atoms. Our simulations suggest that H₂ shows the lowest preference for the binary heterostructure, which is in agreement with the low gas response reported for this gas from the experiments. We found that the incorporation of the TiO₂(111) layer to the sensing device reduces the number of symmetrically inequivalent adsorption sites to one type of O atom and one type of Ti atom. The binding energy of the alcohol with the lowest (largest) molecular weight is reduced by approximately 0.22 (0.08) eV on the Ti positions of the ternary heterostructure with respect to the Cu atom of CuO($\bar{1}11$)/Cu₂O(111). However, the energetic preference of H₂ becomes larger by just 0.01 eV on the O site of TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) compared to the exposed anion on the binary heterojunction. The decreasing order of binding strength found for the alcohols on the Ti site and for H₂ on the O position is $E_{\text{ads}}(\text{H}_2) \gg E_{\text{ads}}(n\text{-C}_4\text{H}_9\text{OH}) > E_{\text{ads}}(\text{C}_2\text{H}_5\text{OH})$.

Figure 11 and **Figure 12** illustrate the most favourable molecular adsorption modes of H₂ and the alcohols on the surface of the binary and ternary heterojunctions, which is in agreement with previous studies^{78,79}. The small H₂ molecule is adsorbed almost perpendicularly to the surface on a 3-fold O site lying in the ridge along the $[\bar{2}\bar{1}\bar{1}]$ direction. This is the weakest interaction reported in this study, which is characterised by an interfacial H–O distance of 2.43 Å. H₂ was also initially placed above a 4-fold O atom in the groove along the $[\bar{2}\bar{1}\bar{1}]$ direction, but it moved

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outwards during optimisation. In the resulting adsorption, H₂ forms a bidentate-binuclear mode, since it lies flat to the surface spanning the gap between two opposite 3-fold O atoms in parallel ridges, as displayed in **Figure 11a**. The results from the calculations indicate that the H–O distance between H₂ and the surface is 0.14 Å larger across the 3-fold Cu side than the 4-fold Cu side. Nevertheless, the interaction with the surface led to a negligible stretching by 0.01 Å of the intratomic H–H distance with respect to the isolated molecule, but only for the adsorption at the groove site. We did not find evidence of H₂ dissociation in our calculations and thus, we speculate that this process is unlikely from a thermodynamic point of view. We found that the O of C₂H₅OH binds the surface 3- and 4-fold Cu atoms at 2.15 and 2.43 Å, respectively, as shown in **Figure 11a**. Given its small size, this molecule is able to place itself parallel to the surface grooves along the [2 $\bar{1}$ 1] direction, by forming hydrogen bonds to the exposed 3-fold O atoms at the top of the ridge. Interestingly, we calculated the shortest distance of 1.95 Å for the interaction between the hydroxyl H and the surface O of the [CuO_{3/2}] unit nearest to the coordinated 3-fold Cu. However, C₂H₅OH forms hydrogen bonds with the O belonging to the same [CuO_{4/2}] unit of the coordinated 4-fold Cu atom at the largest distance of 2.02 Å. The shortest distances for the interaction of C₂H₅OH at the 3-fold Cu position explain our strongest calculated DFT adsorption energy for this site. We observed a difference of just 0.08 Å for the interfacial bond distances between the 3- and 4-fold Cu sites and the O atoms for the heavier *n*-C₄H₉OH molecule, see **Figure 11c**. Moreover, the hydrogen bond distances between the OH group and the surface 3-fold O atoms is 1.92 Å for both adsorption configurations investigated here. Unlike C₂H₅OH, the large and non-polar hydrocarbon chain of *n*-C₄H₉OH prefers to adsorb perpendicular to the surface plane for the interaction with the 3- and 4-fold Cu sites. Thus, this provides *n*-C₄H₉OH with the appropriate orientation and separation from the heterostructure to form its hydrogen bond to the surface O atom directly bound

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to the coordinated Cu atom. We have rationalised the trend of the adsorption energies of the VOCs based on their interatomic distances to the surface, which are inversely proportional.

We have integrated the Bader charges, which indicate that the largest transfer of $\Delta q = -0.119 e^-$ was found for the adsorption of the electron-rich H₂ molecule at the tetrahedral sp^3 dangling bond position of the ridge 3-fold O atom, as summarised in **Table S5**. Based on the large electronegative difference of $\Delta\chi_{\text{HO}} = 1.24$ (Pauling scale)⁸⁰, this adsorption geometry promoted an effective charge transfer mechanism from the adsorbate to the surface. Surprisingly, the smallest charge transfer takes place from the surface to the bidentate binuclear H₂ in the adsorption at the groove position, since the adsorbate is not in a configuration that facilitates an appropriate orbital overlap with the exposed O atoms. **Figure 11a** illustrates that the interfacial charge transfer vanishes for this adsorption mode, with the electron-rich region located between the H atoms, similar to the filled σ_{1s}^{b2} bonding molecular orbital, and electron-depleted regions σ_{1s}^* representing the empty antibonding molecular orbital, all lying on the symmetry axis. Moreover, all VOC molecules lost electronic density upon adsorption onto the binary heteroepitaxial junction, with the observed values of charge flow in line with the trend of simulated adsorption energies. **Figure 11a** and **11b** displays the charge rearrangements between the OH group of the VOCs and the surface Cu and O atoms of the binary heterostructure, which is more noticeable for C₂H₅OH than for *n*-C₄H₉OH.

The incorporation of TiO₂ does not change noticeably the most stable adsorption geometries for the VOCs with respect to the binary sensor, but affects the configuration of the interaction for H₂, see **Figure 12a**. For the most stable adsorption, we found that H₂ prefers to sit perpendicularly to the TiO₂(111) layer at 2.56 Å above any of the quasi-equivalent O atoms. H₂ is only able to coordinate a single anion, since this surface does not form grooves with under-coordinated O atoms at the top of the ridges that allow the flat bidentate binuclear adsorption of

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the molecule. **Figure 12b** illustrates that C₂H₅OH also coordinates an exposed Ti ion *via* its hydroxyl O atom, but the molecule adsorbs 0.09 Å further away from the surface site than in the case of CuO($\bar{1}11$)/Cu₂O(111). Given the close proximity of the surface Ti and O atoms, C₂H₅OH cannot form a strong interfacial hydrogen bond, with the H···O distance stretched to the atypical value of 2.30 Å. *n*-C₄H₉OH also forms a more detached adsorption mode on the surface of TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) than in the binary device, which even prevents the formation of a hydrogen bond with the exposed O atoms, as shown in **Figure 12c**. We found that the computed charge transfers between the adsorbates and the ternary device explain their adsorption energy values, see **Table S5**. For example, the charge transfer is almost *nil* for H₂, suggesting physisorption and in agreement with its lowest adsorption energies, whereas the VOC molecules donate electron density to the surface in agreement with the strength of their adsorptions. The charge transfers for the VOCs are also smaller for the ternary device than for the binary heterostructure, which compares well and explains their relative adsorption energy values. **Figure 12a** shows the intramolecular charge redistribution resembling the molecular orbitals of H₂, with a negligible influence of the surface of the sensor. The Ti ion coordinating the OH group of C₂H₅OH loses some electronic density charge, whereas the cation coordinating the O atom of *n*-C₄H₉OH is the least perturbed for any VOC molecule, see **Figure 12b** and **12c**.

We have provided strong evidence that the change in sensitivity is the consequence of applying a layer of TiO₂(111) to the CuO($\bar{1}11$)/Cu₂O(111) sensing device. Our DFT calculations show that the ternary heteroepitaxial junction displays smaller adsorption energies towards the VOCs than the binary nanodevice, which agrees with the gas response experiments discussed in section 3.4. We have rationalised the different behaviour of the CuO($\bar{1}11$)/Cu₂O(111) and TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) heterojunctions based on their atomic charges and surface structures. The Bader charges calculated for both $q_{\text{Ti}} = +2.235 e^-$ and $q_{\text{O}} = -1.130 e^-$ in the ternary

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device are larger than $q_{\text{Cu}} = +0.867 e^-$ and $q_{\text{O}} = -0.965 e^-$ in the binary heterostructure. This supports a stronger Coulombic attraction between an equally spaced OH group of the VOCs and the surface of TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) than the facet of the binary material. On the other hand, the separation of 3.29 Å between the exposed Cu and O atoms in neighbouring [CuO_{3/2}] units allows the efficient formation of both coordinate and hydrogen bonds between the CuO($\bar{1}11$)/Cu₂O(111) heteroepitaxial material and the OH group of the VOCs. However, the relative position and 1.90 Å separation between the surface counterions prevents the simultaneous formation of the Ti–OH and OH···O bonds between the adsorbate molecule and the TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) heterojunction. Our DFT simulations suggest that the alcohols show a larger preference towards formation of the Ti–OH coordinate bond instead of the hydrogen bond with the ternary heterostructure, given the larger Coulombic attraction of the former type of interaction compared to the latter one. Moreover, the modelling indicates that the surface structure and relative position of the atoms plays a more important role than the Bader charges in determining the geometries and energies for the adsorption of the VOCs. The non-polar H₂ is the only molecule that shows both a physisorbed mode on the CuO($\bar{1}11$)/Cu₂O(111) sensor with larger adsorption energy than the chemisorbed configuration. Although the least favourable of all adsorbates explored here, H₂ is also the only molecule that displays a slightly larger adsorption energy for its physisorption mode on the ternary than on the binary device. Thus, the calculations that we have carried out offer an additional and complementary understanding, from a molecular point of view, into the sensing mechanism of the binary and ternary heteroepitaxial junctions.

4. Conclusions

In this paper we have studied the heterostructure of mixed titania/cuprite/cupric oxide (TiO₂/CuO/Cu₂O) phases obtained *via* a facile, cost-effective, simple and reliable spray-

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sputtering-annealing approach. The investigated mixed oxide semiconductor phases are nanocrystalline and possess direct optical band gaps that are preferred for sensing applications, owing to direct recombination with the release of the energy. We have discussed the material properties and enhanced gas detecting characteristics of the CuO/Cu₂O nanolayered heterojunction device structure based on two precise phases, i.e. CuO and Cu₂O. The effect of the different thicknesses on the sensor performances is reported for the Cu₂O/CuO nanolayered crystalline heterojunction as well. The method presented in this work allows for the preparation of high-quality nanomaterials with different compositions and crystal phases. CuO/Cu₂O nano-layers with a thickness between 20 and 30 nm, which were prepared using conventional thermal annealing at 420°C for 0.5 h, stand out having promising sensing characteristics in terms of good selectivity and high response to volatile organic compounds. The best results, with a response of about 150% to 100 ppm of ethanol were achieved at a working temperature of 350°C.

The CuO/Cu₂O samples show a high selectivity and response to ethanol, which improved, alongside the long-term stability of the sensor, after surface coverage with a thin layer of TiO₂ titania. We also noted that, even after the deposition of titanium oxide, the response to ethanol did not change significantly. In general, *p-type* semiconducting oxides, such as CuO/Cu₂O, will allow a complete reaction of ethanol, which will result in increasing sensing performances to its vapors. Also, we found that the response to ethanol vapors has bisected for the CuO/Cu₂O samples, whereas it remained unchanged in the case of the TiO₂/CuO/Cu₂O samples significant changes were not observed due to the self-cleaning effect of the ternary heterostructure^{55,56}. These results provide evidence that the deposited thin TiO₂ film protects the sensor, increasing its lifespan. Thus, the titania (TiO₂) ultra-thin layers deposited on top of Cu₂O/CuO using spray pyrolysis method can be used for long-term stabilization of functionalities of sensors to provide protection against corrosion at high temperatures^{27–30,55,56}

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We have used first-principle techniques to simulate the formation of the binary CuO($\bar{1}11$)/Cu₂O(111) and ternary TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) heterostructures, as well as their reactivity towards H₂, C₂H₅OH and *n*-C₄H₉OH. We have investigated the deposition of the monoclinic CuO($\bar{1}11$) and TiO₂(111) thin film overlayers on the Cu₂O(111) surface substrate with hexagonal symmetry and found that despite the lattice mismatch, these are thermodynamically stable systems. The binary and ternary heterostructures are thermodynamically stable. The value of the work function rises with the number of components of the heterojunctions, which is consistent with their different sensitivities. The study of the affinity of the VOCs and H₂ towards the binary and ternary heterostructures shows that the adsorbates interact molecularly with the surface of the sensors. H₂ and C₂H₅OH release the smallest and largest adsorption energy in their ground state interaction configurations, respectively, on the surfaces of both nanodevices in agreement with their similar selectivity. The electronic properties of the ground state adsorption configurations alongside the surface structure are important descriptors that explain the trends of binding energies.

■ ASSOCIATED CONTENT

Supporting Information: Details on the surface properties of the binary and ternary heteroepitaxial interfaces simulated using DFT calculations as implemented within the VASP. Cross-section view of set #1 devices made from nanolayers of CuO/Cu₂O (Device 1), and sets #2 based on TiO₂/CuO/Cu₂O heterolayers (Device 2) are presented. Micro-Raman spectra of ultra-thin crystallite films of CuO/Cu₂O and of TiO₂/CuO/Cu₂O thermal annealed at 420°C in air with five different thicknesses are shown, respectively. Comparison of Micro-Raman spectra for CuO/Cu₂O with TiO₂/CuO/Cu₂O thermal annealed at 420°C for 30 min in air is mentioned. The cuprite oxide (Cu₂O) lattice in simple cubic according to the translational group it is described by

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over-all cubic symmetry. The zone centre normal modes for TiO₂, CuO and Cu₂O are explained.

Transmission spectra of TiO₂/CuO/Cu₂O heterostructure and plot of absorption near the UV edge vs. wavelength for same heterostructures are shown. Plot the $(\alpha h\nu)^2$ vs. photon energy $h\nu$ for CuO/Cu₂O, TiO₂/CuO/Cu₂O heterostructure is reported. Dynamic response to 1 ppm concentration of ethanol for CuO/Cu₂O and TiO₂/CuO/Cu₂O samples with thicknesses of 20 nm (Cu20) and 30 nm (Cu30). Dynamic response at different concentration of ethanol for CuO/Cu₂O (Cu20) samples, TiO₂/CuO/Cu₂O (Cu20) samples, CuO/Cu₂O (Cu30) samples and TiO₂/CuO/Cu₂O (Cu30) samples. Gas response at different concentration of hydrogen versus the type of samples for CuO/Cu₂O samples and TiO₂/CuO/Cu₂O samples with different thicknesses of 20 nm (Cu20), 30 nm (Cu30), 40 nm (Cu40), 50 nm (Cu50) and 60 nm (Cu60), respectively, measured at 350 °C. The sensing mechanism is proposed. Energy band diagrams of heterostructures TiO₂/CuO/Cu₂O in air and in ethanol vapors are illustrated. Calculated unit cell lattice parameters (a , b and c), atomic charges (q), atomic magnetic moments (m) and band gap energy (E_g) for the cubic cuprite Cu₂O, monoclinic tenorite CuO and tetragonal anatase TiO₂. Calculated staggering parameter y and angle β are reported for CuO. Oxidation state (OS) and experimental values for a , b , c , y , β , m and E_g are indicated for all phases. The calculated Bader charges are displayed. Side (top panels) and top (bottom panels) views of the optimised structures for terminations (a) A and (b) B of the Cu₂O(111) surface. Crystallographic directions and stacking sequence of the atomic layers are indicated. Side (top panels) and top (bottom panels) views of the optimised structures for terminations (a) A and (b) B of the CuO($\bar{1}11$) surface. Crystallographic directions and stacking sequence of the atomic layers are indicated. Side (top panel) and top (bottom panel) views of the optimised structures of the TiO₂(111) surface, crystallographic directions and stacking sequence of the atomic layers are indicated. Surface energies before (γ_u) and after relaxation (γ_r) and the percentage of relaxation (R) for terminations A and B of the pristine

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Cu₂O(111), CuO($\bar{1}11$) and TiO₂(111) surfaces. The average atomic charges (q), average magnetic moments (m) and work function (Φ), are also indicated for each surface. Geometric misfit parameter (ζ) and interfacial free energy (σ_{int}) for CuO($\bar{1}11$) on Cu₂O(111) and TiO₂(111) on CuO($\bar{1}11$)/Cu₂O(111). The average atomic charges (q), average magnetic moments (m) and work function (Φ), are also indicated for each surface. Adsorption energies (E_{ads}) and charge transfers (Δq) for H₂, C₂H₅OH and *n*-C₄H₉OH on the heterostructures CuO($\bar{1}11$)/Cu₂O(111) and TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111). The adsorption site of the adsorbate on the surface of the heterostructures is also indicated. A negative value of Δq denote that the adsorbate loses electronic charge. Response to 100 ppm of ethanol versus operating temperature of the samples CuO/Cu₂O and TiO₂/CuO/Cu₂O with different thicknesses of 20 nm (Cu20), 40 nm (Cu40), 50 nm (Cu50) and 60 nm (Cu60), respectively. Response to 100 ppm of hydrogen versus operating temperature of the samples CuO/Cu₂O and TiO₂/CuO/Cu₂O with different thicknesses of 20 nm (Cu20), 40 nm (Cu40), 50 nm (Cu50) and 60 nm (Cu60), respectively. Response to different gases (hydrogen, *n*-butanol, 2-propanol, ethanol, acetone and ammonia) versus operating temperature of TiO₂/CuO/Cu₂O sample set with thickness of 30 nm (Cu30) and dynamic response to 2-Propanol of TiO₂/CuO/Cu₂O sample set with thickness of 30 nm (Cu30) to different operating temperature. Response to different gases (hydrogen, *n*-butanol, 2-propanol, ethanol, acetone and ammonia) versus applied voltage of TiO₂/CuO/Cu₂O samples with thickness of 20 nm (Cu20). Response to different gases (hydrogen, *n*-butanol, 2-propanol, ethanol, acetone and ammonia) versus applied voltage of TiO₂/CuO/Cu₂O samples with thickness of 30 nm (Cu30). Dynamic response to *n*-butanol of TiO₂/CuO/Cu₂O sample set with thickness of 20 nm (Cu20). SEM images of the TiO₂/CuO/Cu₂O samples at lower magnification with different thickness of 20 nm and 40 nm. The Supporting Information is available free of charge on the ACS Publications website at <http://pubs.acs.org>

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Notes

The authors declare no competing interest.

AUTHOR CONTRIBUTIONS

O.L., N.A., M.H., and L.Z. have synthesized the CuO/Cu₂O/Cu nanomaterials and developed the synthesis methodology. A.V. and F.F. performed all XPS investigations, analyzed the XPS results and drafted part for the article. O.L., N.A., E.C., V.G. and S.N. fitted a technological procedure for micro-nanomaterial integration into the devices for detecting VOC. N.A., V.G., E.C. and O.L. realized the measurement of the detecting characteristics of the CuO/Cu₂O/Cu materials, analyzed the data and drafted part for the article. Th.P. and O.L. performed the optical measurements. Th.P. performed the titania deposition. O.L., D.S.-C., N.A., A.V., E.C., N.H.d.L., R.A., V.G. and S.H. analyzed the data from experiments and worked on the manuscript. D.S.-C. and N.H.d.L. carried out the DFT simulations and drafted computational part for the article. O.L., R.A., E.C., F.F., N.A., L.C., S.N., N.H.d.L. and D.S.-C. conceived and designed the study and gave the final approval of the draft to be submitted and to be published. O.L., E.C., L.C. and R.A. performed the design and conception for the studies, and approval. The work was written based on contributions by all authors, who all reviewed the manuscript.

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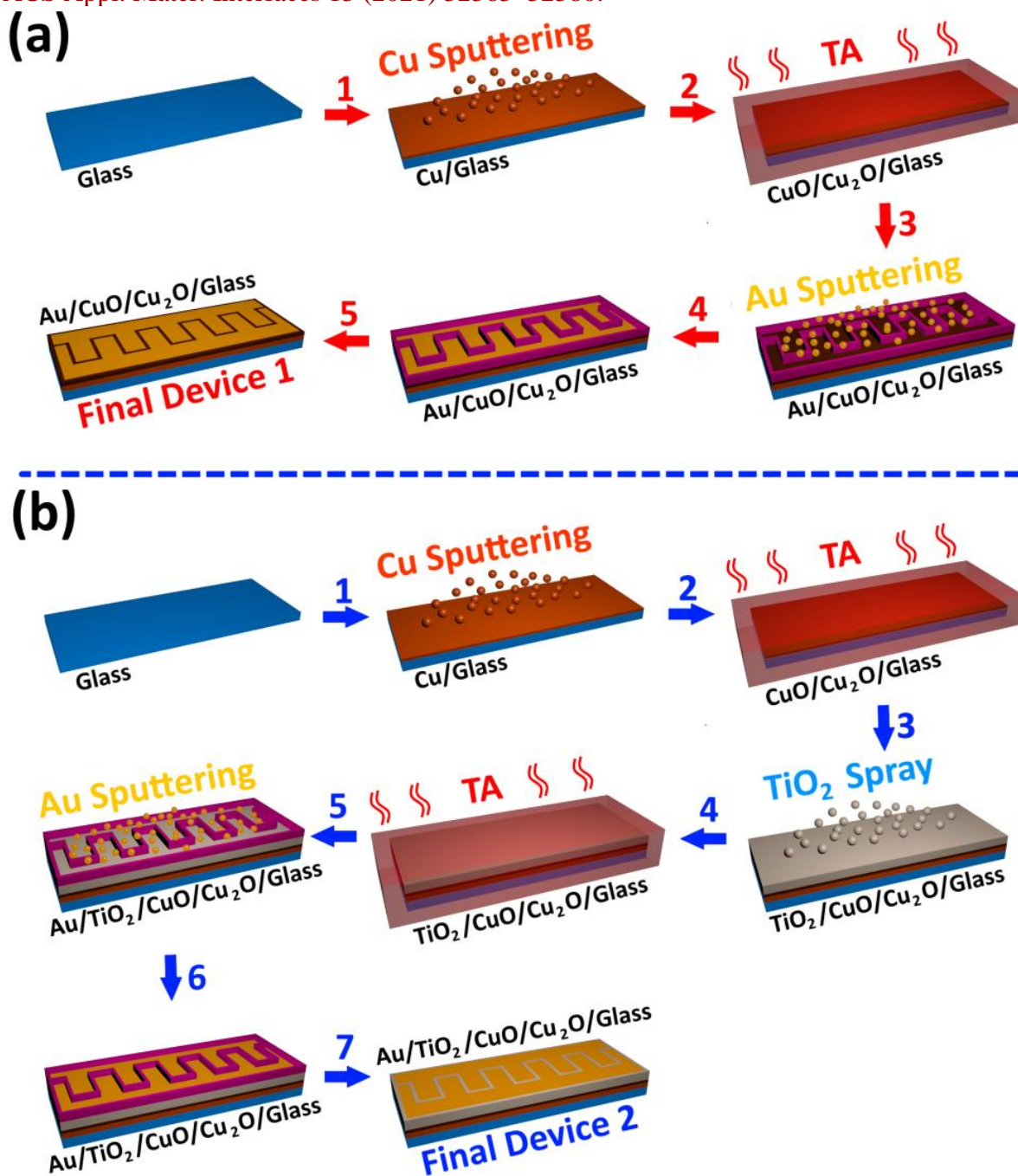


Figure 1. The technological flow chart for the manufacture of the (a) CuO/Cu₂O (Final device #1); and (b) TiO₂/CuO/Cu₂O (Final device set #2) nanostructured layered films sensor devices.

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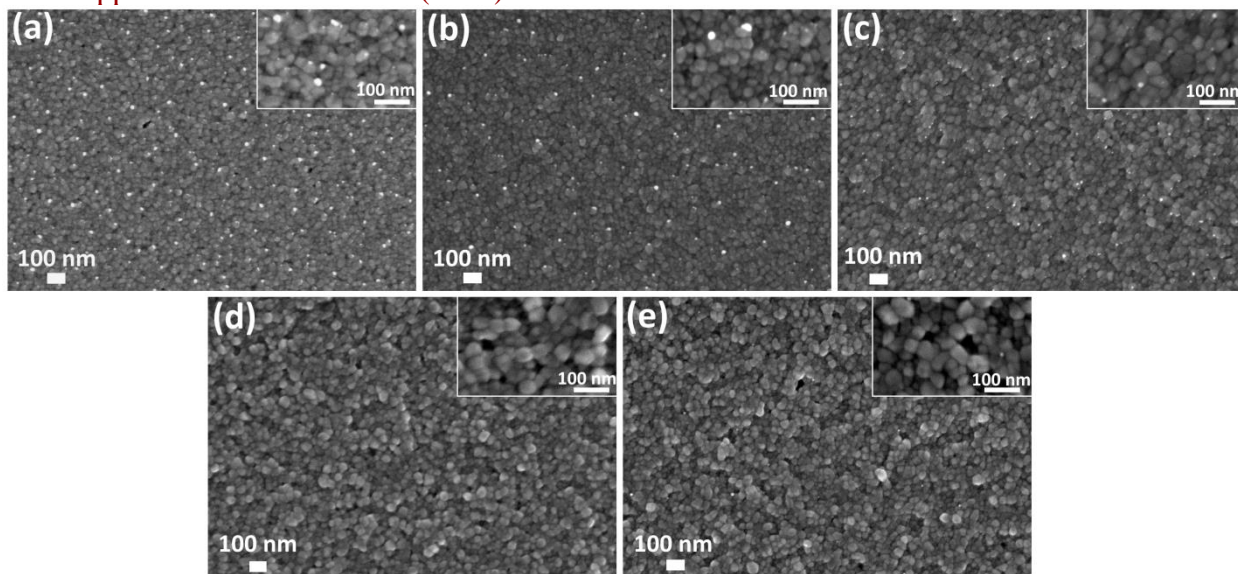


Figure 2. SEM images of the nano-crystallite CuO/Cu₂O samples grown using the sputtering-annealing approach and thermally treated at 420°C, for 30 min: (a) 20 nm; (b) 30 nm; (c) 40 nm; (d) 50 nm; and (e) 60 nm. The inset shows a higher magnification of the SEM images.

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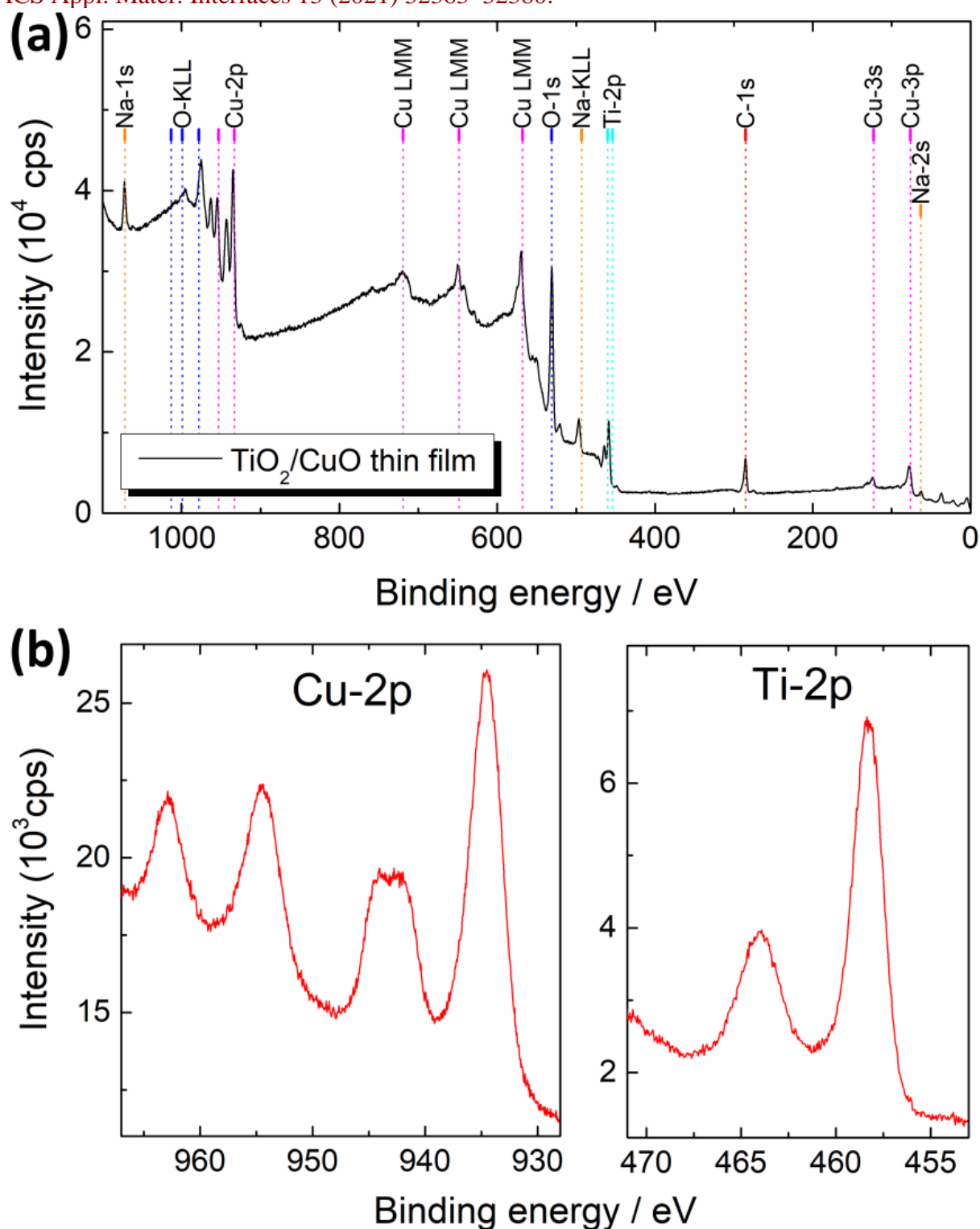


Figure 3. XPS spectra of a TiO₂-CuO (red line) thin film sensor: a) overview spectrum; b) high resolution spectra of the Cu-2p and Ti-2p lines.

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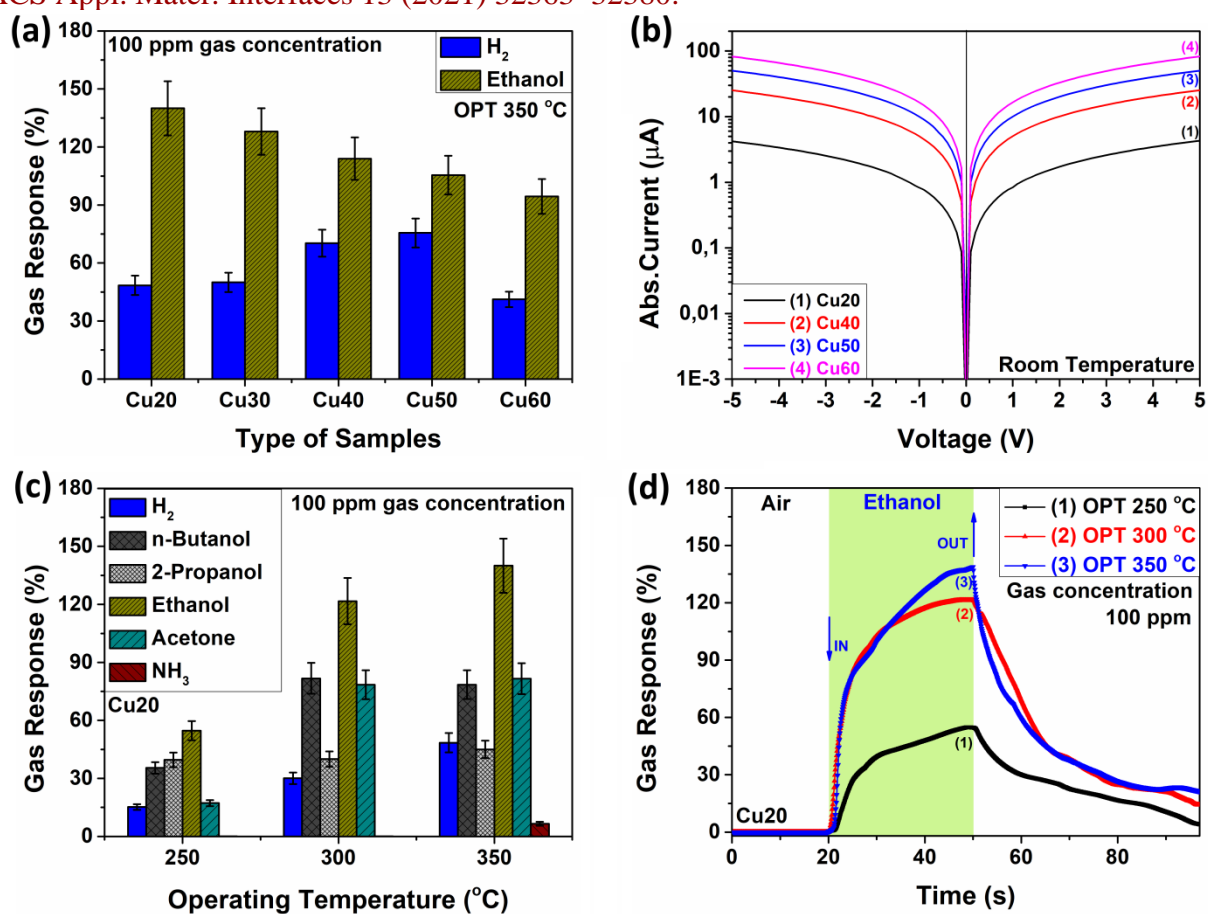


Figure 4. (a) Hydrogen and ethanol response of the CuO/Cu₂O samples with different thicknesses of 20 nm (Cu20), 30 nm (Cu30), 40 nm (Cu40), 50 nm (Cu50) and 60 nm (Cu60), respectively at the operating temperature of 350°C; (b) The *I-V* current-voltage characteristic of the CuO/Cu₂O Cu20, Cu4, Cu50 and Cu60 samples, measured at room temperature; (c) Response to different gases (hydrogen, *n*-butanol, 2-propanol, ethanol, acetone and ammonia) versus operating temperature of the CuO/Cu₂O Cu20 samples; (d) Dynamic response to ethanol of the CuO/Cu₂O Cu20 samples.

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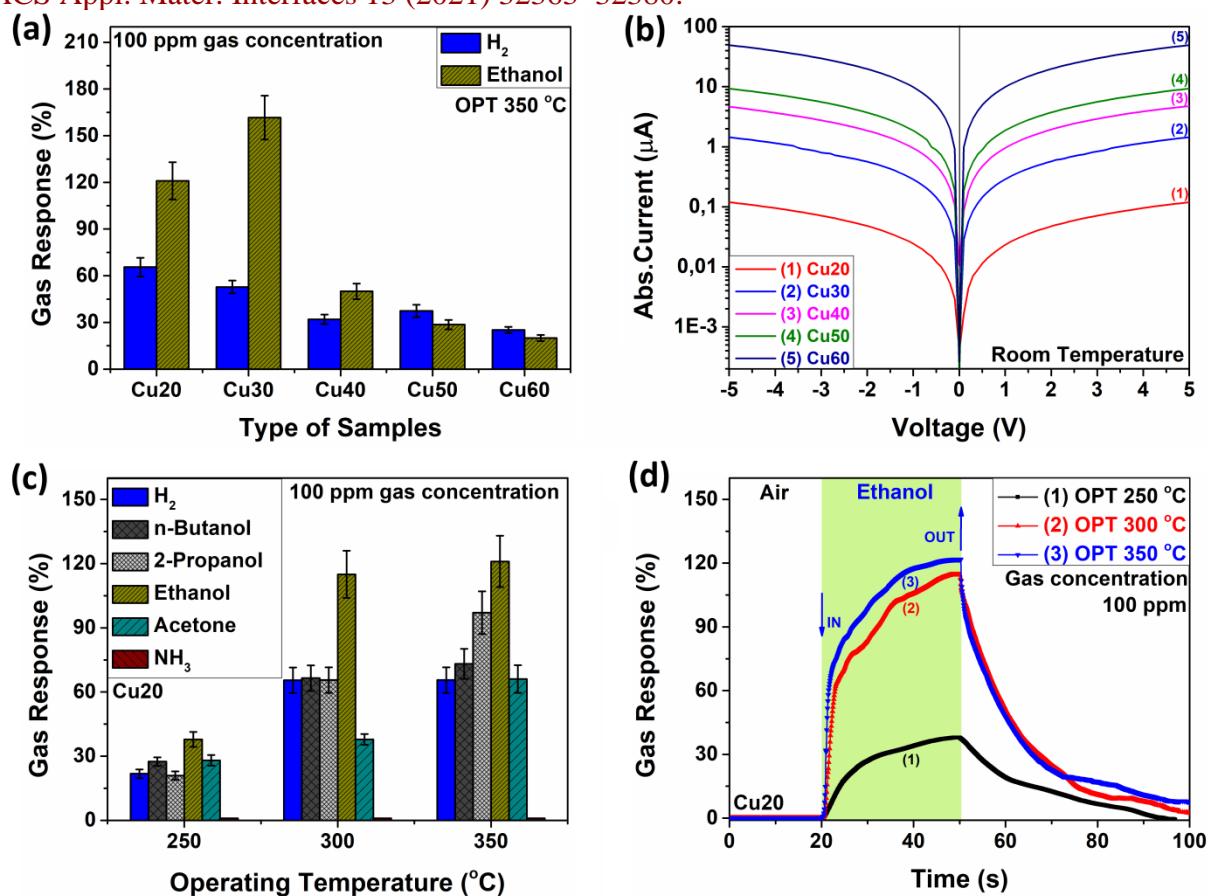


Figure 5. (a) Hydrogen and ethanol responses of the TiO₂/CuO/Cu₂O samples with different thicknesses of 20 nm (Cu20), 30 nm (Cu30), 40 nm (Cu40), 50 nm (Cu50) and 60 nm (Cu60), respectively, measured at 350°C; (b) The current-voltage characteristic at room temperature; (c) Response to different compounds (hydrogen, *n*-butanol, 2-propanol, ethanol, acetone and ammonia) versus operating temperature of the TiO₂/CuO/Cu₂O Cu20 samples; (d) Dynamic response to ethanol of the TiO₂/CuO/Cu₂O Cu20 samples.

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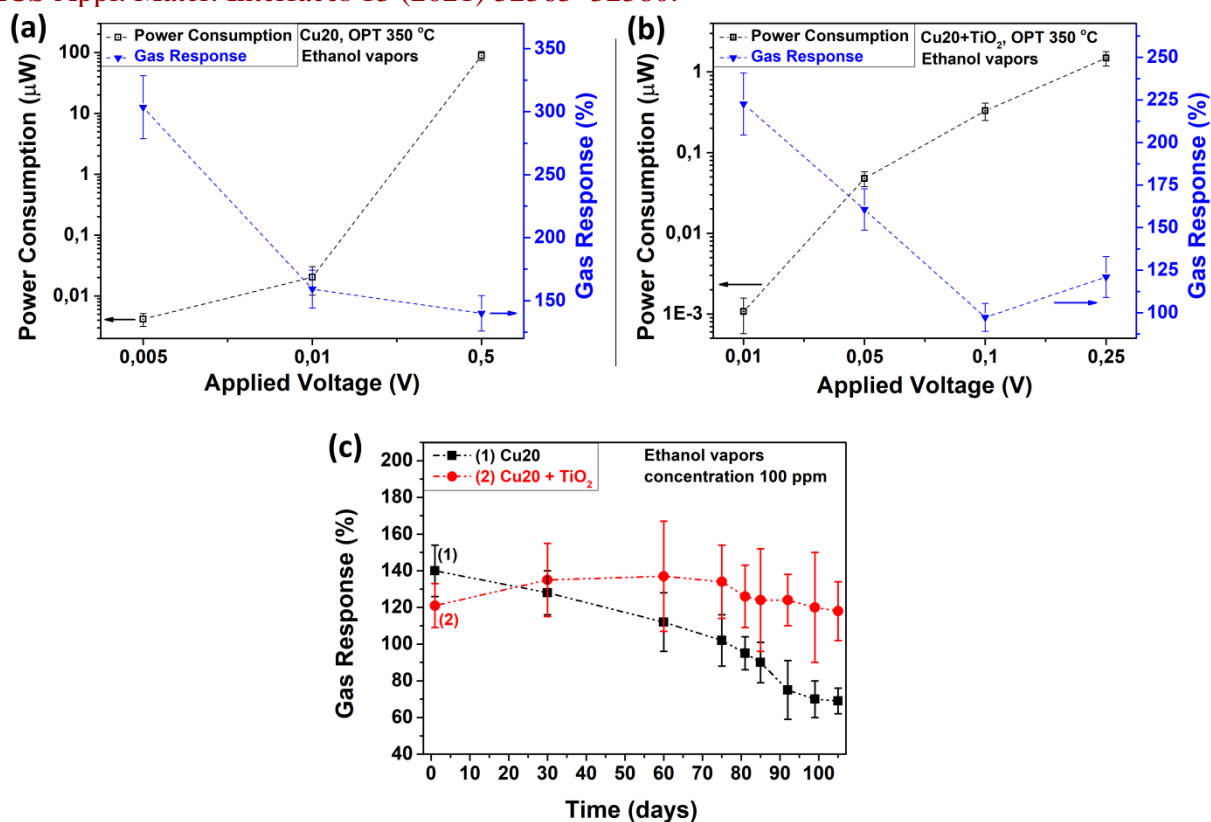


Figure 6. Dependence of the power consumption and the gas response with respect to the applied voltage for the (a) CuO/CuO₂ and (b) TiO₂/CuO/CuO₂ samples with thickness of 20 nm (Cu20); (c) Variation of gas response to ethanol vapors over time for the CuO/CuO₂ and TiO₂/CuO/CuO₂ samples with thickness of 20 nm (Cu20).

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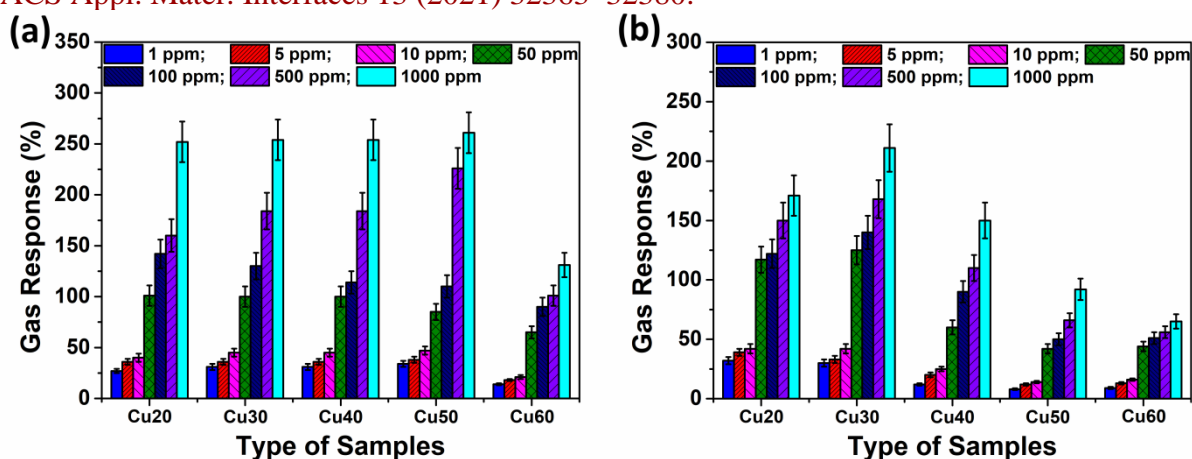


Figure 7. Gas response to various concentrations of ethanol vapor measured at 350 °C versus the type of samples for the (a) CuO/Cu₂O; and (b) TiO₂/CuO/Cu₂O samples with different thicknesses of 20 nm (Cu20), 30 nm (Cu30), 40 nm (Cu40), 50 nm (Cu50) and 60 nm (Cu60).

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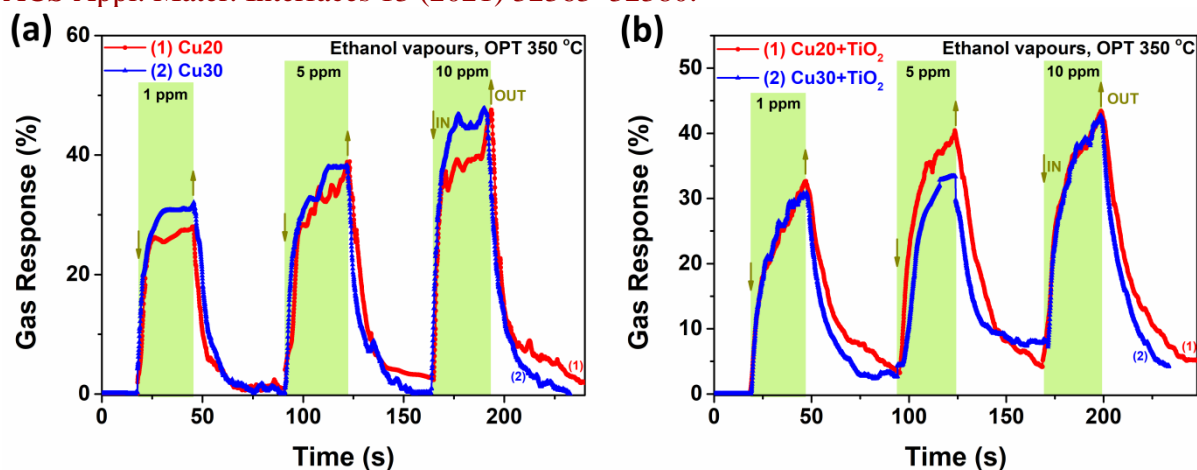


Figure 8. Dynamic response at 1, 5 and 10 ppm of ethanol vapor for the: (a) CuO/Cu₂O; and (b) TiO₂/CuO/Cu₂O samples with various thicknesses of 20 nm (Cu₂₀) and 30 nm (Cu₃₀).

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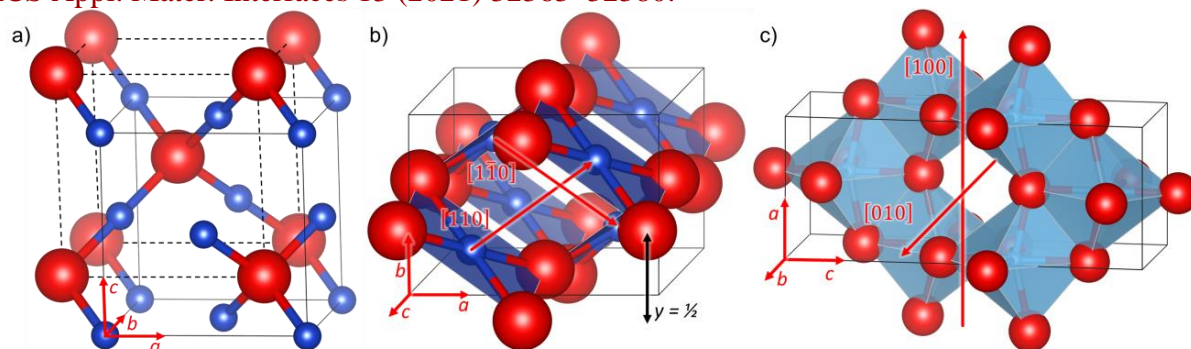


Figure 9. (a) Conventional cubic unit cell containing two formula units (f.u.) of cuprite Cu₂O. Solid lines represent the face-centred cubic (*fcc*) sublattice of Cu ions and dashed lines indicate the body-centred cubic (*bcc*) sublattice of O atoms. (b) Conventional monoclinic unit cell containing 4 f.u. of tenorite CuO. The $\infty^1[\text{CuO}_{4/2}]$ chains are shown along the $[110]$ and $[1\bar{1}0]$ directions, while the ideal staggering parameter y is represented in direct coordinates. (c) Conventional tetragonal unit cell containing 4 f.u. of anatase TiO₂. The channels along the $[100]$ and $[010]$ directions are shown. Crystallographic directions are indicated for all structures. O atoms are in red, Cu atoms are in dark blue and Ti atoms are in light blue. Polyhedral representations are used for the square planar Co atoms and the distorted octahedral Ti atoms.

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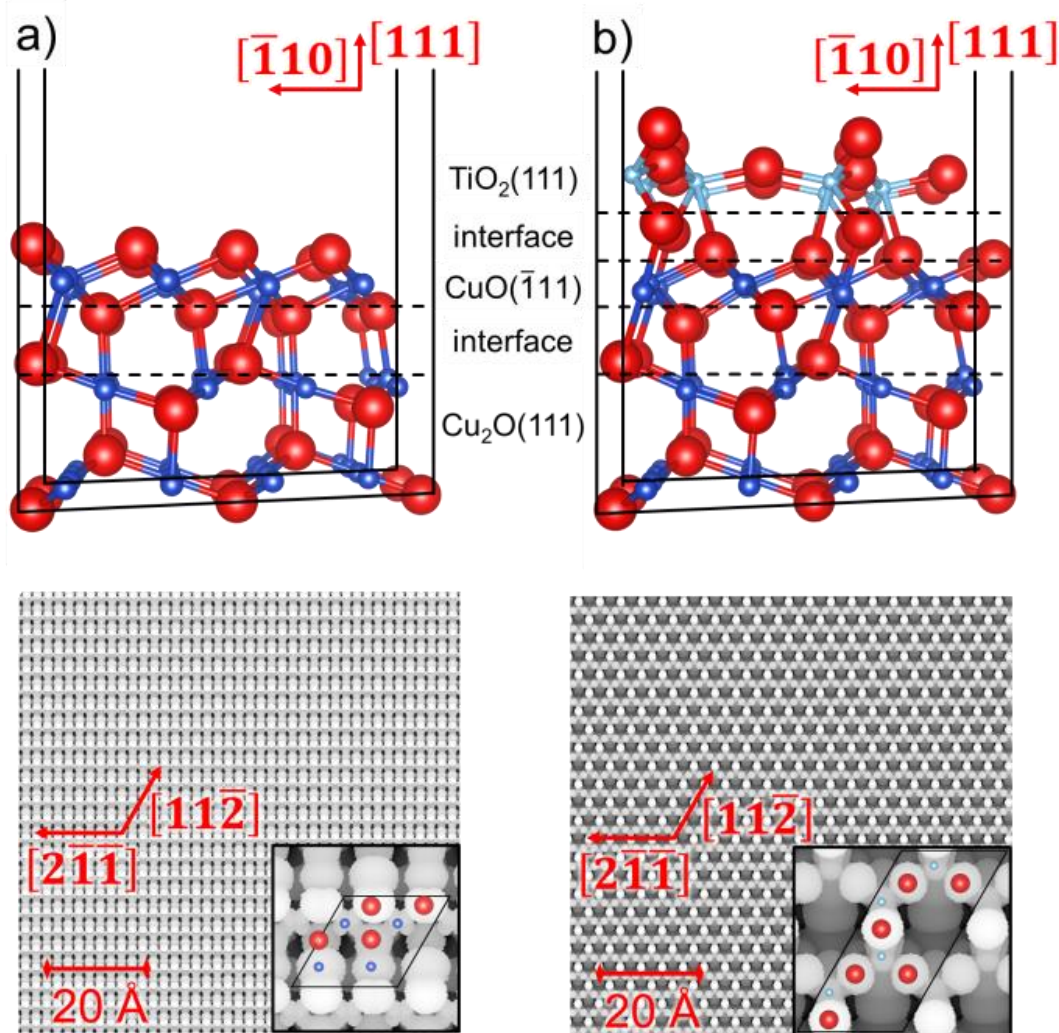


Figure 10. Top panels show the structure of the (a) binary CuO($\bar{1}11$)/Cu₂O(111) and (b) ternary TiO₂(111)/CuO($\bar{1}11$)/Cu₂O(111) interfaces. Bottom panels display the simulated scanning tunnelling microscopy (STM) images using a bias of: (a) $V = 1.0$ eV; and (b) $V = -0.5$ eV, a density of (a) $\rho = 0.0025$ e \AA^{-3} ; and (b) $\rho = 0.0010$ e \AA^{-3} as well as a tip distance of: (a) $d = 1.21$ \AA ; and (b) $d = 1.62$ \AA . Crystallographic directions are indicated with respect to the Cu₂O(111) substrate. O atoms are in red, Cu atoms are in dark blue and Ti atoms are in light blue.

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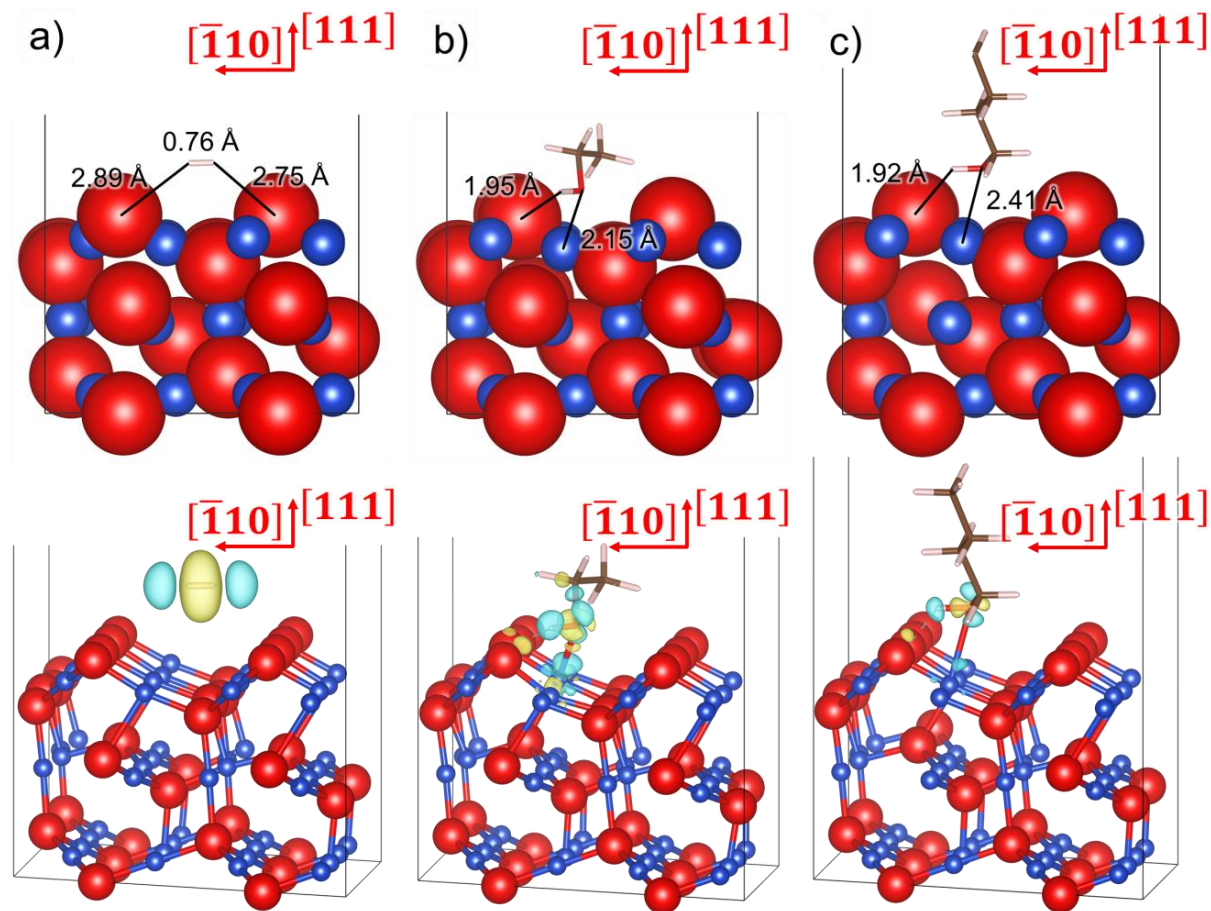


Figure 11. Adsorption of: (a) H₂, (b) C₂H₅OH and (c) *n*-C₄H₉OH on the CuO($\bar{1}11$)/Cu₂O(111) heterostructure. Interatomic distances are indicated in the top panels whilst the charge density flow ($\Delta\rho$) is represented in the bottom panels. Electron density gain and depletion regions are shown in yellow and green colours, respectively. Isosurfaces display a value of $\pm 0.005 \text{ e } \text{\AA}^{-3}$. Crystallographic directions are indicated with respect to the Cu₂O(111) substrate. The binary heterostructures are displayed using the (top panels) space-filling and (bottom panels) ball-and-stick representation, whereas the adsorbates are shown using the stick representation. O atoms are in red, H atoms are in white, C atoms are in brown and Cu atoms are in dark blue.

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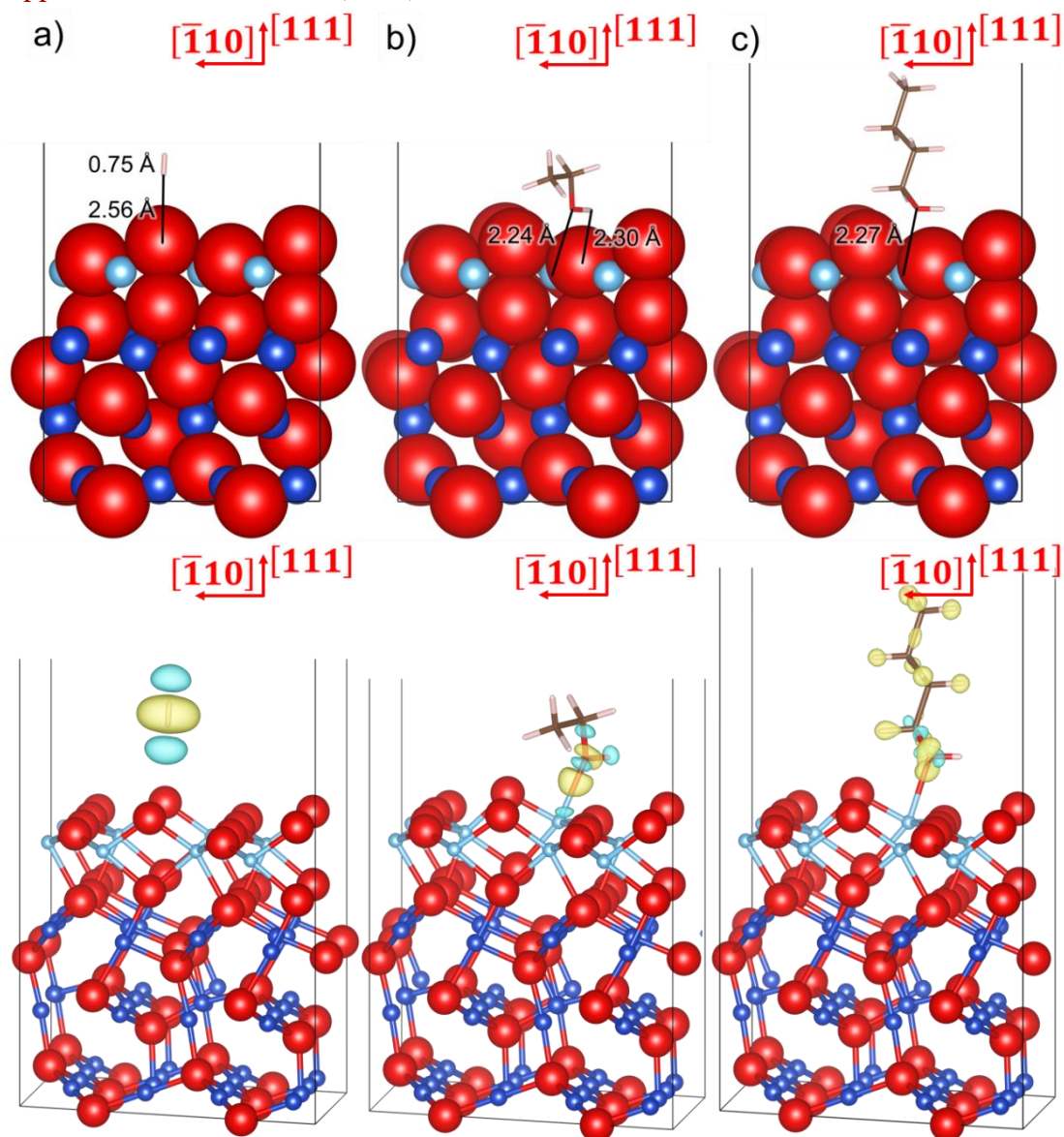


Figure 12. Adsorption of: (a) H₂, (b) C₂H₅OH and (c) *n*-C₄H₉OH on the TiO₂(111)/CuO($\bar{1}\bar{1}\bar{1}$)/Cu₂O(111) heterostructure. Interatomic distances are indicated in the top panels whilst the charge density flow ($\Delta\rho$) is represented in the bottom panels. Electron density gain and depletion regions are shown in yellow and green colours, respectively. Isosurfaces display a value of $\pm 0.005 \text{ e } \text{\AA}^{-3}$. Crystallographic directions are indicated with respect to the Cu₂O(111) substrate. The ternary heterostructures are displayed using the (top panels) space-filling and (bottom panels) ball-and-stick representation, whereas the adsorbates are shown using the stick

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